



**UNIVERSIDADE ESTADUAL DE CAMPINAS
INSTITUTO DE BIOLOGIA**

EVERTON ALVES MACIEL

**LINKING PATTERNS AND PROCESSES IN TREE PLANT COMMUNITIES TO
CONSERVATION AND MANAGEMENT OF THE SOUTH AMERICAN
SAVANNAS**

**VINCULANDO PADRÕES E PROCESSOS EM COMUNIDADES ARBÓREAS AO
PLANEJAMENTO E MANEJO DA CONSERVAÇÃO DE SAVANAS DA AMÉRICA
DO SUL**

**Campinas
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DO SUL**

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Tese apresentada ao Instituto de Biologia da Universidade Estadual de Campinas como parte dos requisitos exigidos para a obtenção do Título de Doutor em Biologia Vegetal.

Orientador: Prof. Dr. Fernando Roberto Martins

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*“Tivesse a noite límpida milhões de estrelas, mas
equidistantes e de igual brilho, como se admirariam o
Cruzeiro do Sul, a Estrela d’Alva, as Três Marias?
A diversidade é o encanto da vida”*

Leopoldo Magno Coutinho

RESUMO

As savanas da América do Sul (SAS) estão entre os biomas mais biodiversos do mundo que, apresenta grande lacuna de conhecimento (Déficit Linneano e Wallaceano) e de conservação, além de estar ameaçado por diversos fatores. No presente estudo, abordamos as seguintes questões na SAS: (i) Qual a importância do déficit florístico da SAS? (ii) A rede de Áreas Protegidas (APs) são suficientes para conservar as espécies arbóreas raras e comuns da SAS? (iii) A mudança no clima e na frequência do fogo e a defaunação tem efeito sinérgico na performance de diferentes grupos funcionais de árvores da SAS? Usamos os registros de árvores de levantamentos florísticos e fitossociológicos não identificados até o nível de espécie como um descritor do déficit florístico; as sete formas de raridade de Rabinowitz para avaliar o status de proteção da flora arbórea da SAS; e, três atributos funcionais – fenologia foliar, capacidade de rebrota e dispersão de sementes – para avaliar o efeito sinérgico dos fatores antrópicos na performance de espécies de árvores da SAS. Descobrimos: i) maior déficit florístico em levantamentos fitossociológicos do que em florísticos; ii) famílias muito diversas como Myrtaceae, mas também pouco diversas como Moraceae, apresentaram grande déficit florístico; iii) maior déficit florístico nas regiões periféricas do Cerrado; iv) o déficit florístico foi um preditor fraco da riqueza de espécies; e v) o déficit florístico não mudou ao longo do tempo. Descobrimos que 40,76% de 2.203 espécies de árvores da SAS eram raras. Dessa 2.203 espécies, 37,49% eram estenotópicas (distribuição restrita), 23,97% eram estenoéncias (especialistas de ecorregião) e 28,01% tinham população escassa. Algumas ecorregiões como Campos Rupestres, apresentaram muitas espécies raras não encontradas em nenhum outro lugar. Nossa estudo, portanto, revela a importância de revisar a lista de ecorregiões prioritárias para conservação (Global200). Existe um efeito sinérgico da mudança climática, do regime do fogo e da defaunação na perda de biomassa vegetal, podendo causar perda de mais de 70% da biomassa acima do solo, sobretudo em espécies decíduas, que não rebrotam após incêndios e são dispersas por animais. Isso mostra que a perda de biomassa vegetal da SAS resulta tanto de efeitos diretos como a incapacidade de lidar com perturbações, e efeitos indiretos como a perda de dispersores de sementes.

Palavras-chave: prioridade da conservação, déficit Linneano, déficit Wallaceano, espécies raras, biomassa acima do solo

ABSTRACT

The South American savannas (SAS) are among the more biodiversity biomes in the world which presents a floristic shortfall (Linnean and Wallace shortfall), a gap of conservation and also face several threats to biodiversity. In this study, we address the following questions in SAS: (i) What is the importance of the floristic deficit of SAS? (ii) Do protected areas enough to conserve rare and common SAS tree species? (iii) How do the synergism of climate change and the frequency of the fire and defaunation affect the performance of different plant functional types of SAS? The records of unidentified tree species were used to quantify the floristic shortfall. The seven forms of rarity were used to analyse the conservation gap in PA. Three traits - leave phenology, resprout capacity, and seed dispersal - was used to assess the synergistic effect of anthropic factors on the performance of SAS tree species. We found: i) greater floristic shortfall in phytosociological than in floristic surveys; ii) huge floristic shortfall in both highly diverse families as Myrtaceae, and low diverse as Moraceae; iii) a floristic shortfall mainly in peripheral regions of the Cerrado; iv) floristic shortfall was a poor predictor of species richness, and v) floristic deficit did not change over time. Of the 2,203 species, 40.76% were rare forms. Of these 2,203 species, 37.49% were stenotopic (restricted distribution in SAS), 23.97% were stenoecius (ecoregion specialists) and 28.01% had a sparse population. Some ecoregion such as Campos Rupestres had many rare species not found anywhere else. Our study, therefore, reveals the importance of reviewing the list of priority ecoregions for conservation (Global200). There is a synergistic effect of the change on climate fire and defaunation on the plant biomass loss. In some cases, the total biomass loss exceeded 70%. Deciduous species, which do not resprout after fires and are dispersed by animals, showed the highest rate of biomass loss. This shows that SAS plant biomass loss results from both direct effects such as an inability to deal with disturbances, and indirect effects such as loss of seed dispersers.

Keywords: priority conservation, Linnean shortfall, Wallacean shortfall, rare species, aboveground biomass

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CAPÍTULO 1: Introdução geral

1.1 Visão geral

A taxa de extinção de espécies no antropoceno já ultrapassa a do registro fóssil (Ceballos et al., 2017). Apenas para angiospermas, cerca de 600 espécies foram registradas como extintas até 2018 (Humphreys et al., 2019). Considerando que a taxa de extinção exata é difícil de precisar (Boucher et al., 2020), é possível que as estimativas atuais estejam subestimadas. Para que essa taxa de extinção seja contida, uma série de estratégias norteadas por princípios da conservação da biodiversidade devem ser desenvolvidas (Lindenmayer et al., 2006; Lindenmayer and Franklin, 2002). Nesse sentido, pesquisadores criaram, durante a décima reunião da Conferência das Partes (COP 10), em Nagoya (Campbell et al., 2014), a estratégia global para a conservação de plantas, com o objetivo geral de conter a perda contínua de espécies que vem ocorrendo (CBD, 2011). Dois objetivos específicos da estratégia global para a conservação de plantas (veja CBD, 2011) incluem: (I) compreender, documentar e reconhecer adequadamente a diversidade de espécies vegetais e, (II) conservar urgente e efetivamente a diversidade de espécies vegetais. Embora a ciência e as práticas de conservação tenham avançado em direção ao alcance desses objetivos (Fonseca and Venticinque, 2018; Zappi et al., 2015), para muitas regiões eles ainda estão longe de serem atingidos (Arruda et al., 2018; Corlett, 2016; Feeley, 2015; Feeley and Silman, 2011; Gomes et al., 2018; A. S. L. Rodrigues et al., 2004).

O conhecimento das espécies e suas distribuições é a base para a conservação da biodiversidade (Mace, 2004; Margules and Pressey, 2000). Não obstante, estima-se que 10-20% das espécies vegetais de angiospermas do mundo ainda não foram descritas (Joppa et al., 2010). Por exemplo, uma análise dos registro de plantas do global biodiversity information facility database (GBIF) depositados até 2013 mostrou que, cerca de 10% da América do Sul

não apresentava registros em coleções biológicas de plantas (Feeley, 2015). Essa falta de dados é observada, inclusive, na flora dos biomas mais estudados daquele continente tais como Amazônia (Ter Steege et al., 2015), Mata Atlântica (de Lima et al., 2015) e o Cerrado (Françoso et al., 2016).

Essa lacuna de conhecimento da flora se reflete na imprecisão taxonômica, ou seja, quando uma ocorrência não é identificada no nível de espécies (Caiafa and Martins, 2007; Troudet et al., 2017) e, passa a ser caracterizada nas amostras de dados como *sp*, *cf* ou *aff*. Essa lacuna de conhecimento afeta nossas conclusões baseadas nos padrões de biodiversidade em diferentes aspectos (Bini et al., 2006; Cardoso et al., 2011; Hortal et al., 2015; Whittaker et al., 2005). Se assumirmos, por exemplo, que os registros não identificados são espécies não descritas (déficit Linneano), isto é, novas espécies (Brown and Lomolino, 1998), então nossas conclusões seriam baseadas em uma fração sub-representativa da biodiversidade existente (Vale and Jenkins, 2012). Por outro lado, se assumirmos que os registros não identificados são parte da distribuição (déficit Wallaceano) de uma espécie já descrita (Brown and Lomolino, 1998), então nossas conclusões seriam baseadas em apenas uma parte da distribuição dessa espécie (Bini et al., 2006; Oliveira et al., 2016). Do ponto de vista pragmático, essa falta de conhecimento da distribuição real de uma espécie pode nos levar a classificar espécies raras como espécies com deficiência de dados (Corlett, 2016).

A riqueza das espécies raras compreende uma parte substancial das assembleias de espécies (Gaston, 2012). Além disso, as espécies raras desempenham um papel funcional na estrutura e função das assembleias, não suportado por espécies comuns (Chapman et al., 2018; Gaston, 2012; Jain et al., 2014; Leitão et al., 2016; Lyons et al., 2005; Mi et al., 2012; Mouillot et al., 2013a; Soliveres et al., 2016). Finalmente, as espécies raras são consideradas espécies focais para a conservação porque serem limitadas por área, processos de dispersão e recursos

(Lambeck, 1999, 1997), além de serem mais susceptíveis as mudanças antrópicas do que espécies comuns (Lambeck, 1997; Sætersdal and Gjerde, 2011; Santoro et al., 2012).

O número de pesquisadores que concordam que espécies raras devam ser consideradas na tomada de decisão tem crescido (Gauthier et al., 2010; Kricsfalusi and Trevisan, 2014; Longton and Hedderson, 2000; Maciel et al., 2016; McIntyre, 1992; Rey Benayas et al., 1999). De acordo com Rabinowitz (Rabinowitz, 1981), raridade é um padrão observado quando a amplitude geográfica, a especificidade ao habitat e o tamanho populacional de uma espécie são combinados. A combinação desses atributos possibilita que as espécies de uma comunidade sejam classificadas em sete formas de raridade e uma forma comum (Rabinowitz, 1981). Na prática da conservação, as espécies raras são usadas numa abordagem espacialmente explícita para avaliar e ampliar a rede de áreas protegidas (Groves et al., 2002; Margules and Pressey, 2000). Em teoria, se os locais e os habitats onde essas espécies ocorrem forem protegidos, então as espécies que lá ocorrem também serão (Arponen, 2012; Kukkala and Moilanen, 2013).

Embora os padrões de raridade em uma comunidade sejam essencialmente importantes para o planejamento da conservação (Groves et al., 2002; Yu and Dobson, 2000), os processos subjacentes que geram tais padrões não devem ser ignorados na tomada de decisão (Marshall et al., 2020). Esses processos subjacentes ou regras de montagem refletem aspectos ecológicos que regulam os padrões emergentes da comunidade (Weiher and Keddy, 2001). Esses padrões emergentes da comunidade podem ser acessados através de estratégias funcionais, por exemplo, usando uma abordagem fisiológica (McGill et al., 2006) como fenologia foliar ou capacidade de rebrota (Nakashizuka, 2001), ou uma abordagem focada na interação como síndrome de dispersão de sementes (Suding et al., 2008; Weiher et al., 1998). No contexto da conservação, essa abordagem pode fornecer uma métrica importante para medir impactos das ações antrópicas na performance dos diferentes grupos funcionais de plantas (McGill et al., 2006; Mouillot et al., 2013b). Enquanto a abordagem baseada em padrões de comunidade

podem nos ajudar no planejamento em escala regional, a abordagem baseada em processos em comunidade pode nos ajudar no manejo da conservação em escala local (Marshall et al., 2020). Assim, padrões e processos em comunidades passam a ser complementares na tomada de decisão (Callicott et al., 1999).

1.2 Savanas da América do Sul

As savanas da América do Sul (SAS) compreendem as seguintes ecoregiões: Cerrado, Llanos, Llanos de Moxos (savana de Beni), Gran Sabana, savanas do Rio Branco-Rupununi, savanas costeiras da Guiana, Llanos del Magdalena (Solbrig, 1996) e Campos rupestres (Alves and Kolbek, 2010; Antar and Sano, 2019), além de encraves de savanas na Amazônia (Barbosa and Fearnside, 2004), Caatinga (Castro and Martins, 1999), Mata Atlântica (Costa-Lima and Chagas, 2019; Moura et al., 2011) e no Chaco (Kunst et al., 2006). Ao longo dessa extensão, a SAS apresenta alta heterogeneidade de habitats (Blydenstein, 1967; Hamilton et al., 2004; Sano et al., 2019; Solbrig, 1996), que proporciona alta diversidade de plantas e elementos florísticos particulares a cada região (Andrade et al., 2018; Bridgewater et al., 2004; Ratter et al., 1996).

No que concerne a fitofisionomia, a savana pode ser caracterizada como um complexo de vegetação composto por dois estratos (Fig. 1): um lenhoso (woody layer), constituído por um componente silvestre esparsa (arbustos e árvores) imerso em um estrato graminoso contínuo (ground layer), onde predominam espécies de via fotossintética C₄ (Ratnam et al., 2011; Solbrig, 1996). Dependendo da razão entre esses estratos e a altura do estrato lenhoso, a savana pode ser classificada em savana aberta (open savanna), savana arborizada (woody savanna) ou savana florestada (savanna woodland). A savana aberta é uma fisionomia com um estrato lenhoso < 7m de altura, composto por formas arbustivo-arbóreas com cobertura do solo de 10%. A savana arborizada apresenta altura do estrato superior entre 7-14m, as árvores são a forma predominante no estrato lenhoso e a cobertura do solo pode variar entre 40% e 60%. Já

a savana florestada apresenta uma altura superior a 15m, sendo as árvores a forma predominante no estrato lenhoso, com mais de 70% de cobertura do solo (Fig. 1).

O balanço fisionômico entre savana aberta e florestada depende de muitos fatores tais como clima, fogo e solo, o que caracteriza a savana como um piropedobioma (Coutinho, 2006; Langan et al., 2017). O clima predominante nas savana é o sazonal, com seis meses secos e seis meses com precipitação variando entre 400 e 1500 mm/ano, e temperatura média anual é 24°C (Walter, 1986). Na estação seca, o estoque de água no solo é o que mantém a vegetação, sendo que a redução desse estoque limita a formação florestal e favorece a formação de savana (Lehmann et al., 2011; Murphy and Bowman, 2012). Os incêndios naturais são parte integrante da savana (Pivello, 2011), estando o maior número de incêndios relacionado a savana aberta, enquanto a supressão de incêndios favorece a formação savana florestada (Durigan et al., 2007). Solos de savana aberta são menos ricos em nutrientes do que os solos de savana florestada, resultado de uma menor produção de matéria orgânica e dos incêndios mais frequentes (Murphy and Bowman, 2012). Assim, o balanço fisionômico entre savana aberta – arbustivo – arbóreas – florestada depende de uma complexidade de fatores ambientais que atuam para determinar a fisionomia predominante (Felfili and Felfili, 2001; Furley, 1992; Langan et al., 2017; Sarmiento and Monasterio, 1975).

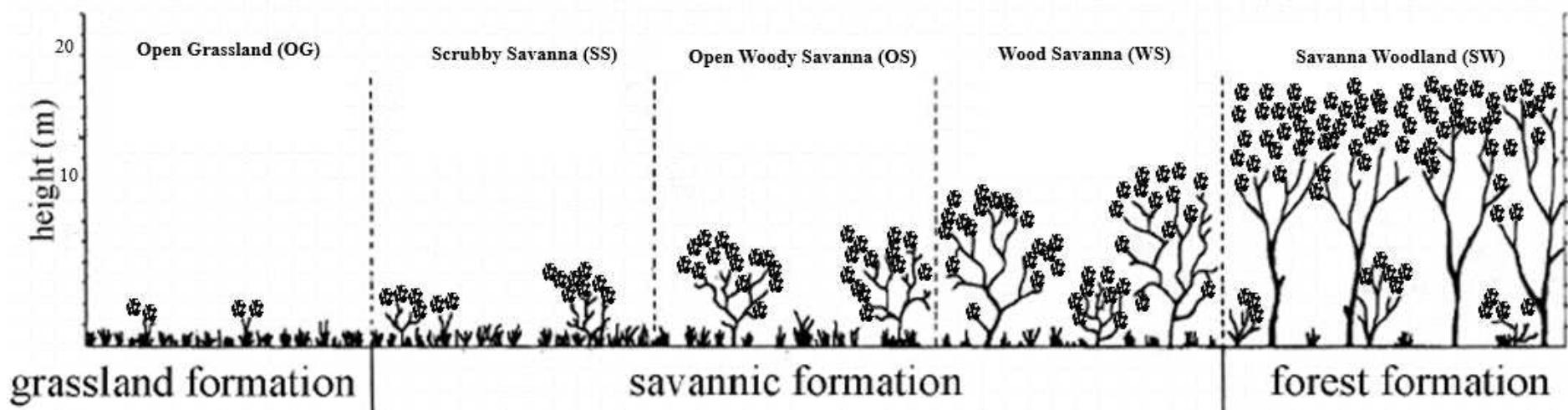


Fig. 1. Esquema de classificação da savana variando de savana aberta para savana florestada. Modificado de Coutinho 1971.

1.3 Escopo da tese

As savanas da América do Sul (SAS) apresentam grande lacuna no conhecimento da flora (Feeley, 2015; Françoso et al., 2016), uma lacuna conservação de espécies (Oliveira et al., 2017; Rodrigues et al., 2004), ao mesmo tempo em que está exposta a muitos fatores que desencadeiam a perda de espécies (Fernandes et al., 2018; Jarvis et al., 2010; Ramirez-Villegas et al., 2012) tais como mudanças climáticas (Torres and Marengo, 2014), mudança no regime de incêndios (Pivello, 2011) e defaunação (Galetti et al., 2018). No presente estudo, analisamos a SAS a luz de três questões: (1) onde se concentra a lacuna do conhecimento da flora? (2) qual o status de proteção das espécies arbóreas raras e comuns? (3) como diferentes fatores antrópicos podem afetar a performance de diferentes grupos funcionais de plantas?

Para responder as perguntas 1 e 2, usamos uma abordagem baseada em padrões em comunidades. Para isso, os registros taxonômicos não identificadas até o nível de espécie (*spp.*, *af.* e *cf.*) foram usados como um descritor da lacuna de conhecimento da flora (déficit florístico); e, as sete formas de raridade de Rabinowitz usadas para avaliar as lacunas de proteção na diversidade de árvores da SAS. Para responder à pergunta 3, usamos os processos de fenologia foliar, capacidade de rebrota e dispersão de sementes para avaliar o efeito das mudanças antrópicas na performance da vegetação arbórea da SAS. Especificamente, o efeito da mudança climática (aumento de CO₂, e da temperatura média anual e diminuição da precipitação), o aumento na frequência de incêndios e a defaunação foram usados como indicador do efeito antrópico. A permanece dos grupos funcionais de plantas foram medidas em termos de mudança na taxa de biomassa acima do solo, antes e pós distúrbio.

Aqui, nós aceitamos que o déficit florístico pode ser atribuído aos déficits Linneano e Wallaceano, conceitos propostos para descrever e quantificar as lacunas de conhecimentos na biodiversidade (Brown and Lomolino, 1998). No primeiro capítulo, fornecemos uma análise

do déficit florístico na SAS com base em estudos fitossociológicos e florísticos. Explicamos como o déficit florístico pode ofuscar nosso conhecimento da diversidade de plantas e afetar a tomada de decisão, e propomos orientações para reduzir esse déficit florístico. Para realizar essa tarefa, colocamos cinco perguntas como diretrizes: i) Os levantamentos fitossociológicos apresentam maior déficit florístico do que os levantamentos florísticos? Esperávamos que os levantamentos fitossociológicos apresentassem maior déficit florístico porque indivíduos férteis e inférteis são amostrados, enquanto em levantamentos florísticos são coletados apenas indivíduos férteis. Como flores e frutos são a base para a identificação completa da maioria das espécies, sua ausência em indivíduos inférteis pode dificultar a identificação correta. ii) Existem alguns táxons com maior déficit florístico? Esperávamos que houvesse maior déficit florístico em táxons mais diversos, como Myrtaceae e Fabaceae, do que seria encontrado em táxons menos diversos. iii) Quais as ecoregiões da SAS têm o maior déficit florístico? Considerando que, a maioria das pesquisas se concentram na área core das ecoregiões, esperávamos encontrar maior déficit florístico em áreas marginais. iv) O déficit florístico depende do total de espécies amostradas em um levantamento? Como muitas espécies permanecem desconhecidas, esperávamos que o déficit florístico aumentasse à medida que o número de espécies amostradas aumenta. v) O déficit florístico mudou ao longo do tempo? Supondo que o conhecimento da flora de uma ecorregião aumenta com o tempo de estudos realizados, esperávamos uma redução do déficit florístico ao longo do tempo.

O status de conservação de uma espécie requer a análise da representatividade da rede de áreas protegidas (APs) em conservá-la (Margules and Sarkar, 2007). Alguns exemplos demonstram que as APs da SAS é insuficiente para proteger espécies ameaçadas e endêmicas de plantas, vertebrados e invertebrados (Goetsch et al., 2019; Oliveira et al., 2017). A eficiência das APs da SAS em conservar as espécies raras e comuns (Choe et al., 2019), no entanto, permanece desconhecida. No segundo capítulo, usamos o esquema de Rabinowitz

(Rabinowitz et al., 1984) e fornecemos a primeira quantificação de espécies raras e comuns das SAS. Mais especificamente, abordamos as seguintes questões: i) quais as formas de raridade da SAS? Considerando que a SAS apresenta alta riqueza de espécies arbórea, muitas das quais com populações pequenas (Castro et al., 1999), endêmicas (Myers et al., 2000) e com alcance geográfico restrito (Bridgewater et al., 2004), esperávamos que as espécies da SAS tivessem todas as sete formas de raridade. Adicionalmente, perguntamos: 2) como essas formas de raridade são distribuídas na SAS? Esperávamos que diferentes ecorregiões tivessem números diferentes de espécies raras. Finalmente, perguntas se 3) as espécies raras e comuns estão adequadamente conservadas na rede das áreas protegidas da SAS? Como as áreas protegidas foram criadas em áreas marginais com pouco valor de conservação (Margules and Pressey, 2000), esperávamos que as áreas protegidas não proporcionassem conservação para todas as espécies raras e comuns.

A SAS é um dos biomas mais ameaçados do mundo (Myers et al., 2000). A SAS coincidem com os *hotspots* climáticos, regiões esperadas para ocorrer o maior prolongamento da estação seca da América do Sul (Torres and Marengo, 2014). Com esse aumento da seca espera-se que a SAS se tornem mais inflamáveis (Alvarado et al., 2020), o que poderá aumentar a frequência de incêndios em algumas regiões (Pivello, 2011). Adicionalmente, a SAS teve entre 50 a 80% da área nativa convertida (Jepson, 2005) para atividades como a agricultura (Alho and Martins, 1995). Esses fatores podem ter efeito direto na performance de plantas, por exemplo, o aumento na frequência de incêndios e o prolongamento da estação seca podem reduzir as taxas populacionais de espécies não adaptadas as condições extremas (Hoffmann et al., 2011; Hoffmann and Solbrig, 2003; Ojeda et al., 2005). A fragmentação e caça furtiva, por sua vez, podem diminuir as populações de dispersores de sementes (Dirzo et al., 2014), com efeito indireto na performance de espécies dispersas por animais (Herrera et al., 1994; Wright et al., 2000, 2001). Argumenta-se ainda que, esses fatores podem ter um efeito sinérgico na

performance de plantas (Brook et al., 2008), mas para a SAS isso ainda é pouco explorado. Assim, no terceiro capítulo, avaliamos o desempenho de diferentes grupos funcionais de espécies arbóreas da SAS em resposta às mudanças climáticas, mudanças na frequência de incêndio e a defaunação. Testamos as seguintes hipóteses: 1) as espécies decíduas, com capacidade de rebrota pós incêndios e não dispersas por animais são menos suscetíveis a distúrbios do que as espécies sempre-verdes, sem capacidade de rebrota e dispersas por animais; (2) as mudanças climáticas, mudanças na frequência de incêndio e defaunação atuam sinergicamente na perda da biomassa de espécies de árvores de savana.

Na medida em que os táxons e ecorregiões de maior déficit florístico da SAS forem identificados (capítulo 1), os esforços para reduzir esse déficit florístico poderão ser direcionados para táxons e regiões que apresentem maior déficit florístico. O primeiro capítulo, por tanto, contribui para conhecer e documentar a diversidade florística da SAS (objetivo I da CBD, 2011). Se detectarmos uma lacuna na rede de áreas protegidas para conservar espécies raras e comuns, então os padrões de raridade que forem identificados e mapeados aqui (capítulo 2), contribuirá para o planejamento estratégico de novas áreas protegidas. Finalmente, se o efeito da mudança climática, do regime do fogo e a dafaunação na mudança da biomassa da vegetação forem conhecidos (capítulo 3), isso contribuirá para as ações de manejo na SAS. Assim, o segundo e o terceiro capítulo contribuirão para desenvolver ações para conservação (objetivo II da CBD, 2011) da diversidade de plantas da SAS.

**CAPÍTULO 2: Linnean and Wallacean shortfalls in the South American savanna tree
flora: consequences for biodiversity conservation¹**

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Abstract

To conserve plants, we must close the knowledge gap about species, either unknown species identity (Linnean shortfalls) or unknown species distribution (Wallacean shortfalls). Because unidentified records are supposed to be a new species, we assume they as Linnean shortfall. On the other hand, it can be a species already described and difficult to identify, then this would be a Wallacean shortfall. Here, we call Linnean and Wallacean shortfalls together as flora shortfalls. We analyzed flora shortfalls based on tree phytosociological and floristic surveys to test five assumptions. Flora shortfalls: i) are greater in phytosociological than in floristic surveys; ii) differ between taxa; iii) are concentrated in some regions of the South American savannas; iv) are dependent on species richness and v) decrease over time. To test these expectations, we created a database with 355 surveys of tree species and quantified the number of species confidently identified and those not identified to binomials. An ANOVA test revealed more flora shortfalls in phytosociological than in floristic surveys. Diverse taxa such as Myrtaceae, but also less diverse taxa such as Lythraceae, had more flora shortfall. An interpolation by the inverse of the distance revealed that flora shortfalls are concentrated in the peripheral Cerrado. The greater the number of species surveyed, the higher the number of flora shortfalls. For both floristic and phytosociological surveys, flora shortfalls have not changed over time. To overcome flora shortfalls, we suggest that researchers specialized in different plant groups should engage with generalist researchers performing floristic and phytosociological surveys. Because many species not yet known may lay unidentified in some herbarium collection, the efforts in identifying herbarium exsiccata should be considered a priority.

Keywords: plant conservation, Linnean shortfall, Wallacean shortfall, flora knowledge gap, data void.

1. INTRODUCTION

Because avoiding extinction rates needs appropriate knowledge of taxonomy and species distribution (Mace, 2004; Morais et al., 2013), the knowledge gap in biodiversity became one of the greatest contemporary challenges of the conservation field (Bini et al., 2006; Brito, 2010; Feeley, 2015; Feeley and Silman, 2011; Morais et al., 2013). The knowledge gap is attributed in part to the number of species not yet described, leading to Linnean shortfalls (Lomolino, 2004; Bini et al., 2006; Brown & Lomolino, 2006; Riddle et al., 2011). The Linnean shortfalls were estimated at 10-20% in the world's angiosperms (Joppa et al., 2010). Even confidently identified plant species do not have their geographic distribution completely known (Oliveira et al., 2016; Sousa-Baena et al., 2014), leading to Wallacean shortfalls (Lomolino, 2004; Bini et al., 2006; Brown & Lomolino, 2006; Riddle et al., 2011).

Flora shortfalls affect especially tropical regions (Corlett, 2016; Feeley, 2015; Feeley and Silman, 2011). For instance, 10% of the South America continent do not have information in plant biological collections (Feeley, 2015). Lack of data is also observed in the flora of the most studied biomes of this continent, such as the Amazon (Ter Steege et al., 2015), Atlantic rain forest (de Lima et al., 2015), and the Cerrado (Françoso et al., 2016). This gap in flora knowledge is reflected in taxonomic inaccuracy, which means that when a record is not identified until species level (Caiafa and Martins, 2007; Troudet et al., 2017), then it is characterized as an indeterminated taxon, such as *sp*, *cf* ou *aff*. This taxonomic inaccuracy has already been documented for both biological collections (Troudet et al., 2017) and phytosociological surveys (Caiafa and Martins, 2007).

A comprehensive analysis of 225 surveys done in the Brazilian Atlantic rainforest revealed that among 2891 records 198 were identified only to family level and 627 to genus level (Caiafa and Martins, 2007). These authors have attributed the non-identified records to limited knowledge of the flora (Caiafa and Martins, 2007). This argument is acceptable if we

consider that the neotropical region has the richest plant biodiversity in the world and also that it is little known (Corlett, 2016; Feeley, 2015). Here, we agree with Caiafa and Martins (2007), but we go beyond them by considering that unidentified records in phytosociological or floristic surveys are Linnean and/or Wallacean shortfalls. A phytosociological survey counts, measures, and collects all individuals fulfilling certain criteria to be included in sampling unities spread across a previously selected vegetation stand, whereas a floristic survey collects reproducing individuals by patrolling a portion of the vegetation. In other words, both reproducing and vegetative individuals are collected in phytosociological surveys, but only reproducing individuals are collected in a floristic survey.

We consider that the complete identification of collected plant material to binomial is a desirable starting point in phytosociological as well as in floristic surveys. In both kinds of surveys, the researchers devote great effort to identify all records, but some remain unidentified to binomials and are included in the floristic list as genus *sp* or even family *sp* (see Caiafa & Martins, 2007). We trust that unidentified records can be Linnean shortfalls because, consequently, new species may remain unknown (Fig. 1). On the other hand, if these unidentified records are considered not new but species already described and difficult to identify, then they would be Wallacean shortfalls because the possible occurrence of the species out of their presently known geographic distribution would not be known (Fig. 1). Hence, if a record is not identified to binomial, it is difficult to distinguish whether it is a Linnean or Wallacean shortfall. Here, we consider these unidentified records as flora shortfalls, a generic designation of Linnean and/or Wallacean shortfall. As it yields an incomplete taxonomic knowledge, the flora shortfalls provide an unrealistic panorama of the spatial distribution of the biodiversity (Schaefer et al., 2011). As a consequence, our understanding of how natural and anthropic pressures influence extinction rates becomes limited (Brito, 2010). Hence, conservation strategies are limited because conservation planning for a sub-representative part

of the biodiversity may not protect that not yet known (Payo-Payo and Lobo, 2016). Then, the complete identification of species plays a key role in conservation and decision-making, since without sufficient taxonomic information, there is less chance of success (Bickford et al., 2007; Mace et al., 2008; Simó-Riudalbas et al., 2017).

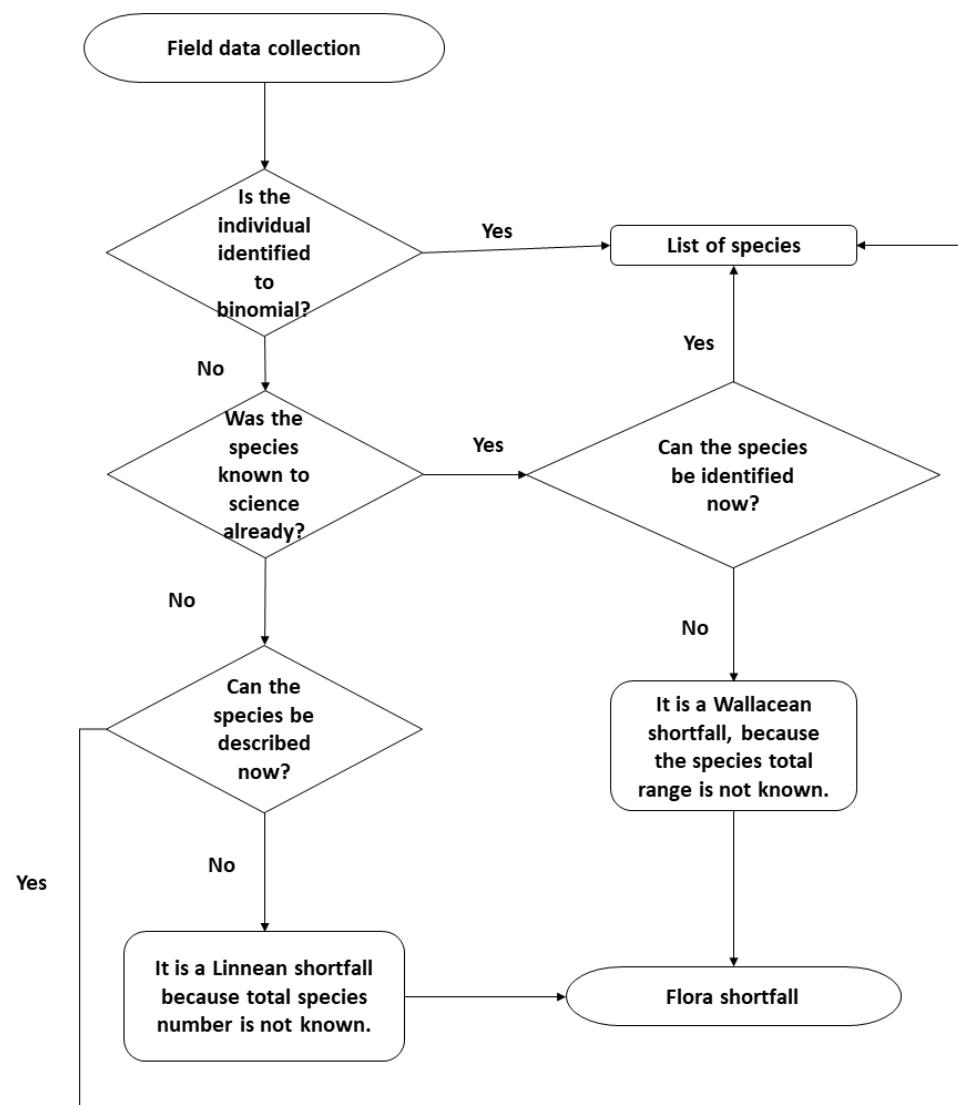


Fig. 1. Flowchart showing steps to Linnean and Wallacean shortfalls, both together called here flora shortfalls.

Here, we provided an analysis of flora shortfalls in the South American savannas based on phytosociological and floristic studies. We explain how flora shortfalls can overshadow what is known about biological diversity, thus affecting decision-making. We also propose directions to reduce flora shortfalls. To accomplish this task, we posed five questions as guidelines (see our scheme for details about our assumptions in Supplementary Fig. 1): i) Do phytosociological surveys have more flora shortfalls than floristic surveys? We expected phytosociological surveys to have more flora shortfalls because both fertile and infertile individuals are sampled and must be identified, whereas in floristic surveys only fertile individuals are collected. Since flowers and fruits are the base for complete identification of most species, their absence in infertile individuals can prevent correct identification. ii) Are there some taxa with a higher incidence of flora shortfalls? We expected more flora shortfalls to occur in more speciose taxa, such as Myrtaceae and Fabaceae. iii) What region of South American savannas has the highest number of flora shortfalls? Considering that the whole extension of the South American savannas is not uniformly sampled, we expected some regions to have more flora shortfalls than others, so that they are worthy of further attention. iv) Are flora shortfalls dependent on the total species sampled in a survey? As many species remain unknown, we expected flora shortfalls to increase as the number of species sampled increases in a survey. v) Have flora shortfalls in phytosociological and floristic surveys changed over time? We expected a reduction of flora shortfalls over time because, supposedly, the knowledge of a flora increases over time. In addressing these issues, we bore in mind that flora shortfalls, the incomplete knowledge of species identity and distribution, can negatively affect what is known about a flora and, hence, the decisions about conservation strategies.

2. METHODS

2.1 Study area

We considered savanna as a vegetation complex that includes three types of physiognomy, namely: open savanna, woody savanna, and savanna woodland (Coutinho, 1978; Furley, 1999, 1992). The open savanna includes open physiognomies with trees <7m in height and a maximum of 10% of shrub and tree crown cover. The woody savanna includes physiognomies with the predominance of trees 7-14m in height and 40% - 60% crown cover. The savanna woodland has forest physiognomies with trees over 15m in height and >70% crown cover. These physiognomies make up a complex mosaic with other vegetation types, especially seasonal forests, all over the South American savanna provinces, whose main ecoregions are the Cerrado, Llanos, Guianan savanna, and Beni savanna (Olson et al., 2001). Although not forming an ecoregion (*sensu* Olson et al., 2001), there are additionally other disjunct, isolated savanna patches scattered in non-savanna vegetation, such as the Amazon forest (northern-northwestern Brazil), Atlantic forest (roughly alongside Brazilian coast) and Caatinga (northeastern Brazil). The inclusion of these disjunct sites widens the savanna occurrence beyond the limits defined by Olson et al. (2001) for the South American Savanna (SAS; Fig. 1).

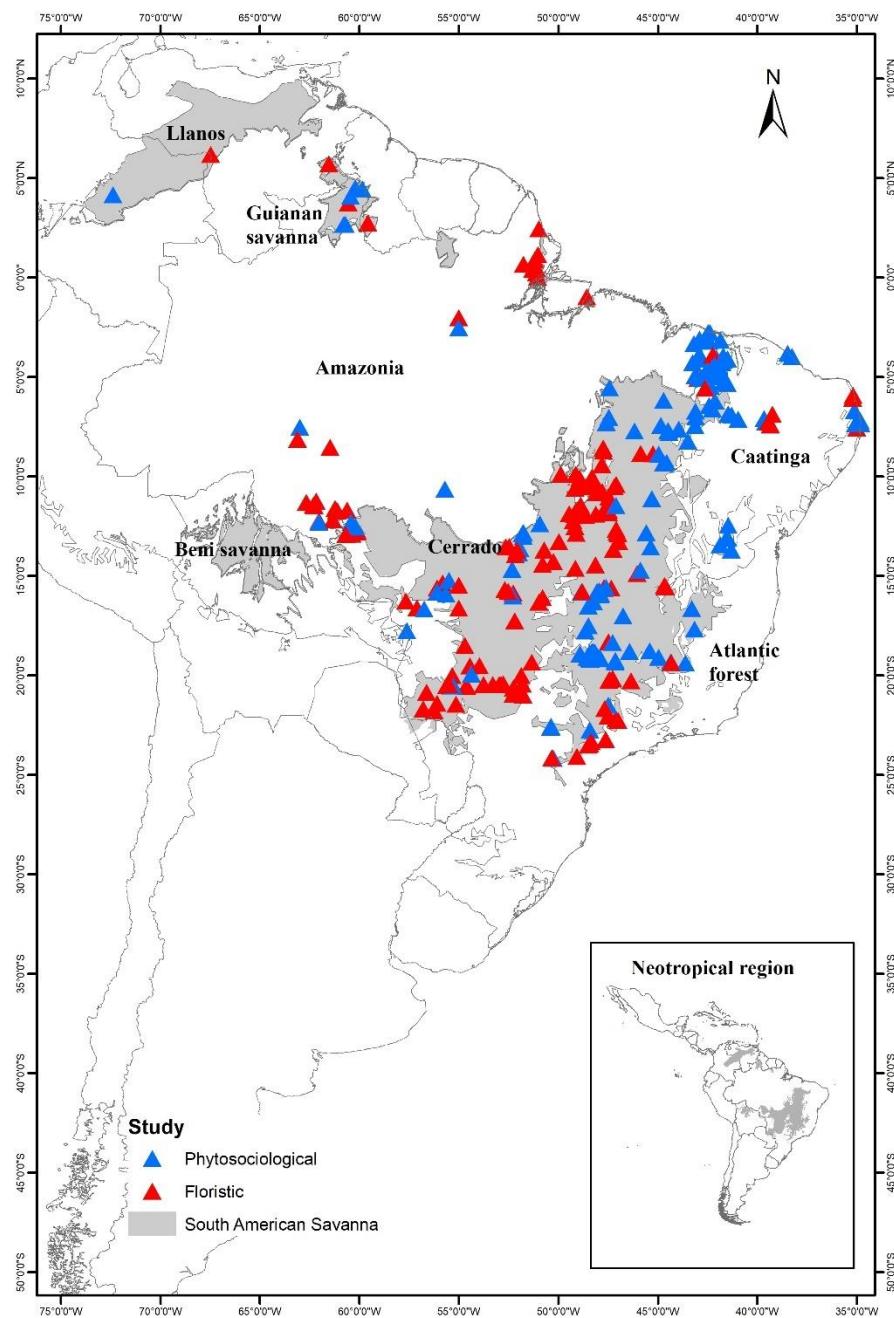


Fig. 1. The South American Savanna (grey) according to Olson et al. (2001). Floristic (red) and phytosociological (blue) surveys with flora shortfalls are scattered in the Llanos, Guianan savanna, Cerrado, and disjunct savanna areas in the Amazonia, Caatinga (northeastern Brazil), and Atlantic forest (alongside Brazilian coast). See Table S3 for literature.

2.2 Database

We searched for papers reporting floristic and phytosociological surveys in the SAS published until February 2016 in *Web of Science*, *Scopus*, *Scielo*, and *Lilacs* using as keywords different combinations of ecoregion + physiognomy + kind of surveys. For example, in searching surveys made in the Cerrado, we used the corresponding keywords ‘Cerrado’, ‘wood savanna’, ‘floristic study’ (see supplementary table 1 for more details about the keywords used). We also considered theses, scientific meeting annals, books, research notes, and unpublished data reported by *Google Scholar*. With the information gathered from this search we created a database containing: 1) the study area (we used the information provided in the papers); 2) the bibliographic reference of each paper; and 3) a list of the taxa sampled/collected in each site. The taxonomic standardization of the names of genera and families followed the *flora* package (Carvalho, 2016) in the R environment (R CoreTeam, 2019). This package directly accesses *flora 2020 online* to generate an output that contains currently accepted taxon names.

2.3 Quantifying the flora shortfalls

We assumed all occurrence of a taxon (whether or not identified to binomial) in floristic and phytosociological survey as a record. Then, we assumed as a flora shortfall any record not identified to binomial, such as unknown, Fabaceae sp., *Andira* sp., spp., aff. (*affinis*), cf. (*conferatum*), and so on (e.g. Caiafa and Martins 2007). We classified data into phytosociological and floristic surveys to allow for comparison between them. For each survey, we quantified the number of records not identified to binomial (absolute shortfalls) and identified to binomial (binomials), and we adopted the same procedure for each family and genus (e.g. Caiafa and Martins 2007). The sum of binomials + absolute shortfalls was considered as the species richness, which was expressed per site or family or genus, according to the case. With these values, we calculated the relative flora shortfall (rfs) as the ratio of the

number of absolute flora shortfalls (afs) to the species richness (S) in each survey or family or genus: $rfs = afs/S$. When analyzing the family or genus with the highest rfs, we pooled floristic and phytosociological surveys together.

2.4 Data analysis

2.4.1 Flora shortfalls in phytosociological and floristic surveys

To analyze differences of relative flora shortfalls (rfs) between the floristic and phytosociological surveys, we performed an analysis of variance (ANOVA). The normality and heterogeneity errors of the variance were tested used the Shapiro and Bartlett test, respectively. Because the rfs did not fulfill the assumptions of normality ($W = 0.725$, $p < 0.00$) and homoscedasticity (Bartlett's K-squared = 45.941, $df = 1$, $p = 0.00$), we applied the Games–Howell test, which is appropriate for unplanned comparisons and nonparametric data (Ruxton & Beauchamp, 2008). The ANOVA with Games–Howell test was applied to compare the number of rfs between floristic and phytosociological surveys (phytosociological ≠ floristic), the analyses being carried using the ‘posthocTGH fucntion’ of the ‘userfriendlyscience package’(Peters et al., 2018) in R version 3.6.1 (R Core Team, 2019). We expected to find a higher number of rfs in phytosociological than in floristic surveys.

2.4.2 Flora shortfalls among the taxa

We considered that a taxon with a high rfs poses a higher difficulty to correct binomial identification and supposed that the relative flora shortfall (rfs) differed among taxa (e.g., Fabaceae ≠ Malvaceae ≠ Myrtaceae, and so on). To test this assumption, we calculated the rfs of each genus and family. Then, we calculated the median relative flora shortfall (mrfs) for each family and genus in n floristic and phytosociological surveys pooled together. Taxa recorded in less than 5 surveys ($n < 5$) were excluded from the analysis. Additionally, we calculated the semi-amplitude of the 95% confidence interval (SCI) around the mrfs for both families and genera. The overlap of confidence intervals indicates statistically similar medians;

otherwise, the medians differ. The analysis was performed using ‘group.CI function’ of the ‘Rmisc packge’ (Hope, 2013) in R version 3.6.1 (R CoreTeam, 2019).

2.4.3 Spatial distribution of the flora shortfalls

To understand how relative flora shortfalls (rfs) are distributed across the SAS, first we built a data matrix with the geographical coordinates of the floristic and phytosociological surveys and the respective rfs. We used this matrix to create a tow data layer in a vector format (floristic and phytosociological layers). Then, we superposed this layer onto a South America map, so as to locate each floristic and phytosociological survey with their respective rfs. Finally, we applied an interpolation technique that uses known point values to estimate values of points not yet known. We compared four different interpolation techniques: idw, spline, natural neighbor, and kriging. To choose the best of them, we adjusted a regression model between the known rfs and the modeled rfs provided by each technique. Hence, we chose the inverse distance weighted (IDW) technique (Watson & Philip, 1985) because it produced the best fit between known data versus modeled data (Table S2). For this procedure, we used the ArcGis (ESRI, 2011) and the World Geodetic System 1984 (WGS 84) as reference systems. As a result of this analysis, we mapped the rfs in the space and identified gaps of knowledge in the SAS tree flora. This helped us to identify the overall pattern of rfs within the SAS.

2.4.4 The relationship between species richness and flora shortfalls

To analyze the relationship between the number of absolute flora shortfalls (afs) and the species richness (S), we quantified them in each phytosociological and floristic survey separately. We used standardized major axis (SMA) regression to evaluate the relationship between the number of afs (y-axis) and the species richness (x-axis). We used the SMA because it is indicated to define a line of best fit for a bivariate relationship (Smith, 2009). To calculate the intercept, slope, 95% confidence interval (CI), and r-square we used *lmodel2* package (Legendre & Oksanen, 2018) in R version 3.6.1 (R CoreTeam, 2019). Because many different

surveys may have the same number of afs, the result is susceptible to recall bias. To correct this bias we calculated the permutation number for repeated elements (Antoch and Hušková, 2001). This procedure helped us to conclude whether or not the observed pattern has arisen by chance even if the dataset was not representative of a statistical population (Legendre and Legendre, 2012). To analyze how rfs varied over time, we used the intercept (β_0) and slope (β_1) of the best line. We first tested if β_1 was different from zero for floristic ($\beta_{1\text{flor}}$) and phytosociological surveys ($\beta_{1\text{phyto}}$). The $\beta_1 = 0$ would indicate that afs did not change with S, while the $\beta_1 \neq 0$ would indicate an increase ($\beta_1 > 0$) or decrease ($\beta_1 < 0$) in it. We expected the number of afs to increase as the number of records increased, hence a positive ($\beta_1 > 0$) relationship between afs and S. We used a t-test to compare the intercepts ($\beta_{0\text{flor}} vs. \beta_{0\text{phyto}}$) and slope ($\beta_{1\text{flor}} vs. \beta_{1\text{phyto}}$) between the two kinds of surveys. This helped us to understand whether afs variation differed between floristic and phytosociological surveys with the increase of species surveyed (S). To analyze how much of the afs variation was explained by S variation, we used the r square value. As our assumption was that the greater the S the higher the number of afs, a high value of r was expected.

2.4.5 Temporal variation in flora shortfalls

To investigate whether relative flora shortfalls (rfs) varied over time, we considered rfs as the dependent variable and the number of years (yr) of study as the independent factor ($rfs = a + b \text{ yr.}$). We followed Warton's approach to choosing the best goodness of fit model (Warton et al., 2016). As rfs is a proportion, we performed two general linear models (GLM). For the first model, we used rfs without transformation and carried out a GLM with quasi-Poisson distribution and logit link. For the second model, we applied a logit transformation on the rfs (logit rfs) and used logit rfs to adjust a GLM with Gaussian distribution. We took the first model because it produced a better fit model-predicted versus adjusted residuals. In addition, it has the advantage to preserve the data on the same observation scale without transforming

them (St-Pierre et al., 2018)) in R version 3.6.1 (R CoreTeam, 2019). All models were performed by *GLM* function (Wickham, 2009), with $p < 0.05$ as the significant threshold. To analyze how rfs varied over time, we used the intercept (β_0) and slope (β_1) of the best line. We first tested if β_1 was different from zero for floristic ($\beta_{1\text{flor}}$) and phytosociological surveys ($\beta_{1\text{phyto}}$). The $\beta_1 = 0$ would indicate that the rfs has not changed over time while the $\beta_1 \neq 0$ would indicate an increase ($\beta_1 > 0$) or decrease ($\beta_1 < 0$) in it. Because we supposed that a flora becomes better known as it is studied along time, we expected the rfs to decrease over time ($\beta_1 < 0$). We used a t-test to compare the intercepts ($\beta_{0\text{flor}}$ vs. $\beta_{0\text{phyto}}$) and slope ($\beta_{1\text{flor}}$ vs. $\beta_{1\text{phyto}}$) between the two kinds of surveys. This helped us to understand whether rfs variation differed between floristic and phytosociological surveys over time.

3. RESULTS

We found a total of 114 papers with flora shortfalls (Table S3) corresponding to 355 surveys, of which 161 floristic and 194 phytosociological surveys (Table S4). Of these 114 papers, 57.8% were published articles, 15.78% thesis, 13.5% unpublished data, 7.8% technical bulletins, technical reports, or research notes, and 5.3% were books (Table S4). The total number of records in all papers was 20,968, of which 18,671 binomials and 2,297 records not identified to binomials, yielding a relative flora shortfall of 10.9%. Among the 2,297 absolute flora shortfalls, 631 (27.48%) were in floristic surveys, and 1,666 (72.52%) were in phytosociological surveys (supplementary table 3). In the 114 papers, we found 119 families and 637 genera, of which 73 families and 289 genera had flora shortfalls.

3.1 Phytosociological surveys have more flora shortfalls

The mean relative flora shortfall (rfs) was 0.07 ($SD \pm 0.08$) and 0.16 ($SD \pm 0.13$) in the floristic and phytosociological surveys, respectively. The ANOVA with Games-Howell test

revealed that the mean rfs was significantly higher in phytosociological than floristic surveys ($F = 48.93$, $p < 0.001$).

3.2 Some taxa have more relative flora shortfalls

Of the 119 families, 73 had flora shortfalls and 32 of these were present in 5 or more surveys (5 was considered the minimum sample size based on 355 surveys and 114 papers). The median relative flora shortfall (mrfs) in these 32 families ranged from mrfs=0.12 (SCI \pm 0.01) to 0.50 (SCI \pm 0.07) (Table 1), meaning that these families could present 12% to 50% of records without confident identification to genus or binomials in some surveys. Less frequent families among savanna trees, such as Verbenaceae (mrfs = 0.50 \pm 0.07), Asteraceae (0.42 \pm 0.05), and Sapindaceae (0.42 \pm 0.07), had higher mrfs than the most frequent families, such as Myrtaceae (0.33 \pm 0.03), Malpighiaceae (0.25 \pm 0.03) and Fabaceae (0.12 \pm 0.01).

Table 1. Families with flora shortfalls occurring in five or more surveys. The median relative shortfalls (mrfs) and the respective semi-amplitude of the 95% confidence interval (SCI) in the number (n) of surveys where they appear are shown. A total of 355 surveys and 114 references were analyzed.

| Family | <i>n</i> | mrfs | SCI |
|------------------|----------|------|------------|
| Myrtaceae | 63 | 0.33 | \pm 0.03 |
| Fabaceae | 58 | 0.12 | \pm 0.01 |
| Malpighiaceae | 45 | 0.25 | \pm 0.03 |
| Rubiaceae | 31 | 0.23 | \pm 0.03 |
| Bignoniaceae | 26 | 0.20 | \pm 0.03 |
| Apocynaceae | 24 | 0.31 | \pm 0.03 |
| Annonaceae | 21 | 0.33 | \pm 0.04 |
| Asteraceae | 20 | 0.42 | \pm 0.05 |
| Melastomataceae | 18 | 0.25 | \pm 0.05 |
| Sapotaceae | 17 | 0.33 | \pm 0.06 |
| Malvaceae | 16 | 0.34 | \pm 0.05 |
| Arecaceae | 14 | 0.38 | \pm 0.31 |
| Euphorbiaceae | 14 | 0.38 | \pm 0.05 |
| Ochnaceae | 14 | 0.33 | \pm 0.06 |
| Chrysobalanaceae | 13 | 0.40 | \pm 0.06 |
| Erythroxylaceae | 13 | 0.33 | \pm 0.05 |

| Family | n | mrf_s | SIC |
|----------------|----------|------------------------|------------|
| Salicaceae | 12 | 0.37 | ± 0.06 |
| Combretaceae | 11 | 0.16 | ± 0.06 |
| Lauraceae | 11 | 0.33 | ± 0.04 |
| Verbenaceae | 9 | 0.50 | ± 0.07 |
| Vochysiaceae | 9 | 0.27 | ± 0.06 |
| Anacardiaceae | 8 | 0.29 | ± 0.1 |
| Calophyllaceae | 8 | 0.31 | ± 0.09 |
| Celastraceae | 8 | 0.29 | ± 0.1 |
| Connaraceae | 8 | 0.42 | ± 0.07 |
| Ebenaceae | 8 | 0.23 | ± 0.11 |
| Lamiaceae | 8 | 0.25 | ± 0.1 |
| Sapindaceae | 8 | 0.42 | ± 0.08 |
| Moraceae | 7 | 0.33 | ± 0.07 |
| Nyctaginaceae | 7 | 0.25 | ± 0.05 |
| Polygalaceae | 5 | 0.40 | ± 0.05 |
| Solanaceae | 5 | 0.33 | ± 0.07 |

Of the 637 genera, 289 had flora shortfalls, and 74 of these were present in 5 or more surveys. The mrf_s in these 74 genera ranged from 0.17 ± 0.36 to 1 ± 0.21 (Table 2). In other words, these genera could present 17% to 100% of their records without confident identification to binomials in some surveys. Only 9 genera had mrf_s < 1: *Brosimum*, *Byrsonima*, *Campomanesia*, *Hymenaea*, *Machaerium*, *Pseudobombax*, *Psychotria*, *Senna*, and *Symplocos*. All the other genera had a mrf_s flora shortfall of 1, such as *Casearia*, *Dalbergia*, and *Duguetia*.

Table 2. Genera with flora shortfalls occurring in five or more surveys. The median relative flora shortfalls (mrf_s) and respective semi-amplitude of the confidence interval (SCI) in the number (n) of surveys where they appear are shown. A total of 355 surveys and 114 references were analyzed.

| Genera | n | mrf_s | SIC |
|---------------|----------|------------------------|------------|
| Myrcia | 75 | 1 ± 0.07 | |
| Eugenia | 57 | 1 ± 0.07 | |
| Byrsonima | 54 | 0.5 ± 0.09 | |
| Psidium | 50 | 1 ± 0.05 | |
| Erythroxylum | 47 | 1 ± 0.08 | |
| Miconia | 46 | 1 ± 0.08 | |
| Andira | 30 | 1 ± 0.06 | |

| Genera | n | mrf_s | SIC |
|---------------|----------|------------------------|------------|
| Pouteria | 29 | 1 ± | 0.06 |
| Bauhinia | 26 | 1 ± | 0.05 |
| Kielmeyera | 25 | 1 ± | 0.14 |
| Callisthene | 22 | 1 ± | 0.05 |
| Ouratea | 22 | 1 ± | 0.08 |
| Himatanthus | 21 | 1 ± | 0.05 |
| Ocotea | 21 | 1 ± | 0.14 |
| Tabebuia | 21 | 1 ± | 0.08 |
| Casearia | 19 | 1 ± | 0 |
| Hymenaea | 19 | 0.5 ± | 0.06 |
| Diospyros | 18 | 1 ± | 0.06 |
| Manihot | 17 | 1 ± | 0 |
| Aspidosperma | 16 | 1 ± | 0.11 |
| Guapira | 15 | 1 ± | 0.11 |
| Campomanesia | 14 | 0.75 ± | 0.08 |
| Croton | 14 | 1 ± | 0.1 |
| Salacia | 14 | 1 ± | 0.1 |
| Duguetia | 13 | 1 ± | 0 |
| Licania | 11 | 1 ± | 0 |
| Vitex | 11 | 1 ± | 0 |
| Heteropterys | 10 | 1 ± | 0.15 |
| Lippia | 10 | 1 ± | 0 |
| Schefflera | 10 | 1 ± | 0.15 |
| Syagrus | 10 | 1 ± | 0 |
| Vernonia | 10 | 1 ± | 0.11 |
| Annona | 9 | 1 ± | 0.13 |
| Cordia | 9 | 1 ± | 0.13 |
| Dalbergia | 9 | 1 ± | 0 |
| Lantana | 9 | 1 ± | 0 |
| Vismia | 9 | 1 ± | 0.06 |
| Alibertia | 8 | 1 ± | 0.15 |
| Eriotheca | 8 | 1 ± | 0.2 |
| Jacaranda | 8 | 1 ± | 0 |
| Myrsine | 8 | 1 ± | 0.15 |
| Cassia | 7 | 1 ± | 0 |
| Combretum | 7 | 1 ± | 0 |
| Copaifera | 7 | 1 ± | 0 |
| Eremanthus | 7 | 1 ± | 0.17 |
| Inga | 7 | 1 ± | 0 |
| Pseudobombax | 7 | 0.75 ± | 0.27 |
| Baccharis | 6 | 1 ± | 0.21 |
| Brosimum | 6 | 0.5 ± | 0.21 |
| Chrysophyllum | 6 | 1 ± | 0 |
| Eupatorium | 6 | 1 ± | 0 |
| Guatteria | 6 | 1 ± | 0.21 |

| Genera | n | mrf_s | SIC |
|-----------------|----------|------------------------|------------|
| Matayba | 6 | 1 ± | 0 |
| Senna | 6 | 0.17 ± | 0.36 |
| Solanum | 6 | 1 ± | 0 |
| Vochysia | 6 | 1 ± | 0.36 |
| Coccoloba | 5 | 1 ± | 0.28 |
| Connarus | 5 | 1 ± | 0 |
| Davilla | 5 | 1 ± | 0 |
| Ficus | 5 | 1 ± | 0 |
| Ilex | 5 | 1 ± | 0 |
| Machaerium | 5 | 0.67 ± | 0.37 |
| Maytenus | 5 | 1 ± | 0 |
| Psychotria | 5 | 0.4 ± | 0.48 |
| Sapium | 5 | 1 ± | 0 |
| Stryphnodendron | 5 | 1 ± | 0.28 |
| Symplocos | 5 | 0.5 ± | 0.39 |
| Tapirira | 5 | 1 ± | 0 |
| Vatarea | 5 | 1 ± | 0 |

3.3 The flora shortfalls concentrated in some ecoregions

Both phytosociological and floristic surveys showed flora shortfalls distributed throughout the SAS but concentrated in some ecoregions. The relative flora shortfall (rfs) ranged from 0.011 to 0.38 in floristic surveys and from 0.009 to 0.541 in phytosociological surveys. The highest rfs (> 0.5) in floristic surveys were found in the Llanos, Cerrado-Caatinga transition, the southeastern sector of the Cerrado, and savanna disjunctions (Amazon and Atlantic forest) (Fig. 3a). In phytosociological surveys, the highest rfs were found in the Cerrado-Amazon and Cerrado-Caatinga transitions, and Beni savanna (Fig. 3b). On the other hand, more than 50% of the total number of surveys have been done in the Cerrado (Table S5), although a great number of surveys have also been done in other ecoregions, such as the Guianan savanna and the disjunct savannas in the Maranhão Babaçu forest, Madeira-Tapajós moist forest, and Caatinga.

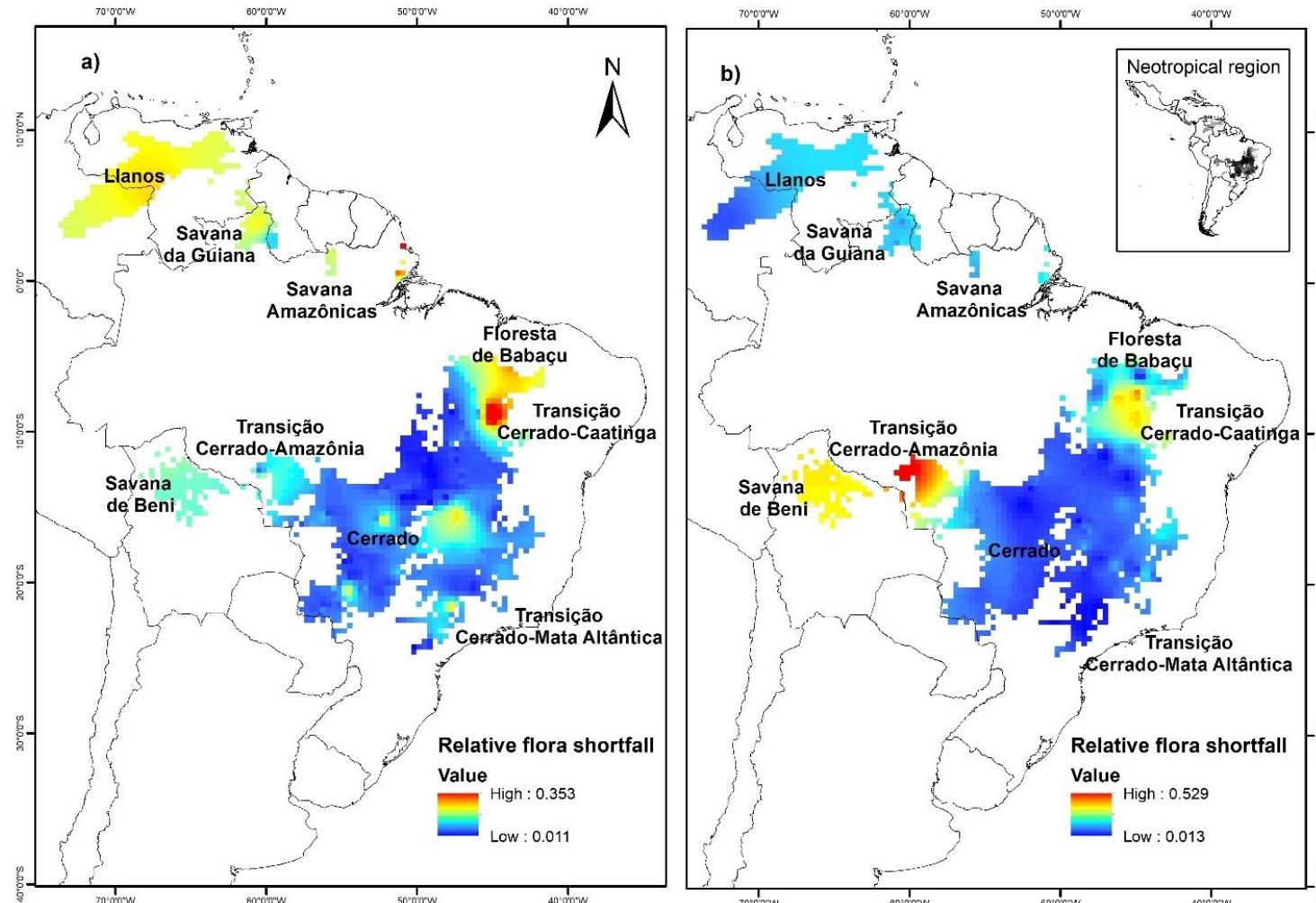


Fig. 3. Interpolation map by the inverse distance weighted (IDW) technique for floristic (a) and phytosociological (b) surveys in the South American savannas (adapted from Olson et al. 2001) including the Llanos, Guianan savanna, Cerrado, and disjunct savanna areas in Amazonia, Caatinga (northeastern Brazil), and Atlantic forest (alongside Brazilian coast). The relative flora shortfall increases from blue to red color (highest relative flora shortfall).

3.4 The number of flora shortfalls is weakly related to the species richness

The SMA slope showed a positive correlation between the number of absolute flora shortfalls (afs) and species richness (S) as can be seen on the β_1 different from zero for both floristic ($\beta_{1\text{flor}} = 0.18$; $p = 0.006$) and phytosociological surveys ($\beta_{1\text{phyto}} = 0.34$; $p = 0.009$). T-tests found no significant differences in the slopes ($t = 3.25$, $df = 1$, $p\text{-value} = 0.19$) or the intercepts ($t = -5.6798$, $df = 1$, $p\text{-value} = 0.1109$) between the floristic and phytosociological surveys. According to the goodness-of-fit test, for both floristic ($R^2=0.04$) and phytosociological ($R^2=0.12$) surveys, the proportion of the variation in afs explained by S was quite small. In the floristic surveys, only 0.5% of the afs could be attributed to the increasing richness, while this number increases to 12% in phytosociological surveys. The results of the regression analysis are set out in figure 4.

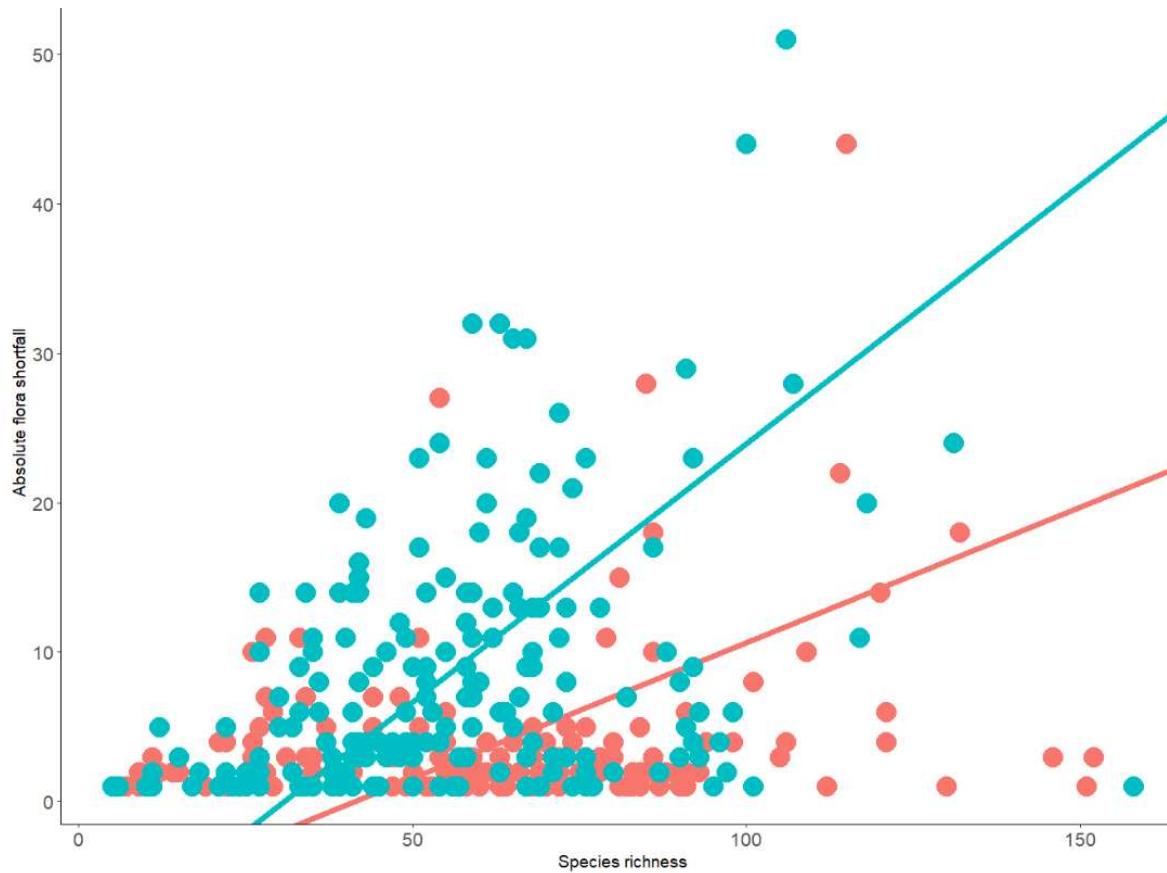


Fig. 4. Relationship between absolute flora shortfall and the species richness of floristic (red) and phytosociological (blue) surveys. The lines represent the best-fit regression from the Standardized Major Axis (SMA) for floristic (red) and phytosociological (blue) surveys.

3.5 Flora shortfall over time

Our data comprised almost 4 decades of surveys between 1980 to 2015, and we expected the flora shortfalls to diminish with time, as the flora became progressively better known. However, the slope (β_1) of the regression line was not different from zero for both floristic ($\beta_{1\text{flor}} = -0.02$; $p = 0.08$) and phytosociological surveys ($\beta_{1\text{phyto}} = 0.01$; $p = 0.3$). This means that there was no relation of the rfs variation with time, whether considering floristic or phytosociological surveys. Also, T-tests found no significant differences in the slopes ($t = 1.73$, $df = 3$, $p = 0.19$) or intercepts ($t = 0.23344$, $df = 1$, $p\text{-value} = 0.854$) of floristic or phytosociological surveys. These results show that the proportion of rfs in both floristic and phytosociological surveys kept a similar magnitude and did not diminish along time. The results of the Generalized Linear Model analysis are set out in figure 5.

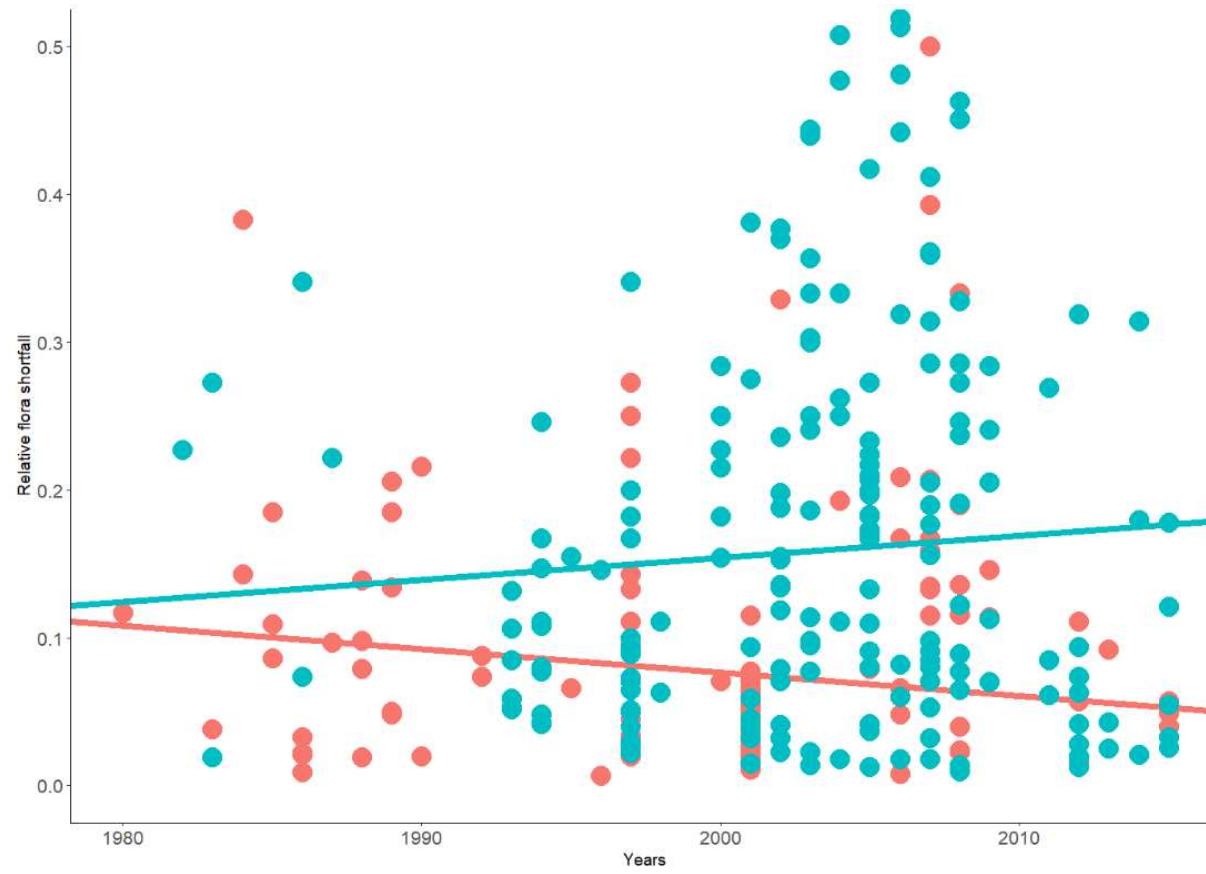


Fig. 5. Dispersion model of the relative flora shortfalls over time in floristic (red) and phytosociological (blue) surveys. The lines represent the best-fit regression from the Generalized Linear Model (GLM) with quasi-Poisson logit link for floristic (red) and phytosociological (blue) surveys.

4. DISCUSSION

Several reports have shown that Linnean and Wallacean shortfalls represent a big challenge for conservation biodiversity, but none of them investigated these shortfalls in the South America savanna (SAS). Here, we provide the first analysis using the records unidentified until binomial as a descriptor of the Linnean and Walleacean shortfalls of the SAS tree flora. Of the 20,968 records analyzed, 2,297 were flora shortfalls, which occurred mainly in some taxa and regions of the SAS.

4.1 Phytosociological surveys have more flora shortfalls

We found more flora shortfalls in phytosociological than floristic surveys. In the phytosociological surveys, all individuals fulfilling previously agreed sampling criteria are collected, independently of their phenophase. On the other hand, only individuals with fruits and/or flowers are collected in floristic surveys. Depending on the family or genus, the identification of some species is very difficult, even when they bear flowers and fruits; and when they are in a vegetative phenophase, the identification to binomial is at least harder, if possible. Traditionally, flowers and fruits are the basis for the taxonomic description of species and genera (Rejmánek and Brewer, 2001). In his book *Philosophia botanica*, Linneus (1975) declared that '*habitus occulte consulendus est*', i.e., plant habitus (non-reproductive structures) are surreptitiously analyzed, so that they would not be part of the basis for separating genera and species (Cain, 1995).

Even today, non-reproductive structures are still seldom considered in identification keys of traditional taxonomy, and most identification keys consider only reproductive attributes (Rejmánek and Brewer, 2001). To reduce flora shortfalls, we argue that identification keys and the respective species description should provide more detailed characters and consider both reproductive and vegetative attributes (Costion and Plunkett, 2016; Mantovani et al., 1985; Urbanetz et al., 2010). For example, Costion and Plunkett (2016) proposed an identification

key for the genus *Osmoxylon* (Araliaceae) that uses both vegetative and reproductive characters.

4.2 Some taxa have higher proportions of relative flora shortfall

Some savanna taxa have wide morphological variations. For example, a study comparing individual variations of *Myrcia splendens* (Sw.) DC. (Myrtaceae) from different populations revealed differences in many attributes, such as leaf area, length, thickness, and width (Amorim and de Melo Júnior, 2016). This makes the process of delimiting species very difficult even for specialists (Barroso, 1984; McVaugh, 1968). Due to these morphological variations, there are cases in which a unique species was considered to be various. On the other hand, some taxa are very similar and lack of variation can lead to more than one species being classified as a unique, a well-known problem in savanna flora (e.g. Goodland and Ferri, 1979). Due to morphological similarity, about 10-20% of the species that are identified by traditional taxonomy are supposedly to be two or more (Janzen et al., 2017).

We observed that two kinds of families have higher proportions of flora shortfalls. On one side, the most speciose families in the savanna tree flora, such as Fabaceae, Myrtaceae, Vochysiaceae, Malpigiaceae, Bignoniaceae, Apocynaceae, and Rubiaceae (Castro et al., 1999; Françoso et al., 2016) have many records without confident identification to binomials. On the other side, families with few species in the savanna tree flora also show many shortfalls, as is the case of Chrysobalanaceae, Ebenaceae, and Ochnaceae. There is in the research literature a historic tendency to focus efforts on the more diverse taxa while the less diverse taxa have been neglected (Di Marco et al., 2015). Many authors have focused discussion on the most diverse families such as Fabaceae and Myrtaceae (e.g. Dutra et al., 2015; Zappi et al., 2015; Terra et al., 2017), while other smaller families are seldom discussed. Our results show that both large and small families pose similar problems in identifying species. In both floristic and phytosociological surveys, the most speciose families are sampled with many species, not

all of them known by the researcher, although these large families are the most studied in taxonomic investigations. The consequence is that many records of these families are not confidently identified to binomials because very few specialists know every species of a very large family (Hopkins, 2005). However, there are families with few species in the savanna tree flora, and because the small taxa usually do not attract much taxonomic attention (Di Marco et al., 2015), they are seldom studied and their species are generally not completely known.

The use of vegetative characters as additional information in the plant keys is just one of the necessary actions because diminishing the taxonomic uncertainty will only be possible with the application of modern techniques together with traditional ones (Draper et al., 2020). These modern techniques may include, for example, DNA sequencing and image recognition (Draper et al., 2020). DNA sequencing is a powerful technique for the discrimination of cryptic species (Prata et al., 2018; Vicentini, 2016), i.e., two morphologically similar species. A recent example using Fourier Transformed Near-Infrared Spectroscopy for taxonomic identification from tree branches proved to be efficient in identifying genus and family (Lang et al., 2017).

4.3 Flora shortfalls are concentrated in some regions

The pattern that we observed in the spatial analysis of the flora shortfalls in the SAS showed that the Brazilian Cerrado is the most problematic savanna in South America since it had the greatest number of flora shortfalls. On the one hand, the Brazilian Cerrado is the richest savanna in the world, especially in South America (Pennington and Lavin, 2016), meaning that it can still have many unknown species. On the other hand, the Cerrado has also been the most surveyed savanna, and few surveys have been done in other savanna ecoregions. Hence, the smaller proportions of flora shortfalls in non-Cerrado savannas in South America may be the outcome of a small number of surveys. The small number or lack of surveys in non-Cerrado South American savannas has been reported as a great problem in the Neotropics, leading to knowledge gaps of the flora of entire regions (e.g. Kier et al., 2005). These Neotropical regions

with gaps in the flora knowledge have great potential for the discovery of new species (Bini et al., 2006; Brito, 2010). Then, both Cerrado and non-cerrado savannas still require more surveys and special attention to their flora.

Our results also show that the Cerrado-Amazonia and Cerrado-Caatinga transition zones concentrate many flora shortfalls. The savanna transition zones in South America have been reported as regions representing great gaps of knowledge on the flora (Castro et al., 1999; Françoso et al., 2016), and at the same time they are regions for the potential appearing of new species (Schmida and Wilson, 1985; Castro et al., 1999). The Cerrado transition zones are generally far from the research centers that traditionally make researches on the Cerrado flora, which are located in regions with great population density. A spatially explicit analysis by Diniz-Filho et al. (2005) showed that the number of described anuran species in the Cerrado positively correlates with the size of the regional human population. According to the model proposed by Diniz-Filho et al. (2005), biodiversity knowledge increases in response to the population size and knowledge of a region. In the case of the Cerrado, its occupation has begun in the south heading towards the north (Klink and Moreira, 2002) and may partially explain the spatial pattern we found for the flora shortfalls. We found that most flora shortfalls concentrated in the northern portions of the Cerrado, that is, Cerrado-Amazonia and Cerrado-Caatinga transitions, indicating a delay in the knowledge of the flora in these regions in the relation to the southern portions of the Cerrado. Therefore, intensifying floristic and phytosociological surveys in the South American savanna transition zones is recommended (Haidar et al., 2013; Maciel et al., 2016; Smith et al., 1997).

Nevertheless, the many tree flora shortfalls observed in the savanna transition zones may be a consequence of phenotypic variation in response to the variable environmental conditions. Plant populations may show great variability among individuals in response to multiple, different factors that are frequent in heterogeneous environments, such as transition

zones (Menezes et al., 2017; Slovák et al., 2012; Valladares et al., 2007). Therefore, individuals in transition zones may show great variation in attributes that largely differ from those of individuals of the same species in a core area (Smith et al., 2001). We believe that at least part of the flora shortfalls we observed in transition zones comes out from the peculiar environmental variation among savannas.

4.4 The number of flora shortfalls is weakly related to the species richness

Given the fact that many species are still unknown in the world (Linnean shortfalls), we expected a strong correlation between richness and flora shortfall. We expected that as species richness in a survey increased, the chance for new species to appear also increased. Indeed, we found that the number of flora shortfalls increased as the number of surveyed species increased. However, this correlation was weak, meaning that only a small part of the flora shortfalls could be attributed to an increasing number of records surveyed. Therefore, we conclude that most of the flora shortfalls result from hardly identifiable, widespread species. In consequence, the geographic distribution of many taxa (Wallacean shortfalls) in the South American savannas remains unknown.

Although our results suggest that the flora shortfalls are more likely to be Wallacean than Linnaean, they are still a problem. In fact, if these biodiversity data are a sample of already known species, this suggests a bias in the distribution of existing biodiversity (Oliveira et al., 2016; Sousa-Baena et al., 2014). The main methods for the analysis of the threat status of a species are based on its distribution (Mace et al., 2008). The incomplete knowledge of the spatial distribution of taxa affects the programs of biodiversity monitoring (Pereira and Cooper, 2006). Likewise, the analysis of a species conservation status can only be done through the analysis of its distribution relative to the network of protected areas (Margules and Pressey, 2000; A. S. L. Rodrigues et al., 2004). Thus, the Wallacean shortfalls have serious implications for flora conservation.

4.5 Flora shortfalls over time

We admitted that along time there would be at least some advance in the knowledge of a flora, so we expected a decrease of the flora shortfalls over time. However, our expectation was not confirmed for both floristic and phytosociological surveys. The slopes of the GLM line of both floristic and phytosociological surveys were not significantly different from zero, meaning an absence of a tendency between flora shortfalls and time. Also, the intercepts did not significantly differ between the two regression lines, meaning that in both floristic and phytosociological surveys a similar proportion of non-identified taxa held over time. This means that the non-identified records of already surveyed areas keep on being flora shortfalls, and new shortfalls appear as new areas are sampled. The result is the maintenance of a proportion of non-identified records along time, thus perennating Linnean and Wallacean shortfalls.

Traditionally, the botanical material sampled in floristic and phytosociological surveys becomes part of regional herbarium collections. Eventually, some unidentified material can be identified, and data become available for all people around the world through platforms such as the Global Biodiversity Information Facility (GBIF). These data are the opportunity to answer questions on a large geographical and temporal scale (Troudet et al., 2017). The complete identification of the species list from a region made at a specific time is the opportunity to analyze how species richness changes in response to human actions when compared within a given time interval (Steinbauer et al., 2018). The accurate recording of a species over time underlies the many issues that emerge from the global change scenario. Thus, we believe that researchers will have to strive for this deficit to decrease in the coming decades.

5. CONCLUSION

Our results showed more frequent flora shortfalls in phytosociological than floristic surveys. Both very speciose and poor families showed high numbers of flora shortfalls, confirming that taxonomic difficulties vary among taxa. Cerrado-Amazonia and Cerrado-Caatinga transition zones had the highest numbers of flora shortfalls, demanding greater attention. These results show that the proportion of relative flora shortfalls in both floristic and phytosociological surveys kept a similar magnitude and did not diminish along time. Our findings suggest several courses of action to scale down flora shortfalls. (1) Specialists acquainted with different plant groups should integrate with specialists who carry out phytosociological and floristic surveys. (2) Because many species not yet known may lay unidentified in some collection, concentrating efforts in identifying herbarium exsiccata should be considered a priority. (3) It is do not possible to know all species in the world, but efforts should be made towards identifying as many binomials as possible.

CAPÍTULO 3: Rarity patterns in the tree flora of the South American savannas: are the protected areas enough?

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Abstract

The interest in quantifying rare species has been increasing recently, but less attention has been paid to the analysis of their conservation status. Here, we used the Rabinowitz' method, based on geographical range, habitat specificity and population size, to classify the rarity forms of tree species in the South America savanna (SAS) and to analyse their conservation status. We sought to answer the following questions: (1) What forms of rarity does the SAS tree flora have? (2) How are these forms of rarity distributed over the SAS and (3) especially in the protected savanna areas (PAs)? We considered species with narrow (stenotopic) or wide (eurytopic) geographic range occurring respectively in up to 10% or more of the latitudinal belts in the SAS, generalist (euryoecious) or specialist (stenoecious) species occurring in one or more ecoregions, respectively, and locally scarce or abundant populations with respectively low or high numbers of the records within a grid cell of 1° latitude \times 1° longitude. These parameters were combined to classify 2,203 SAS species into a common and seven forms of rarity and to analyse these species according to three categories: those occurring only within PAs, within and outside PAs or only outside PAs. Of the 2,203 tree species, 898 were rare, 826 stenotopic, 528 stenoecious and 617 locally scarce. The Cerrado had the highest number of rare species, followed by savannas in the Dry Chaco, Bahia coastal forests, Campos Rupestres, Guianan savanna and Llanos. Of the 2,203 species, 205 species occurred only in PAs, 1,650 within and outside PAs and 348 only outside PAs. Our results reveal a sampling bias on the species records outside PAs relative to those inside PAs. Future studies in PAs could reveal a conservation status of species not yet known.

Keywords: seven forms of rarity, conservation gap, protected areas, ecoregion, priority species.

1. Introduction

The identification of priority species should underly any strategy for biodiversity conservation because it is important to take into account that different species have different vulnerability levels (Groves, 1994; Groves et al., 2002). Consequently, classifying species into different threat levels has become a powerful ally in species conservation (Gärdenfors, 2001; Morais et al., 2012; Possingham et al., 2002). The scheme of the IUCN (International Union for Conservation of Nature) is the most used and accepted system for classifying species in different threat categories (Mace et al., 2008). To classify species according to their extinction risk, the IUCN scheme considers a taxon's spatial and temporal distribution and its life history. However, the IUCN scheme has some biases (Abeli et al., 2009), such as some species with a great risk for extinction being classified as deficient data (Bland et al., 2015), although some researchers have averred that the data-deficient species are more likely to be rare (Corlett, 2016; Roberts et al., 2016).

Rare species in a community can result from a combination of geographic range, population size and habitat affinity (Broennimann et al., 2005; Caiafa and Martins, 2010; Fattorini, 2014; Maciel and Martins, 2019; Manne and Pimm, 2001; Rabinowitz, 1981; Rey Benayas et al., 1999; Yu and Dobson, 2000). Species can have scarce (single or doubleton) or abundant populations, a narrow (stenotopy) or wide (eurytopy) geographic range and low (euryoecy) or high (stenoecy) affinity for a specific habitat(Caiafa and Martins, 2010; Rabinowitz, 1981). The arrangement of these possibilities leads to eight situations where the combination of eurytopic, euryoecious and abundant populations results in common species, whereas the other combinations result in seven forms of rarity (Rabinowitz, 1981).

The interest in rare plant species, classified according to Rabinowitz's scheme, has increased over time(Caiafa and Martins, 2010; Harnik et al., 2012; Reed et al., 2020; Yu and Dobson, 2000), but for some ecoregions, this subject is still poorly explored. In South

America, little is currently known about rare plant species, which has been investigated only in the Amazon (Pitman et al., 1999), the Atlantic forest (Caiafa and Martins, 2010) and the Cerrado (Maciel and Martins, 2019). Nothing is known about the seven forms of rarity in the South American savannas, which comprise the Cerrado, Gran Sabana, Llanos, Beni savanna, Campos Rupestres (topmontane rocky grasslands) and disjunct areas interspersed in the Caatinga (dry forest), the Amazon and the and Amazon and Atlantic forest (Olson et al., 2001; Ratter et al., 2003). It is well established that many savanna plant species have small populations (Castro et al., 1999), whereas others are endemic (Alves and Kolbek, 1994; Pennington and Ratter, 2006) or occur within geographically restricted areas (Bridgewater et al., 2004). South American savannas occur in a wide range of environmental conditions of climates (Blydenstein, 1967), soils (Goodland and Pollard, 1973) and fire frequency (Coutinho, 1990; Murphy and Bowman, 2012). As these factors can limit species on different scales, different forms of rarity are expected in the South American savannas.

These different forms of rarity are differently associated with species extinction rates, with the highest extinction rates being expected for stenoecious species with restricted distribution and low abundance (Davies et al., 2004; Purvis et al., 2000). Savannas are among the 60% of the planet's ecosystems most affected by human activities (Grantham et al., 2020). In contrast to the many threats, the area officially protected in the South American savannas is one of the smallest when compared to that in the Amazonia rain forest (Oliveira et al., 2017). In South America, few investigations have been conducted on species rarity, and they considered only species that were previously recognised as endemic or threatened (Monteiro et al., 2018; Oliveira et al., 2017). It has been claimed that rare species are unlikely to be conserved if conservation strategies are made with an exclusive focus on other groups of species (Groves et al., 2002). Despite the importance of the rare species for conservation policies (Chapman et al., 2018; Lyons and Schwartz, 2001; Mouillot et al., 2013a), no

previous study has investigated if and how each species in each form of rarity is conserved in the protected areas (PA) of the South American savanna.

Here, using Rabinowitz's scheme (Rabinowitz et al., 1984), we provide the first quantification of rare and common species in the South American savannas (SAS). More specifically, we addressed the following questions: (1) What forms of rarity does the SAS tree flora have? Considering that the SAS is rich in species and has a high environmental heterogeneity (Sarmiento, 1984) and that a large number of plant species have small populations (Castro et al., 1999), are endemic (Colli-Silva et al., 2019; Pennington and Ratter, 2006) and have a restricted geographic range (Bridgewater et al., 2004), we expected the species to show all the seven forms of rarity. (2) How are these forms of rarity distributed over the SAS? We expected that different ecoregions have different numbers of species in each form of rarity. (3) Are both rare and common species conserved in the network of the SAS protected areas (PAs)? Because the PAs were created in marginal areas with little conservation value (Margules and Pressey, 2000), we expected that the PAs would not provide conservation for all rare species in the SAS.

2. Material and Methods

2.1. Study area

We considered a savanna a vegetation complex that includes three types of physiognomy, namely: open savanna, woody savanna and savanna woodland (Coutinho, 1978; Furley, 1999, 1992). The open savanna includes physiognomies in which the dominant forms are shrubs and trees < 7 m in height and with a crown cover of up to 10%. The woody savanna encloses physiognomies with the predominance of trees with 7-14 m in height and covering 40–60% of the ground. In the savanna woodland, trees are the dominant form and attain 15 m or more in height and > 70% crown cover. The savanna vegetation complex predominates in

the SAS ecoregions such as Llanos and Cerrado (Olson et al., 2001), but as a result of the expansion and retraction of the savanna through the glacial cycles (Berrio et al., 2002; Bueno et al., 2017), areas of savannas vegetation are found in non-savanna ecoregions, such as the Amazon (Barbosa and Fearnside, 2004), the Caatinga (Castro et al., 1999; Vieira et al., 2019), the Chacos (Kunst et al., 2006; Morales et al., 2019), and Atlantic forests (Costa-Lima and Chagas, 2019; Moura et al., 2011).

2.2. Datasets

Our database was built from two datasets. To build the first dataset, we searched for floristic and phytosociological surveys of trees performed in the SAS. To do this, we used different combinations of keywords such as, for example, ‘cerrado’, ‘dense’, ‘floristic study’ (see Table S1 for the complete keyword combinations). We then searched for papers published until December 2016 in *Web of Science*, *Scopus*, *Scielo* and *Lilacs* and built a dataset containing: 1) the localisation of each study site (using the information on latitude and longitude from the surveys) and 2) a list of the species that were sampled in each site. All studies performed in the savanna vegetation complex were included in our dataset. In total, 104 references were recovered (Table S2), namely, annals ($n = 3$), articles (68), books (6), research note (1), technical bulletins (3), technical report (1), theses (18) and unpublished data (3).

To build the second dataset, we used the NeoTropTree (Oliveira-Filho, 2017), a large database containing the results of a collaboration, to compile data across the tropical region of South America. The NeoTropTree database divides the space into sites with five kilometres in diameter and lists species with their geographic localisation, besides providing physiognomies, domain and environmental variables such as climate and precipitation. We used the savanna physiognomies as filters to pick up the surveys of SAS from the NeoTropTree and to construct our second dataset.

To build our final databank (hereafter referred to as “savanna database”), we merged the two datasets, making up 949 sites surveyed (Fig. S1). For taxonomic standardisation of species and family names, we used the ‘*flora* package’ in the R environment (Carvalho, 2016), which interacts with Flora 2020 online, to generate an output containing the currently accepted taxon names. For species names not standardised by the ‘*flora* package’, we used the Taxonomic Name Resolution Service (Boyle et al., 2013).

2.3. Rabinowitz’s seven forms of rarity

To classify the forms of rarity of the SAS tree species, we followed Rabinowitz’s scheme (Rabinowitz, 1981), who proposed that species rarity can be assessed according to a restricted or wide geographic range, low or high affinity for the habitat and scarce or abundant local population (Fig. 1). Rabinowitz’s scheme is a robust method because it provides a scale of species rarity on multiple levels. The combination of the two states of each of these three parameters ($2 \times 2 \times 2$) leads to eight situations, in which one of the eight forms englobes species with a wide geographic range (eurytopic), low affinity for the habitat (euryecious) and abundant local populations, thus being common species. The other seven situations are considered the seven forms of rarity (Fig. 1). Form 1 species are eurytopic and euryoecious, but have scarce local populations, being rare due to scarcity; Form 2 species are eurytopic and have abundant local populations, but occur in a specific habitat, being rare due to the single habitat. Form 3 includes eurytopic, stenoecious species with scarce local populations, which are rare due to the single habitat and scarcity; whereas Form 4 species are stenotopic, euryoecious and have abundant local populations, being rare due to their narrow geographical range. Form 5 species are stenotopic and euryoecious with scarce local populations, being rare due to their narrow geographic range and paucity; Form 6 species are stenotopic and stenoecious, although with abundant local populations, being rare due to their narrow geographic range within a single habitat. Form 7, the most extreme form of rarity, contains stenotopic, stenoecious species with

scarce local populations, which are rare due to their narrow geographic range, single habitat and paucity. An overview of the database construction and the rarity forms classification can be seen in Figure 1.

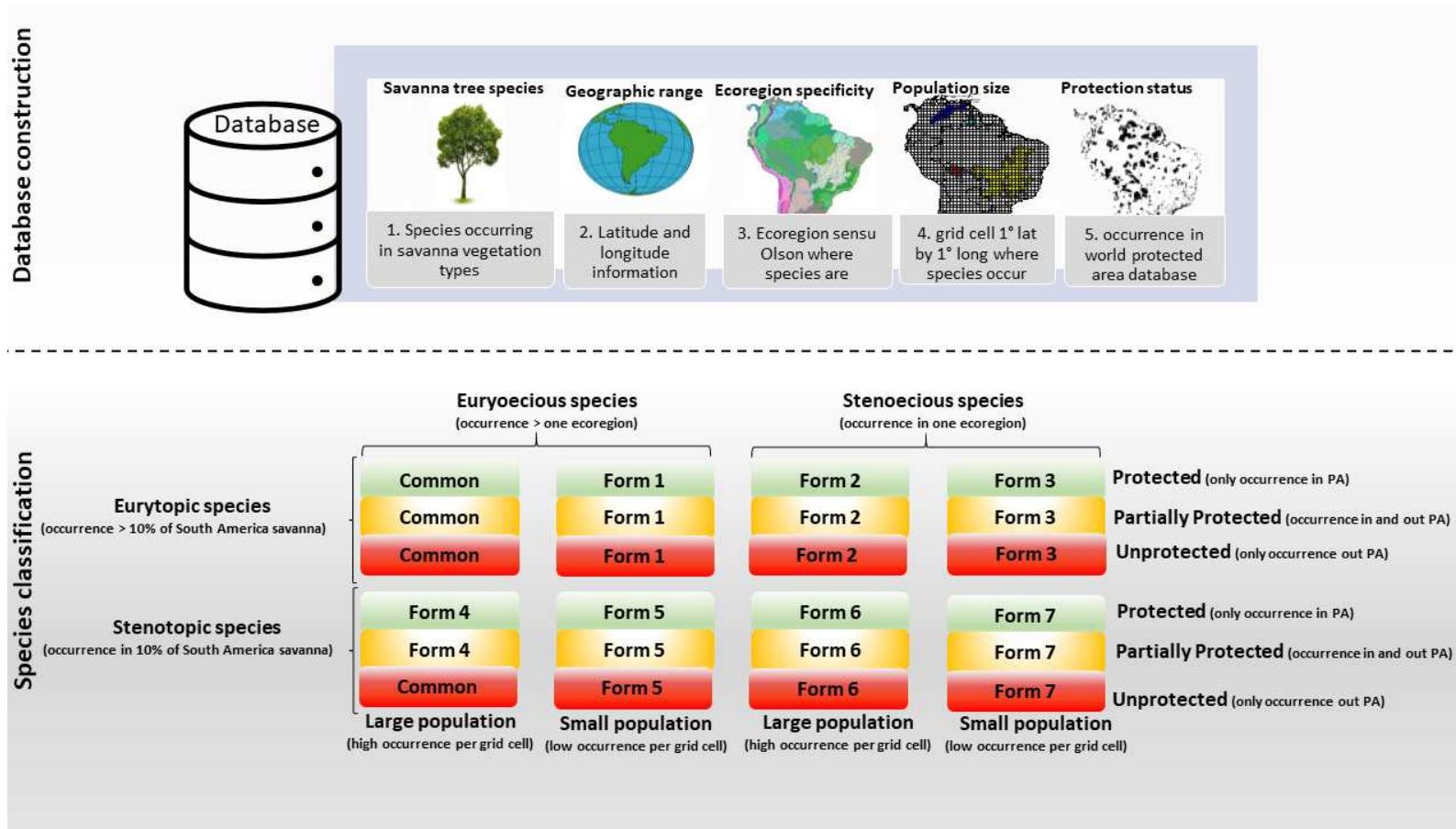


Fig. 1. Database construction including tree species with their geographical coordinates, occurrence in the South American savanna ecoregions, 1 x 1 degree of latitude x longitude cells and protected areas. Species classification according to Rabinowitz's scheme shows the eight combinations that give rise to common species and seven forms of rarity, each of them classified into three protection categories according to their occurrence exclusively in protected areas (protected species), both in protected and non-protected areas (partially protected) as well as only in unprotected areas (unprotected species).

2.3.1. Geographic range

Usually, the geographic range is calculated by considering the number of plots with the species' presence in a geographic region; species occurring in less than 10% of all plots are considered stenotopic (Sætersdal and Birks, 1997). More recently, the geographic range has been calculated from the number of latitudinal belts where the species occur (Caiafa and Martins, 2010). We think that expressing the geographic range of a species as its latitudinal range may better reflect the species distribution limits as a response to the latitudinal gradient. The SAS latitudinal range comprises 35 belts from latitude 9° N to 25° S and is equivalent to a linear distance of around 3,939.82 km (Fig. S1). To attribute the species geographic range, we followed Caifa and Martins (2010) and considered the number of latitudinal belts within which each species occurs. We considered as stenotopic the species occurring in three or less latitudinal belts, as this corresponds to around 10% of the total belts in the SAS. All species occurring in four or more latitudinal belts were considered eurytopic.

2.3.2. Ecoregion affinity

Although habitats are usually defined on a local scale, they can also be defined at the landscape scale (Tews et al., 2004). It is also accepted that the ecoregion concept provides a different way to classify habitat-specialist species. For example, in Canada, one study has shown that habitat-specific species were exclusive to one ecoregion (Kricsfalusi and Trevisan, 2014). Here, we use the ecoregion concept to classify habitat specificity (from now on, ecoregion specialist). Ecoregions are “relatively large units of land containing a distinct assemblage of natural communities and species, with boundaries that approximate the original extent of natural communities prior to major land-use change”(Olson et al., 2001). An ecoregion is an area where ecological processes strongly interact (Orians, 1993) and includes different levels of biological complexity for conservation planning (Olson et al., 2001; Smith et al., 2018). Currently, conservation planning is based on the ecoregion concept, not anymore

on phytophysiognomy. Therefore, the ecoregion concept can provide a more coherent base to classify the forms of rarity because it is more congruent with current international conservation strategies (Loyola et al., 2009; Smith et al., 2018). Thus, we considered as stenoecious species those occurring in only one ecoregion, whereas species that occur in two or more ecoregions were considered euryecious. For this classification, we considered all the 38 South American ecoregions where savanna communities occur (Olson et al., 2001).

2.3.3. Population size

We used cells of 1° latitude by 1° longitude to calculate the population size per species (Ratter et al., 2003; Simon and Proen  a, 2000). Although abundance was originally proposed as a measure of the number of individuals in an area of 1 hectare, we accepted the record of each species per grid cell because it showed a positive correlation with abundance per ha (Spearman test, $n = 225$; $r = 0.7$, $p = < 0.05$; Fig. S2). To avoid that low sampled grid cells resulted in low population sizes due to methodological artifacts, we assessed how each species was relatively “diluted” or “concentrated” in a grid cell. We expressed the species “content” in a grid cell as the ratio between the number of records of a species and the total number of records of all species in a grid cell and considered this ratio to be the response or dependent variable. The “dilution” or “concentration” of this content was estimated considering the ratio between the number of grid cells with the species presence and the total number of grid cells (a constant); we considered this ratio to be the explanatory or independent variable. Then, we adjusted the General Linear Model with the best fit for these two variables in R version 4.0.2 (R CoreTeam, 2019). Finally, we used a piecewise regression (Muggeo, 2008a, 2008b) to detect the breakpoint position in the saturation curve and estimate the parameters of the linear regressions under and above the breakpoint. The breakpoint is an abrupt threshold where there is a sharp change in the relationship between the response and explanatory variables. Species with a population size below the breakpoint were considered to

have a scarce population (Fig. S3). The analysis was performed in the *segment* package (Muggeo, 2008b) in R version 4.0.2 (R CoreTeam, 2019).

2.4. Data analysis

2.4.1. Forms of rarity in the SAS tree flora

To investigate the forms of rarity and the proportions of rare and common tree species in the SAS, we first built a data matrix with the species names (rows) and the following four parameters (columns): latitude, longitude, number of ecoregions where the species occur and population size calculated as described above (Fig. 1). Then, we classified each species in one of the eight categories of Rabinowitz (1981) by applying the package *rare7* (Maciel and Arlé, 2020) in R version 4.0.2 (R CoreTeam, 2019). First, the *raredata* function calculates the total amplitude of all species based on the data input. Then, this same function quantifies, for each species, the number of latitudinal belts and ecoregions where it occurs and the maximum number of individuals as a discrete variable. Since we did not have the number of individuals, one solution we found was to assign the number 1 to the species that stayed below the breakpoint and the number 3 to the species above the breakpoint. This allowed the *rarityseven* function to classify the species below the breakpoint as locally rare. Finally, we obtained a list of species with the following parameters: maximum number of belts, maximum number of ecoregions and population size (Table S3). Using this list as input data, the *rareseven* function was able to operate and classify species as rare or common.

2.4.2. Differences of rarity among SAS ecoregions

To investigate whether the number of species in each Rabinowitz's form differed among ecoregions, we created a matrix with the number of species in each Rabinowitz' category in each ecoregion. According to the Shapiro-Wilk test, the number of species did not fulfil the assumption of normality. Then, to compare the number of rare species in each ecoregion, we performed the Friedman test using the *t* function in R version 4.0.2 (R

CoreTeam, 2019). All statistical analyses were performed at a significance level of 0.05 and a confidence interval of 95%. This procedure helped us to find out how the forms of rarity were distributed among the ecoregions and how much each ecoregion stood out for its contribution to rare and common species.

2.4.3 Species protection status

We considered as protected areas (PAs) those listed in the World Database on Protected Areas (WDPA), which provides information about the number, size, year of creation, among other data, of protected areas (Fig. 1). A full description of WDPA protected areas is available on the International Union for Conservation of Nature (IUCN) site (<https://www.iucn.org>).

The species protection status considered the species occurring within or outside the PAs. We classified the species into three protection status: protected, partially protected and unprotected. Protected species occurred only within Pas, partially protected species occurred both within and outside of the Pas, and unprotected species occurred only outside PAs. To assess the species protection status, we considered each form of rarity in each SAS ecoregion. This procedure allowed us to assess the proportion of the species with any record within PAs relative to all SAS species (Cao et al., 2010).

For the partially protected species, we calculated the proportion of records within PAs relative to all records. For each species partially protected, we carried out the following analysis: $r = opa/to$, where r was the ratio between the number of records within PAs (opa) and the total number of records (to). At the end of this analysis, each partially protected species had an r value between 0 (no record within PAs) and 1 (all records within PAs). Then, we calculated the median of r (rMD) and the semi-amplitude of the 95% confidence interval (SCI) around the MD for all species together (protected, partially protected and unprotected) and for each form of rarity.

2.4.4 Sampling effect on the species protection status

To find out whether the protection status of a species was influenced by the sampling effort within PAs, for each grid cell, we calculated the proportion of harboured (protected + partially protected) species, that is, the ratio between the number of species within PAs and the total number of species. We took the proportion of harboured species in each grid cell as the dependent or response variable. To represent the sampling effort within each PA, we calculated the proportion of records within PAs in relation to all records per grid cell and took this as the explanatory variable. Then, we hypothesised that the proportion of harboured species is directly related to the sampling effort within PAs and tested this hypothesis by applying a general linear model. Because both y and x values were proportions ranging from 0 to 1, we used the *glm* function (Warton et al., 2016) in R version 4.0.2 (R CoreTeam, 2019) to adjust a general linear model with quasibinomial distribution after applying a logit transformation to the data. A positive correlation between y and x would indicate that the proportion of harboured species within PAs was dependent on the sampling effort. This helped us understand whether the species protection status was influenced by the relatively low sampling effort within PAs.

3. Results

3.1. The seven forms of rarity and common species in the SAS

We found all seven forms of rarity and common species as proposed by Rabinowitz et al. (1981) in the SAS (Table 1). Among the total 2,203 species, 40.76% (898) were classified as rare and 59.24% (1,305) as common. Form 7, the most extreme form of rarity, attained the highest proportion (16.98%) among all species. Form 3 (wide geographic range, ecoregion specificity and scarce population) had the least proportion, with only one species. Of all 2,203 species, 37.49% showed a restricted geographic range (stenotopic species), while 62.51% sowed a wide geographic range (eurytopic species); 23.97% were stenoecious species, whereas

76.03% were euryoecious species. Further, 71.99% had abundant populations, whereas 28.01% had scarce populations (Table 1).

Table 1 Number and percentage (between parentheses) of forms of rarity found in the South American savannas.

| Geographic distribution | | Wide | | Restricted | |
|----------------------------|----------------------|-------------------|--------------------|--------------------|--|
| Habitat preference | Various | Single | Various | Single | |
| Abundant population | 1,305 (59.2%) common | 26 (1.18%) Form 2 | 128 (5.81%) Form 4 | 127 (5.76%) Form 6 | |
| Scarce population | 45 (2.04%) Form 1 | 1 (0.05%) Form 3 | 197 (8.94%) Form 5 | 374 (17%) Form 7 | |

3.2. Distribution of the rare and common species among the SAS ecoregions

Our comparisons revealed differences in the numbers of rare and common species among ecoregions ($F_{(37)} = 2.927$, $p < 0.001$). The number of rare species in each ecoregion ranged from 276 (Cerrado) to 2 (Bolivian Yungas and Caatinga), and that of the common species ranged from 1,023 (Cerrado) to 30 (Bolivian montane dry forests). Two ecoregions did not show any rare species (Tocantins-Pindaré moist forests and Serra do Mar coastal forests). All seven forms of rarity were present in the Cerrado. Form 4 was the most frequent one and present in 31 ecoregions, whereas Form 2 had the lowest frequency, occurring in only one ecoregion (Cerrado). The Cerrado had the highest number of rare species (all forms of rarity), followed by the Dry Chaco, Bahia coastal forests, Campos Rupestres montane savanna, Guiana savanna and Llanos (Fig. and Table 2). The ecoregion specialists (species occurring in just one of the 38 ecoregions) ranged from 142 (Cerrado) to 1 (Bolivian Yungas, Monte Alegre várzea, and Tapajós-Xingu moist forests). Seven ecoregions had over 20 stenoecious species: Cerrado (142), Bahia coastal forests (104), Dry Chaco (49), Llanos (42), Maranhão Babassu forests (32), Guianan savanna (31) and Campos Rupestres montane savanna (28), whereas seven other ecoregions did not contain any specialist species (Fig. and Table 2).

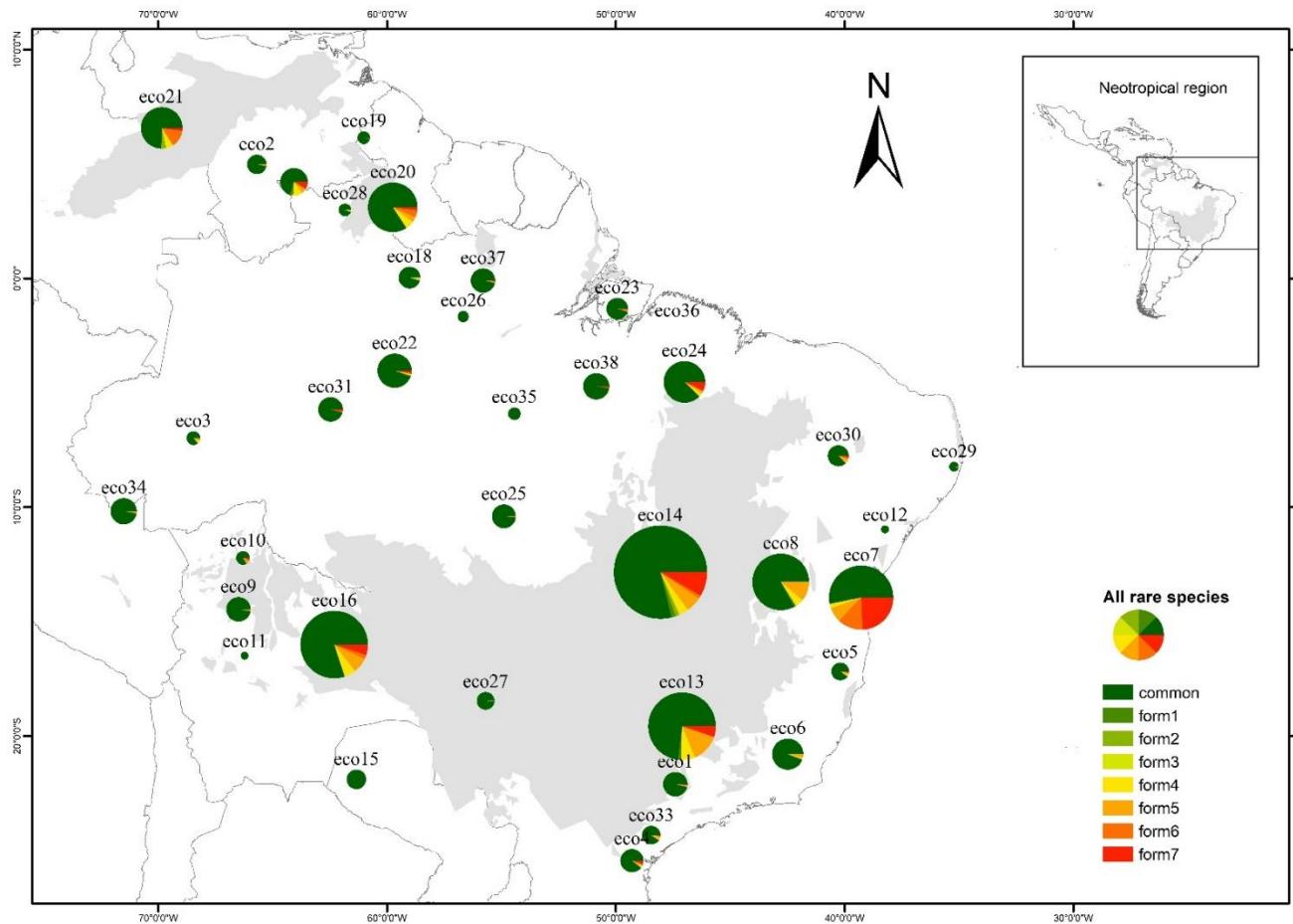


Fig. 2. Pie charts indicating the proportions of common and rare species in the 38 ecoregions (in grey) of the South American savannas. The pie codes are not the same as Olson's and are used here as a linchpin between this figure and Table 2. Pie size is proportional to the total number of rare species. The colours indicate Rabinowitz's forms of rarity.

Table 2. Numbers of total species (S), common species (common), rare species (rare), the seven forms of rarity (Form 1 to Form7), ecoregion-specialist species (es), as well as protected (prot), partially protected (part) and unprotected (unp) species in the 38 ecoregions (Olson et al. 2001) of the South American savannas (SAS). The codes in this table are not the same as Olson's codes and are used here as a lynchpin between this table and the chart in Figures 3 and 4.3.3. Protection status of species.

| codes | eco | S | common | rare | form1 | form2 | form3 | form4 | form5 | form6 | form7 | es | prot | part | unp |
|-------|---|------|--------|------|-------|-------|-------|-------|-------|-------|-------|-----|------|------|-----|
| eco1 | Alto Paraná Atlantic forests | 491 | 472 | 19 | 2 | 0 | 0 | 5 | 5 | 0 | 7 | 7 | 469 | 2 | 17 |
| eco2 | Amazon-Orinoco-Southern Caribbean mangroves | 237 | 225 | 12 | 1 | 0 | 0 | 1 | 7 | 1 | 2 | 3 | 230 | 3 | 4 |
| eco3 | Apure-Villavicencio dry forests | 46 | 40 | 6 | 0 | 0 | 0 | 4 | 0 | 2 | 0 | 2 | 41 | 0 | 9 |
| eco4 | Araucaria moist forests | 165 | 148 | 17 | 0 | 0 | 0 | 4 | 3 | 2 | 8 | 10 | 153 | 4 | 8 |
| eco5 | Atlantic Coast restingas | 106 | 96 | 10 | 0 | 0 | 0 | 4 | 4 | 0 | 2 | 2 | 97 | 1 | 8 |
| eco6 | Atlantic dry forests | 496 | 465 | 31 | 3 | 0 | 0 | 11 | 17 | 0 | 0 | 0 | 484 | 0 | 13 |
| eco7 | Bahia coastal forests | 280 | 149 | 131 | 0 | 0 | 0 | 6 | 21 | 36 | 68 | 104 | 225 | 31 | 24 |
| eco8 | Bahia interior forests | 573 | 471 | 102 | 6 | 0 | 0 | 28 | 63 | 0 | 5 | 5 | 542 | 5 | 27 |
| eco9 | Beni savanna | 433 | 414 | 19 | 5 | 0 | 0 | 2 | 9 | 1 | 2 | 3 | 423 | 1 | 9 |
| eco10 | Bolivian montane dry forests | 36 | 30 | 6 | 0 | 0 | 0 | 0 | 4 | 0 | 2 | 2 | 61 | 5 | 0 |
| eco11 | Bolivian Yungas | 67 | 65 | 2 | 1 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 66 | 0 | 1 |
| eco12 | Caatinga Enclaves moist forests | 112 | 110 | 2 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 0 | 111 | 0 | 1 |
| eco13 | Campos Rupestres montane savanna | 540 | 395 | 145 | 8 | 0 | 0 | 36 | 73 | 0 | 28 | 28 | 500 | 9 | 32 |
| eco14 | Cerrado | 1299 | 1023 | 276 | 19 | 18 | 1 | 44 | 71 | 14 | 109 | 142 | 1144 | 31 | 128 |
| eco15 | Chiquitano dry forests | 596 | 584 | 12 | 2 | 0 | 0 | 4 | 4 | 1 | 1 | 2 | 587 | 0 | 9 |
| eco16 | Dry Chaco | 705 | 560 | 145 | 5 | 0 | 0 | 40 | 51 | 14 | 35 | 49 | 621 | 15 | 73 |
| eco17 | Guianan Highlands moist forests | 80 | 56 | 24 | 3 | 0 | 0 | 9 | 5 | 0 | 7 | 7 | 67 | 12 | 0 |
| eco18 | Guianan moist forests | 230 | 215 | 15 | 1 | 0 | 0 | 10 | 2 | 1 | 1 | 2 | 222 | 0 | 8 |
| eco19 | Guianan piedmont and lowland moist forests | 89 | 84 | 5 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 88 | 1 | 0 |
| eco20 | Guianan savanna | 473 | 395 | 78 | 4 | 0 | 0 | 27 | 16 | 22 | 9 | 31 | 417 | 43 | 14 |
| eco21 | Llanos | 214 | 159 | 55 | 0 | 8 | 0 | 12 | 1 | 29 | 5 | 42 | 172 | 0 | 42 |
| eco22 | Madeira-Tapajos moist forests | 569 | 532 | 37 | 7 | 0 | 0 | 8 | 3 | 6 | 13 | 19 | 537 | 16 | 16 |
| eco23 | Marajó varzea | 270 | 255 | 15 | 1 | 0 | 0 | 1 | 7 | 1 | 5 | 6 | 265 | 3 | 2 |
| eco24 | Maranhão Babassu forests | 432 | 377 | 55 | 1 | 0 | 0 | 12 | 10 | 1 | 31 | 32 | 394 | 5 | 36 |
| eco25 | Mato Grosso seasonal forests | 617 | 599 | 18 | 3 | 0 | 0 | 1 | 7 | 1 | 6 | 7 | 603 | 5 | 10 |

| codes | eco | S | common | rare | form1 | form2 | form3 | form4 | form5 | form6 | form7 | es | prot | part | unp |
|--------------|--|----------|---------------|-------------|--------------|--------------|--------------|--------------|--------------|--------------|--------------|-----------|-------------|-------------|------------|
| eco26 | Monte Alegre várzea | 138 | 134 | 4 | 3 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 136 | 0 | 2 |
| eco27 | Pantanal | 359 | 349 | 10 | 2 | 0 | 0 | 4 | 1 | 0 | 3 | 3 | 351 | 0 | 8 |
| eco28 | Pantepui | 89 | 84 | 5 | 0 | 0 | 0 | 4 | 1 | 0 | 0 | 0 | 88 | 1 | 0 |
| eco29 | Pernambuco coastal forest | 81 | 78 | 3 | 0 | 0 | 0 | 2 | 1 | 0 | 0 | 0 | 75 | 0 | 6 |
| eco30 | Pernambuco interior forests | 106 | 92 | 14 | 1 | 0 | 0 | 2 | 7 | 0 | 4 | 4 | 94 | 0 | 12 |
| eco31 | Purus-Madeira moist forests | 323 | 304 | 19 | 4 | 0 | 0 | 3 | 1 | 4 | 7 | 11 | 309 | 10 | 4 |
| eco32 | Serra do Mar coastal forests | 96 | 96 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 96 | 0 | 0 |
| eco33 | Southern Atlantic mangroves | 117 | 106 | 11 | 0 | 0 | 0 | 4 | 5 | 0 | 2 | 2 | 104 | 0 | 13 |
| eco34 | Southwest Amazon moist forests | 412 | 390 | 22 | 6 | 0 | 0 | 2 | 9 | 1 | 4 | 5 | 400 | 3 | 9 |
| eco35 | Tapajós-Xingu moist forests | 285 | 280 | 5 | 4 | 0 | 0 | 0 | 0 | 1 | 0 | 1 | 281 | 1 | 3 |
| eco36 | Tocantins/Pindare moist forests | 40 | 40 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 40 | 0 | 0 |
| eco37 | Uatuma-Trombetas moist forests | 399 | 380 | 19 | 4 | 0 | 0 | 3 | 7 | 1 | 4 | 5 | 386 | 5 | 8 |
| eco38 | Xingu-Tocantins-Araguaia moist forests | 498 | 476 | 22 | 7 | 0 | 0 | 3 | 3 | 1 | 8 | 9 | 480 | 13 | 5 |

3.3. Protection status of species

We found only 9.30% (205) of the total species occurring only within protected areas; 74.89% (1,650) were partially protected, occurring both within and outside PAs, whereas 15.79% (348 species) were unprotected, only occurring outside PAs (Fig. 3A and Table S3 and S4). The number of rare species protected varied from 6 (common and Form 4) to 135 (Form 7). Rarity Forms 2 and 3 had no protected species. The number of partially protected species ranged from 1 (Form 3) to 1,254 (common species). The number of unprotected species ranged from 2 (Form 2) to 165 species (Form 7) (Fig. 3B). The protected species were concentrated in some ecoregions across the SAS, whereas the partially protected species were distributed over almost the entire SAS (Fig. 3B and D). Except for the common species and the rarity Forms 1 and 7, the unprotected species were also found in specific regions of the SAS (Fig. 3E).

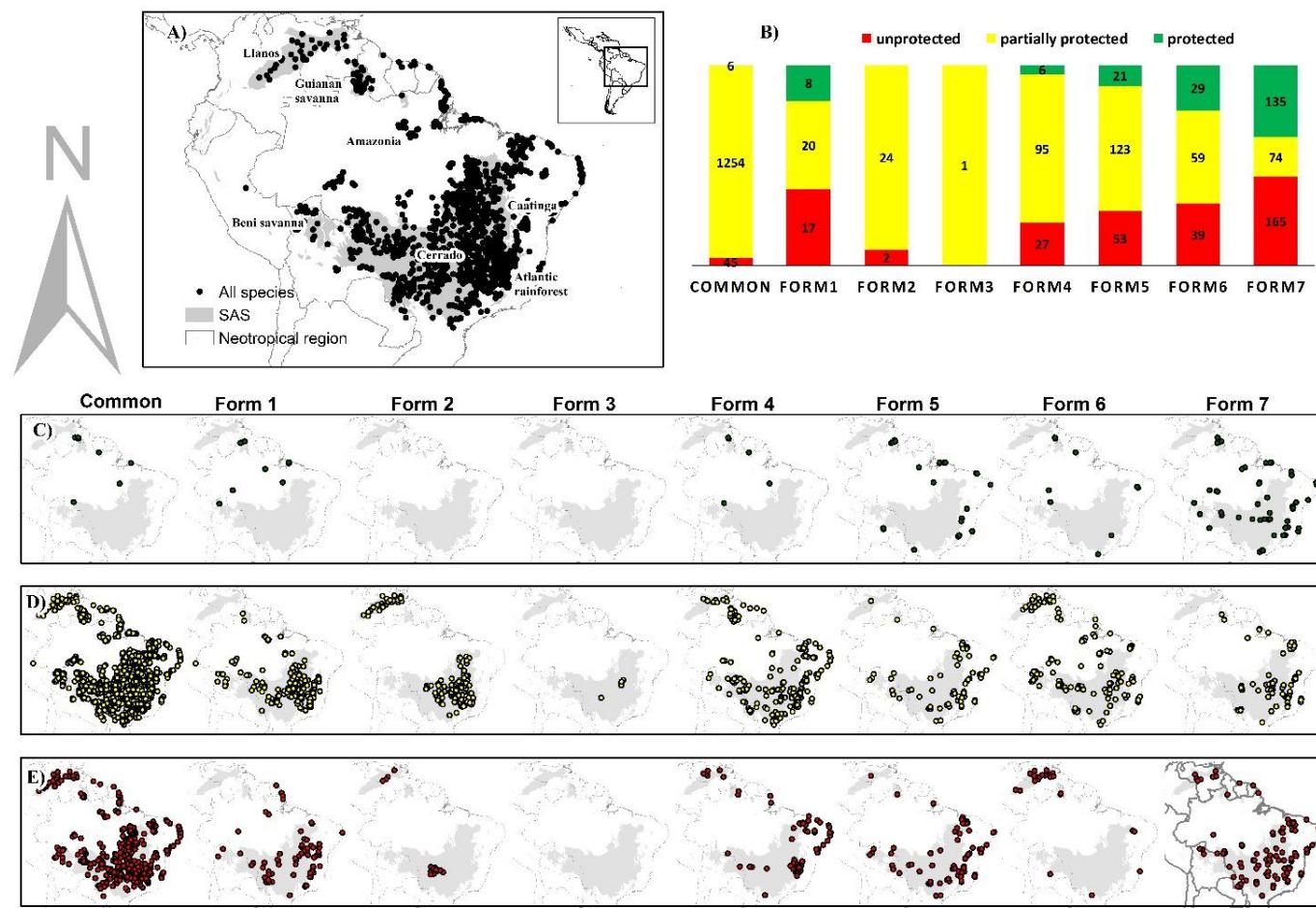
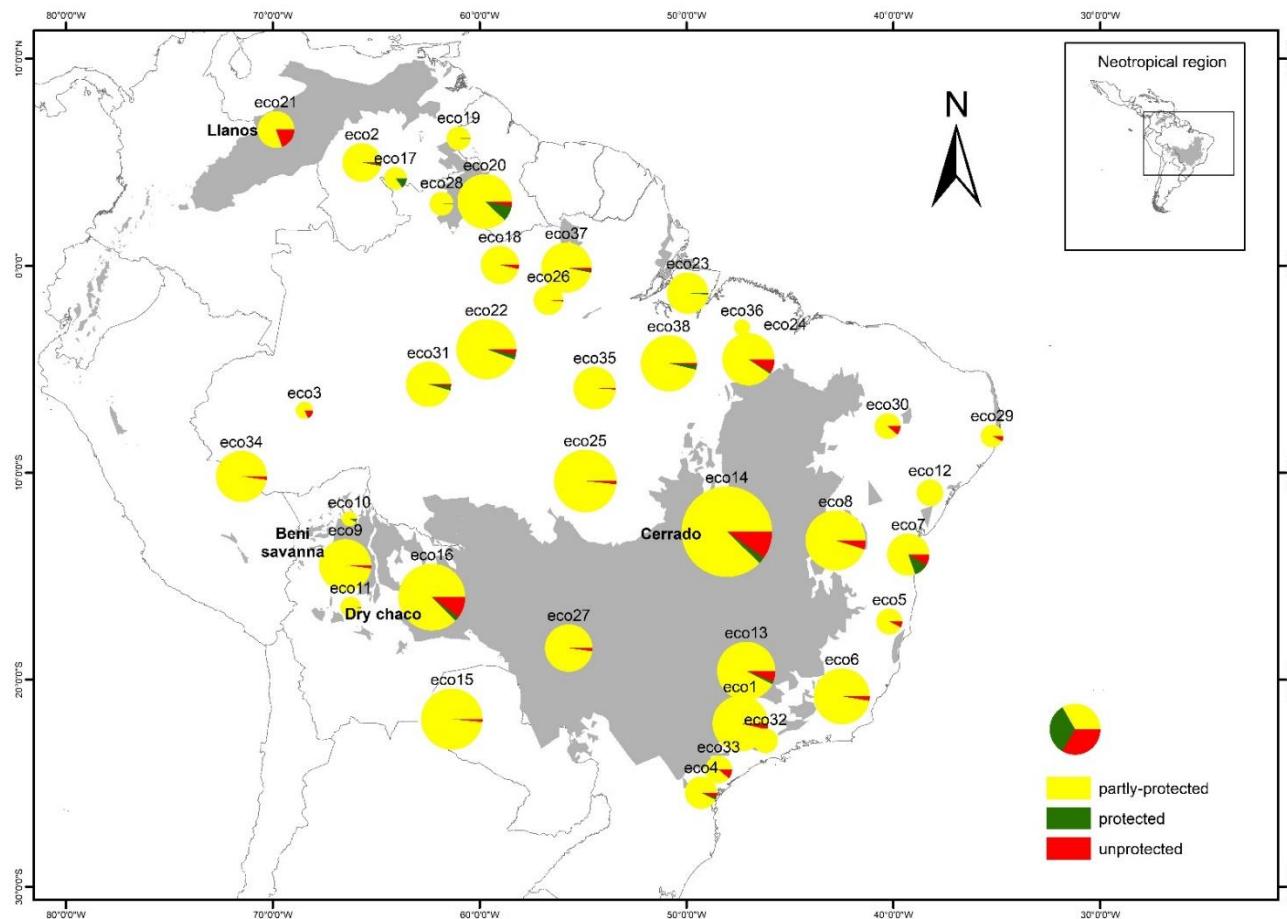


Fig. 3. (A) Tree species protection status (protected in green, partially protected in yellow and unprotected in red) in the network of protected areas in the South American savannas. (B) Number of species in each protection status (C, D, and E) and the geographic distribution of each form of rarity according to the species protection status.

1 Species in different protection categories were distributed differently among the
2 ecoregions. Just one protected species occurred in the Atlantic Coast restingas, whereas 43
3 occurred in the Guiana savanna. Partially protected species ranged from 40 in the Tocantins-
4 Pindaré moist forests to 1,144 species in the Cerrado, and unprotected species ranged from 1
5 in the Bolivian yungas and Caatinga to 128 species in the Cerrado (Fig. 3). The largest number
6 of unprotected species was found in the Cerrado (128 species), followed by the Dry Chaco
7 (73), Llanos (42), Campos Rupestres montane savanna (32) and Bahia interior forests (24).

8 When all partially protected species were considered (1,650 species), the median ratio
9 (rMD) between the records within PAs and total records was $rMD = 0.14949$ (semi-amplitude
10 of the 95% confidence interval - $SIC \pm 0.0003$), and the minimum ($rMin$) and maximum ($rMax$)
11 ratios were 0.02 and 0.88, respectively (Table S5). Of all the 2,203 species, 1,317 species had
12 maximum r above rMD , whereas 333 species had it below rMD . Rarity Form 1 showed the
13 lowest rMD ($rMD = 0.13$, $SCI \pm 0.023$), whereas Form 6 had the highest value ($rMD = 0.6$,
14 $SCI \pm 0$).



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Fig. 4. Pie charts showing the proportions of the protection status (protected in green, partially protected in yellow and unprotected in red) of common and rare species in each of the 38 ecoregions (in grey) of the South American savannas. The pie codes are the same as Olson's (2001) and are used here as a lynchpin between this figure and Table 2. Pie size is proportional to the total number of species.

3.4. Species recorded within PAs versus sampling effort

We found a sampling effect on the species protection status. The greater the ratio between the number of species within PAs and the total number of species the higher was the ratio between records within PAs and all records per grid cell (Fig. 5). The mean ratio between the number of species within PAs (harboured species) and the total number of species in a grid cell was 0.23, whereas the mean ratio between species records within PAs and the total number of records in the grid cell was 0.33. Our GLM model showed that the higher the ratio of records within PAs, the higher the proportion of harboured species in each grid cell (Fig. S4). The high value of the R square ($R^2 = 0.86$) revealed a good fit of the model, suggesting that higher numbers of harboured species per grid cell can be attributed to higher sample efforts.

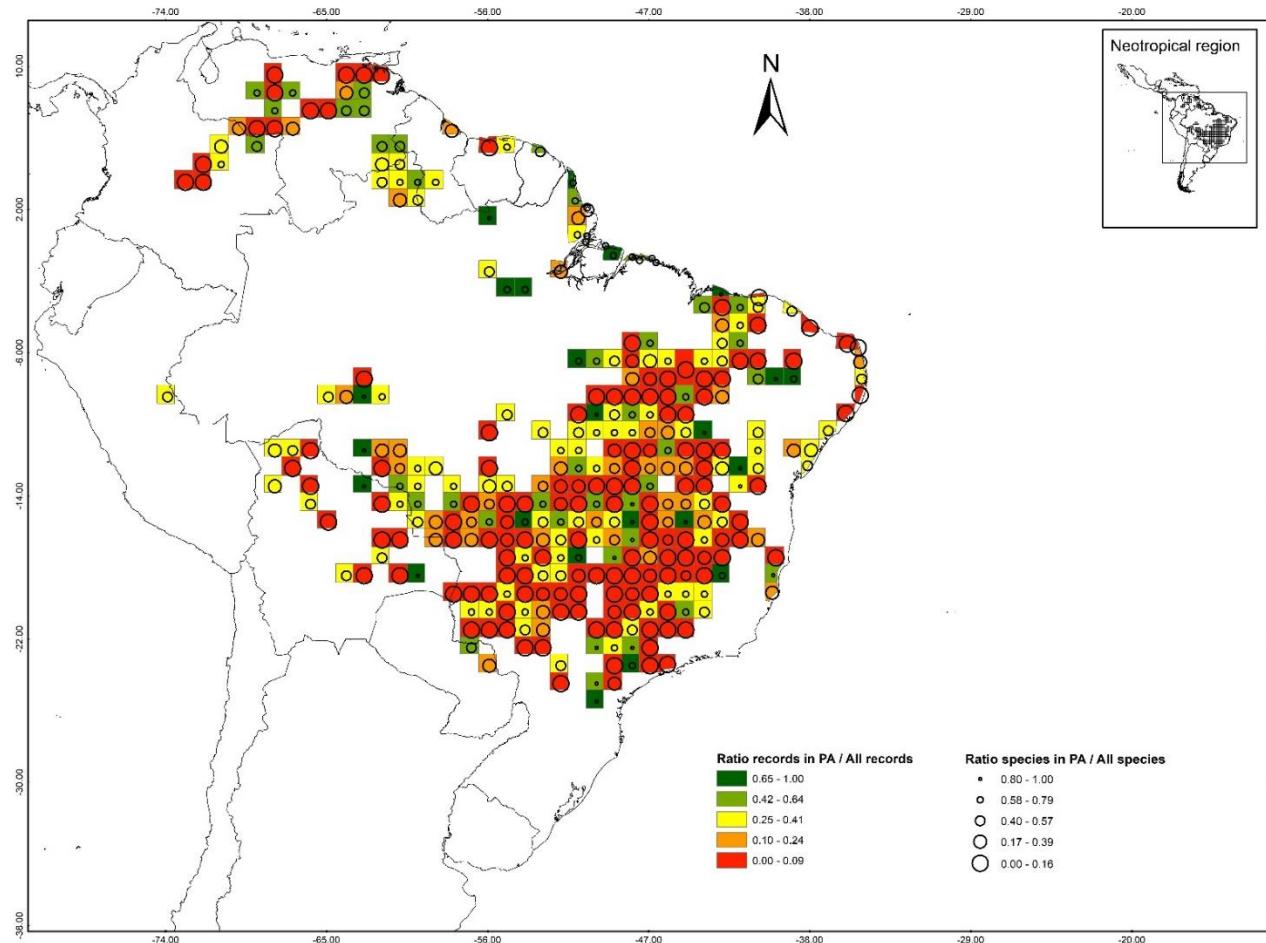


Fig. 5. Ratio between the number of records within the protected areas and all records in each grid cell of 1° of latitude by 1° of longitude. The ratio of records within PAs decreases from green to red colour (larger sampling gap within PAs).

4. Discussion

The importance of rare species is recognising as part of biodiversity conservation strategies (Groves et al., 2002; Le Berre et al., 2019; Longton and Hedderson, 2000; Maciel et al., 2016; Maciel and Arlé, 2020; McIntyre, 1992; Rey Benayas et al., 1999). This is justified because the rare species comprises a huge part of the community (Gaston, 2012), and they play a crucial role in the assemblies' structure and function, not supported by common species (Chapman et al., 2018; Gaston, 2012; Jain et al., 2014; Leitão et al., 2016; Lyons et al., 2005; Mi et al., 2012; Mouillot et al., 2013a; Soliveres et al., 2016). Moreover, because rare species are limited by area, dispersion, resources, or ecological processes (Lambeck, 1999, 1997) this makes them prone to extinction due to atrophic and stochastic processes (Lambeck, 1997; Sætersdal and Gjerde, 2011; Santoro et al., 2012). Here, we found rare species distributed throughout the Cerrado, Llanos, Beni savanna, Dry Chaco, Guiana savanna, and disjunct savannas in the Caatinga and Amazon and Atlantic rain forest. Therefore, our results provide an opportunity for plant biodiversity conservation strategies in South American savanna.

4.1. Forms of rarity among the tree species

Geographic range. The majority of the SAS rare species are geographically restricted (Forms 4, 5, 6 and 7; Table 1), with 37.49% (826) of all species being observed in only 10% or less of the SAS latitudinal range. Therefore, as in the Atlantic rain forest (Caiafa and Martins, 2010), stenotopy is the most important variable determining rarity in the SAS. Many species that are now considered characteristic of the Cerrado experienced their maximum expansion during the Last Interglacial (Bueno et al., 2017). Regions that today have disjunct savannas had suitable climatic conditions for the expansion of the savanna in the past (Bueno et al., 2017; Haffer, 1969). In some regions, the climate conditions such as precipitation and temperature are quite different than it has been in the Quaternary period (Auler and Smart, 2001). As a result, some species that have been able to overcome the climate changes condition are now

wide distribution, while species that were not able to inhabit some current conditions have become restricted over time (Foster, 2001; Hunter Jr et al., 1988; Pearman et al., 2008; Saupe et al., 2015).

Ecoregion preference. We observed 23.97% (528) of all species occurring in only one ecoregion. Each ecoregion has unique features that could have favoured the evolution of different species lineages over time (Liu et al., 2016). Presently, many ecoregions face numerous threats and, therefore, a high risk of habitat loss (Taylor et al., 2016). The distribution of plants is considered to be suited to the range of conditions found in an ecoregion, such as water table and rainfall (Oldfather, 2019). The areas of the SAS are among the richest environments, with high heterogeneity of soil, climate and relief (Coutinho, 1990; Sarmiento and Monasterio, 1975; Solbrig, 1996), thus explaining the high proportion of habitat-specialist species that we found.

Population size. We found that 28.01% (617) of all species have scarce populations (Forms 1, 3, 5 and 7), i.e., species that occur with few individuals in each savanna grid cell. The great proportion of species with scarce populations may be a result of the synergy of different factors. For instance, the high limitation of seed dispersal has been reported as a filter leading to the maintenance of small local populations while also maintaining high diversity (Salazar et al., 2012). Also, in hyperseasonal savannas, waterlogging is the main factor determining the low density of species that do not have the ability to cope with saturated soils (Batalha et al., 2005). In addition, fire is considered an important factor reducing species populations, especially of those that do not have resprout capacity (Ojeda et al., 2005; Salazar and Goldstein, 2014).

4.2. Compositions of rare species differ among ecoregions

Our analyses showed that some ecoregions have exclusive rare species that do not occur anywhere else. The Cerrado and Llanos make up the largest ecoregions of the South American

savannas. They also are part of the Global200 ecoregions (Olson et al., 2001), the priority ecoregions for conservation in the world. In the case of the Cerrado, it has been recurrently emphasised as an important ecoregion in the tropics (Giam et al., 2011; Redo et al., 2013). Nevertheless, other ecoregions, such as the Campos Rupestres, Maranhão Babassu forests and Dry Chaco, also contain large numbers of rare species, but they are not part of the Global200. The Campos Rupestres, for example, has a high floristic diversity that is driven by many factors (Neves et al., 2018) and as considered a conservation priority due to the threatened flora of the mountaintops in Brazil (Monteiro et al., 2018). Since the Global200 ecoregions are considered a priority, they should receive more funding for conservation than non-priority ecoregions. These resources are intended to maintain protected areas or sustain projects in these ecoregions. However, our findings show that more savanna ecoregions should be included in the Global200, as they have many unique species that are probably vulnerable to extinction. Therefore, we strongly suggest an urgent revision of the list of ecoregions included in the Global200.

4.3. The status of protected species of the South America Savanna

Of all analysed species, 15.84% (350) were unprotected species, meaning that these species are not inside any protected area at all. Some of these species, such as *Annona pickelii* (Diels) H. Rainer (Form 7), were classified as ecoregion specialists, geographically restricted, with scarce population, and have previously been classified as vulnerable (IUCN, 2019). Thus, our results agree with those of other authors who found that the South American network of protected areas has a conservation gap that fails to conserve a part of the biodiversity (Forero-Medina and Joppa, 2010; Jenkins and Joppa, 2009; Neves et al., 2018; Ribeiro et al., 2018; A. S. L. Rodrigues et al., 2004; Silva and Bates, 2002). Like the threatened and endemic plant species (Dryflor et al., 2016; Neves et al., 2018; Oliveira et al., 2017), we found that part of SAS rare species is also not protected in the present network of protected areas. However, the

conservation gap concerning rare species has not yet received attention in the South American protection network (Maciel et al., 2016). Efforts for flora conservation in South America have considered only threatened and endemic species (Oliveira et al., 2017; Ribeiro et al., 2018). We do not advocate that well-established groups of species should be replaced for rare species in conservation planning, but we agree with the importance of also considering the rare species in conservation planning (Broennimann et al., 2005; Maciel and Eisenlohr, 2016; Mehlman et al., 2004).

Another issue is the low proportion of species populations occurring within protected areas. Of the species occurring within any protected area, 851 (rare and common species) had less than 10% of their occurrences within PAs. This is quite disturbing for species that have scarce populations, such as, for instance, *Campomanesia xanthocarpa* (Mart.) O. Berg. Of 12 records of *C. xanthocarpa* in the entire databank, only one was within a protected area. In our analysis, *C. xanthocarpa* had, on average, two or fewer individuals per grid cell. Globally, the population size decrease of species is one of the most important triggers for local extinction (Belaid et al., 2018; Khanum, 2017). Therefore, maintaining a viable population size of both common and rare species is one of the great conservation challenges. Although the existing protected areas play a great role in the conservation of species populations, a great deal of commitment from decision-makers and managers is required.

4.4. Species records within PA versus sampling effort

We showed that the protected areas have a good performance in harbouring species, but not in keeping abundant populations. In addition, we showed that the proportion of the number of records within protected areas to the total number of records within a grid cell is low, that is, PAs have been little sampled. We found a strong positive correlation between the proportion of species and records within PAs in each grid cell, meaning that the number of harboured species depends on the sampling effort in the PAs; if more samples are taken within PAs, the

number of harboured species can increase. However, these statements cannot be conclusive because many grid cells do not have any PAs. Protected areas are mainly located in the south of the Llanos and Cerrado core area and completely absent in many other grid cells of the SAS, where many species occur in all forms of rarity. Thus, although the sampling effort within the PAs should be encouraged, the creation of new protected areas is still necessary since there are large conservation gaps in many other savanna areas in South America.

4.5. Implications for conservation

Habitat specialist species with a restricted geographic range and small populations are more likely to become extinct (Purvis et al., 2000; Staude et al., 2020). Drivers such as climate, habitat fragmentation, fire regime (Pivello, 2011; Torres and Marengo, 2014) and others, which may trigger the process of extinction of plant species, are intensifying in the SAS. These changes occur so rapidly that many species may not have time to adapt to them. As a result, the loss of these species is expected. Thus, the high proportion of rare species that we found in the savannas of South America claims the urgent need to include them in conservation planning. Besides, our results reinforce the need to conserve ecoregions (Smith et al., 2018), thus conserving habitats and, as a consequence, the many exclusive rare plant species that only occur therein.

We detected that many rare species are restricted to only one of these Olson's ecoregions. Some of these ecoregions are restricted to the country, such as the Campos Rupestres (Brazil), or embrace more than one country, such as Llanos (Colombia and Venezuela). Collaboration among countries aimed at conservation is not frequent due to different policies, and conservation actions are usually done on a national scale. In consequence, species occurring in more than one country have the chance of being conserved by different governments, while restricted species may not be conserved anywhere. In this sense, cross-boundary collaboration has been encouraged in the conservation field (Kark et al.,

2015). Our results claim for the opportunity of the application of cross-boundary collaboration to the South America savanna. For example, species restricted to Llanos, Dry chaco, Beni savanna, Cerrado and Campos Rupestres can be prioritised by the countries where these ecoregions occur.

Additionally, we observed many rare species occurring also in ecoregions where the savanna is not the predominant vegetation but occurs as enclaves or refuges (disjunction savanna). These ecoregions of disjunction savanna are located in the Amazon, for example, Madeira-Tapajos moist forest and Xingu-Tocantins-Araguaia moist forests, and in the Atlantic rain forest, for example, Bahia coastal forests and Bahia interior forests. These savanna disjunctions are well known and have been reported as a result of the expansion and contraction of large biomes during Pleistocene climate oscillations. We found that many of these species of disjunction savanna were classified in Forms 4 and 5, i.e., occurring in many ecoregions, but are locally scarce or geographically restricted. Conservation approaches are usually focused on core areas and have fewer focuses on disjunctions. However, these disjunction areas keep species that are crucial for our understanding of the evolutionary processes that result from the floristic exchange between core savanna and disjunctions. We have shown that many of these shared species are rare, and we encourage conservation strategies focused on both core savannas and their disjunctions.

Conclusion

We found that 40.76% of the 2,203 tree species in the South America savanna show a form of rarity. Of these 2,203 species, 37.49% were stenotopic (restricted distribution), 23.97% were stenoecious (ecoregion specialists), and 28.01% had a scarce population. Our findings revealed that many ecoregions have not been considered a priority for conservation, such as the Campos Rupestres. These ecoregions contain many rare species found nowhere else, and

our study emphasises the importance of reviewing the list of priority ecoregions for conservation (Global200). The existing protected areas harbour a high number of savanna tree species, but the proportion of records within PAs relative to the total number of records is low, that is, the sampling effort in PAs is relatively small. Enhancing the number of samples done in PAs can provide more accurate information about the conservation status of savanna tree species in South America. In addition, PAs are concentrated in some savanna core areas and completely absent in peripheral and disjunct areas. Considering the evolutive importance and the uniqueness of species occurring in these areas, new protected areas should be created.

CAPÍTULO 4: Defaunation and changes of climate and fire frequency have synergistic effects on biomass loss in South American savannas²

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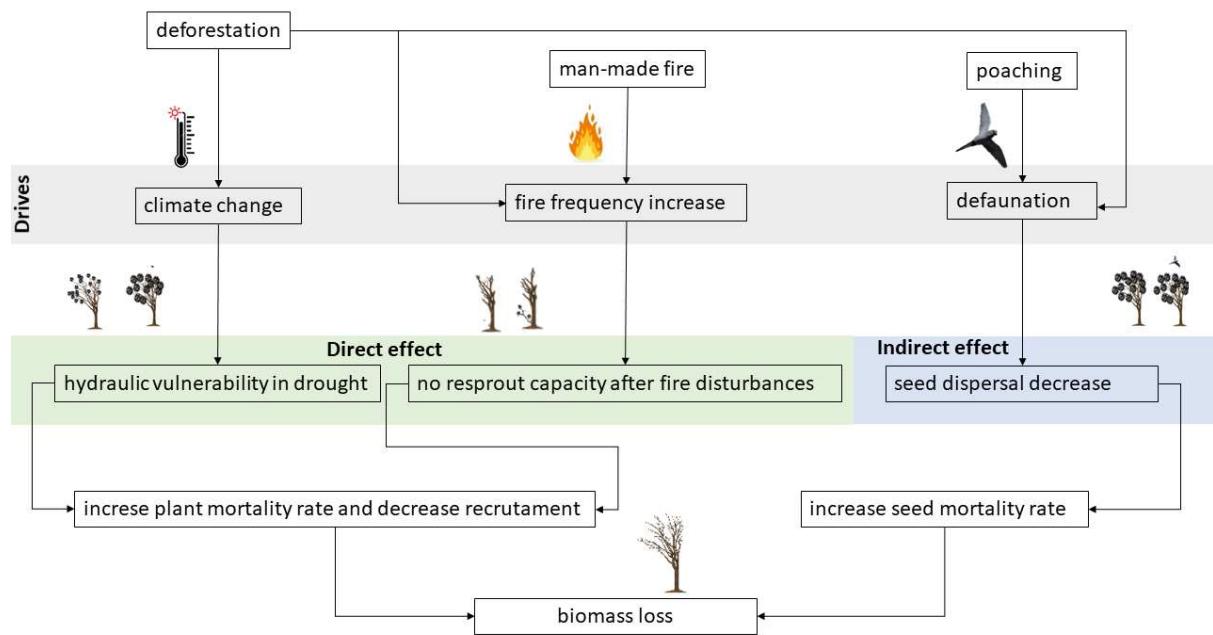
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Graphical abstract



Highlights

- We found a synergic effect of climate and fire regime changes and defaunation on the savanna biomass loss.
- In some cases, the total biomass loss exceeded 70%.
- Different physiognomies and plant functional strategies have different responses to global changes.
- Animal-dispersed, deciduous non-resprouting species have the highest biomass loss among functional groups.

Abstract

As a result of anthropogenic pressure, three effects are expected to occur in Brazilian savannas: an increase in the dry season, more frequent fire events, and defaunation. These drivers are a trigger for biodiversity loss, which hinders ecosystem services such as carbon storage. Our goal was to analyze how these drivers can affect the structure and dynamics of savanna tree species and estimated aboveground biomass (AGB). We analyzed eight areas comprising a vegetation gradient from open savanna to savanna woodland in Brazil. The species were classified by their phenological strategies (deciduous or evergreen), fire resistance (resprouting or non-resprouting), and dispersal syndrome (animal or non-animal). We modeled AGB loss in a dry season (austral winter), five fire events by decadal-series, and a defaunation scenario. Although climate, fire, and defaunation effects impact AGB separately, they also have a synergistic effect. This effect was observed in functional groups and also in the community total AGB, which exceeded 70% in some cases. AGB loss was highest in animal-dispersed, deciduous non-resprouting species. These results show that AGB loss is an outcome of both direct (e.g. inability to cope with disturbance) and indirect effects (e.g. loss of animals for seed dispersal). In the long term, these changes can hinder ecosystem services such as carbon balance and groundwater stock. Our results show that responses to environmental changes are different among phytobiognomies and areas, suggesting that management plans should take into account the savanna heterogeneity.

Keywords: climate change, fire frequency, defaunation, aboveground biomass loss, biodiversity loss, biodiversity management.

1. Introduction

The savanna is a type of vegetation found in Central and South America, Africa, and Australia under tropical climates with a defined dry season when natural fires occur (Coutinho, 1990; Durigan and Ratter, 2016; Solbrig, 1996). The savanna covers poor soils, whose aluminum gradient is the main driver of the different savanna physiognomies (Lopes and Cox, 1977). Although the savanna has a well-defined herbaceous stratum (mainly C4 grasses), it is an important carbon sink and thus a climate regulator (Ratnam et al., 2011). Additionally, the savanna plays a key role in other ecosystem services, such as the provision of food to humans, and the improvement of freshwater and air quality (Batlle-Bayer et al., 2010; Klink and Machado, 2005; Nabou et al., 2011). Nevertheless, ecosystem services provided by the savanna are now at risk of being lost due to anthropic changes in the environment, such as land conversion (Abreu et al., 2017; Cardoso Da Silva and Bates, 2002; Carvalho et al., 2009), the introduction of exotic species, defaunation (Hoffmann and Haridasan, 2008; Mistry, 1998), and climate change with consequent changes in fire frequency (Coutinho, 1990; Kauffman et al., 1994; Klink et al., 1993; Torres and Marengo, 2014), all of which increase plant mortality and lead to changes in the vegetation structure (e.g. Abreu et al., 2017; Rossatto et al., 2012).

Not all savanna woody species are expected to respond similarly to anthropic changes in the environment. For example, the seasonal tropical climate where the savanna occurs has acted as a selective pressure on leaf phenology (Van Schaik et al., 1993). Some species are deciduous and avoid drought by shedding their canopy, either entirely or partially, and hence not transpiring at significant rates during the dry season (Franco et al., 2005). Additionally, deciduous species show a decreased demand for belowground water in the dry season (Eamus, 1999; Franco et al., 2005; Lambers et al., 2008). This functional strategy provides high water use efficiency but comes at the expense of high vulnerability to cavitation due to the high gradient of water potential between the leaves and the soil (Santiago et al., 2016). Another cost

of this strategy is the high investment in leaf nitrogen to maintain high photosynthetic rates during the rainy season (Donovan et al., 2011; Edwards et al., 2014; Rossatto et al., 2013). On the other hand, other species of savanna communities are evergreen, which means they keep their canopy and transpire during the entire dry season (García-Núñez et al., 2019; Goldstein et al., 2008; Rossatto et al., 2013). In face of the climate change resulting in longer dry seasons, evergreens are expected to show higher mortality than deciduous species, leading to deciduous-dominated communities (Engelbrecht and Kursar, 2003; Hasselquist et al., 2010).

Fire is another important selective pressure on savanna tree species (Bueno et al., 2018). Some species reconstitute their aerial parts from the underground system after fire events and are called resprouters (Moreira et al., 2012; Ojeda et al., 2005). In the savannas, the resprouting strategy increases survival in comparison to the non-resprouters (Clarke et al., 2015; de Moraes et al., 2016; Ott et al., 2019; Salazar and Goldstein, 2014) because fire can kill seeds and small individual (de Medeiros and Miranda, 2005a; Hoffmann and Solbrig, 2003) but commonly does not damage underground structures. In a future scenario of increasing fire frequency due to longer and drier periods, resprouters can experience an increase in abundance whereas non-resprouters are likely to become less abundant (Hoffmann, 1998; Ojeda et al., 2005; Salazar and Goldstein, 2014).

While climate and fire have a direct effect on plant performance (Ratnam et al., 2011), defaunation indirectly impacts vegetation structure through changes in plant-animal interactions such as seed dispersal and seed predation. The causes of defaunation are many and complex, including poaching (Ferreira et al., 2015, 2012), modification of natural fire frequency (Abom et al., 2016; Langevelde et al., 2003; Radford et al., 2015), animal trafficking (Souto et al., 2017), and habitat fragmentation (Marine, 2001). These factors have an overall stronger negative effect on large animals that act as effective seed dispersers and as top-down control of seed predators (Wright et al., 2000). As a result, there is a decline in plant recruitment

(Dantas et al., 2018) and consequently in local plant populations (Carvalho et al., 2016; Rumeu et al., 2017; Terborgh et al., 2008; Valiente-Banuet et al., 2015). Similarly to other tropical vegetation, most tree species of the Brazilian savannas rely on animals for seed dispersal (e.g., Batalha et al., 1997; Mantovani and Martins, 1988), meaning that these species are more susceptible to local extinction than those that do not depend on animals as dispersal agents (Dantas et al., 2018).

It has been shown that climate change (Bustamante et al., 2012), changes in fire frequency (Abreu et al., 2017), and defaunation (Dirzo et al., 2014) separately change ecosystem dynamics in the neotropics. However, factors affecting vegetation dynamics are many and do not act in isolation and therefore there is a growing interest in understanding how drivers act synergistically (Brook et al., 2008; Laurance and Useche, 2009; Midgley et al., 2010), i.e. whether the effects of two or more factors combined are stronger than the effect of each factor acting in isolation. For example, drought and deforestation acting together in South American rainforests cause an alternative community state (Staal et al., 2015). Additionally, poaching and fragment size have a synergistic effect on the population decline of large vertebrates (Peres, 2001). Nevertheless, there is still little knowledge of how the main drivers of the savanna dynamics act synergistically.

Here, using a process-based dynamic vegetation model (FORMIND), we evaluated the performance, by estimating aboveground biomass (AGB), of different plant functional types (PFTs) in Brazilian savannas in response to climate change, changes in fire frequency, and defaunation. We tested the following hypotheses: (1) species that are deciduous, resprouters, or not primarily dispersed by animals are less susceptible to disturbance than evergreen, non-resprouting, and animal-dispersed species; (2) the synergistic effects of climate change, changes in fire frequency, and defaunation on AGB are greater than the isolated effects of each disturbance type.

2. Methods

2.1. Study sites

The Brazilian savanna comprises the core area of the Brazilian Cerrado and the Amazonian disjunction savannas (e.g., Huber, 1987; Salazar et al., 2015). Our data included eight sites of open savanna (OS), woody savanna (WS), and savanna woodland (SW) (Coutinho, 1978; Furley, 1999, 1992). The OS is dominated by shrubs and trees <7m in height and up to 10% of crown cover. The physiognomy of WS is dominated by 7-14m tall trees with crown cover between 40% and 60%. In the SW, trees are taller than 15m and cover more than 70% of the ground (Fig. S1).

Open Savanna. We used a site at the northeastern portion of the state of Roraima, northern Brazil. This physiognomy comprises part of the Rio Branco-Rupununi complex of Brazil, Venezuela, and the Republic of Guyana (Barbosa and Fearnside, 2004). In Roraima, the OS is popularly known as '*Campos do Rio Branco*' or '*lavrado*' and occupies an area of ~40,000 km² (Barbosa et al., 2007). The climate is Awi (tropical humid without a cold season) in the Köppen classification system (Barbosa et al., 2007).

Woody Savanna. Our data come from five sites in different savanna regions. Three sites are located in the Cerrado core in Brazil (WS1, WS2, and WS3) and have been surveyed twice, in 2009 and 2013 (Silva et al., 2019). The Aw-tropical seasonal climate is predominant in these sites, with a dry season lasting from April to September. The annual average temperature varies between 19 and 20°C with an annual mean precipitation of 1600-1900 mm (Alvares et al., 2013). The fourth WS site (WS4) is located in the state of Minas Gerais, southeastern Brazil. The climate is highland tropical, Cwb, with an annual average temperature of 23°C, and annual mean precipitation of 700-2000 mm (Alvares et al., 2013). Our fifth WS site (WS5) is located in the municipality of Carolina in the state of Maranhão, northeastern Brazil (de Medeiros et

al., 2008), in the transition between the Cerrado and the Amazon, thus having ecotone characteristics. The predominant climatic regime is Aw.

Savanna Woodland. We used data from two sites in southeastern Brazil, both located at the municipality of Bauru in the state of São Paulo. The Cwag climate subtype, temperate, moderately rainy with a dry winter, is predominant in the region (Cavassan, 2002, 1990). In Brazil, this climate is characterized by an average annual temperature ranging between 17 and 24°C, and average precipitation of 700-2500 mm (Alvares et al., 2013). In Bauru municipality, the minimum temperature can be as low as 13.6°C and the average annual rainfall is 1262.9mm (Cavassan and Weiser, 2015). An overview of the eight study sites is shown in Table 1 and Fig. S1.

Table 1. Location of the sites used in this study with physiognomies of open savanna (OS), wood savanna (WS), and savanna woodland (SW). Details for each site can be found in the respective references.

| Site | Locality | Forest type | Domain | Reference |
|------|-------------------|-------------------------------------|-----------------------------|----------------------------|
| OS1 | Água Boa-RR | <i>Grassland + Parkland</i> savanna | Amazonia | Barbosa and Fearnside 2005 |
| WS1 | Rio Paraiso-GO | <i>cerrado stricto sensu</i> | Cerrado | Eliseu-Silva et al. 2019 |
| WS2 | Batalhão-GO | <i>cerrado stricto sensu</i> | Cerrado | Eliseu-Silva et al. 2019 |
| WS3 | Lajeado-GO | <i>cerrado stricto sensu</i> | Cerrado | Eliseu-Silva et al. 2019 |
| WS4 | Serra do Cipó –MG | <i>cerrado dense</i> | Cerrado | Sarmento 2014 |
| WS5 | Carolina-MA | <i>cerrado stricto sensu</i> | Transition Cerrado-Amazônia | Medeiros et al. 2008 |
| SW1 | Bauru-SP | Cerradao | Cerrado | Unpublished data |
| SW2 | Bauru-SP | Cerradao | Cerrado | Unpublished data |

2.2. FORMIND model

To model the dynamics of the vegetation, we used FORMIND (e.g. Fischer et al., 2016), which is a dynamic vegetation model that simulates tree growth based on the carbon balance of individual trees (Köhler and Huth, 1998) resulting from basic processes such as competition for light and water, seed dispersal, recruitment, and mortality (Fischer et al., 2016). To model the community dynamics, the species are classified into plant functional types (PFTs) according to their life histories, based on characteristics such as seed dispersal and resource-use strategies. Simulations are performed for a minimum area of 1 ha, and this area is divided into plots of 20 × 20m, where the trees interact within each plot (Fischer et al., 2016). The survival and growth rates of the trees depend on environmental factors such as soil water table, temperature, and disturbance rates (e.g., Köhler and Huth, 2007). The competitive capacity of each PFT determines its basal area and AGB accumulation rate. Therefore, species with greater capacity to acquire resources provide a greater contribution to the community basal area and AGB (Pagel et al., 1991).

2.3. Plant functional types

We classified the species (trees and shrubs) surveyed in our eight study sites into eight PFTs, which were a combination of leaf phenology (deciduous *vs.* evergreen), resprouting capacity (resprouters *vs.* non-resprouters), and seed dispersal strategy (animal *vs.* non-animal) (Table S1). Because plant species with small, medium, and large seeds that are dispersed by animals of different sizes have been reported to decline progressively and quickly in the savanna (Harrison et al., 2013), we did not discriminate seed size in our models.

2.4. Baseline model (Control scenario – C)

For each of the eight study sites, we built a baseline model assuming political intervention that resulted in the stall of anthropic effects. In this scenario, the CO₂ emission was controlled until 2050 and CO₂ concentration remained around the current 400ppm (Moss

et al., 2010). The other characteristics of the C model were parameterized in the submodels below.

Advanced climate submodel: here we included precipitation and temperature for each site. Data were extracted from the Brazilian National Meteorology Institute (INMET - Instituto Nacional de Meteorologia, 2018) using the geographic position of each study site and for the same years that each site was surveyed. The dry season did not exceed six months, with a monthly average of <10-40mm of rain. The annual precipitation reached up to 1,500mm with a minimum monthly average of 200mm. The annual temperature remained at an average of 25°C.

Soil water submodel: in FORMIND, soil water content is hourly calculated according to the volume of precipitation, interception, and runoff (Kumagai et al., 2004). In order to represent realistic gap/successional dynamics within water-limited environments, we included the effect of rooting in soils as a facilitation process for water availability. Therefore, in our simulations we assumed that water infiltration (I) was facilitated by root mass density, using the relationship developed by Hart (2017):

$$I = 4.8 \cdot B_{root} + 0.6 \quad (1)$$

Where I is in cm/hr and B_{root} is root mass density (kg/mg). This allows for larger soil water availability through the open pore structure of strongly rooted soils (Zhou and Shangguan 2007; Ostonen et al. 2007; Bartens et al. 2009) and a dynamic in which plants with lower hydraulic conductivities and photosynthesis rates (evergreen PFTs in our model) are selected for in uncolonized soils with low infiltration rates and high potential evapotranspiration. Once the volume of water in the soil increases through rooting by evergreen species (San José and Medina 1975), these are substituted by deciduous (Nardoto et al. 1998; Ludwig et al. 2004). In our model, deciduous species have lower water use efficiency but a higher photosynthetic capacity than evergreen species (Jackson et al. 1999; Franco et al. 2004); this provides a

functional advantage to deciduous against evergreen species in less water-limited environments.

Fire submodel: in our models, the fire was parameterized from empirical studies, which have shown that the frequency of natural fire is two or three episodes per 10-yr interval (*e.g.* Pivello and Norton, 1996) and that each episode burns ~40% of Cerrado areas (Pereira Júnior et al., 2014; Ramos-Neto and Pivello, 2000). In our models, the mortality rate after the fire disturbance was assumed as inversely proportional to the stem height (de Medeiros and Miranda, 2005b; Massi and Franco, 2016; Vale and Lopes, 2010). Additionally, fire tolerance of PFTs was defined on a scale of 1 (no tolerance) to 4 (tolerant). We specified non-resprouters as 2 (less fire-tolerant) and resprouters as 3 (more fire-tolerant) in our models, because resprouters are in general more successful than non-resprouters after fire events (Hoffmann, 1998; Ojeda et al., 2005).

Seed production and dispersal of mother trees submodel: seed limitation can be affected by the number of seeds produced (source limitation) and by the spatial dispersal capacity of seeds (dispersal limitation) (Nathan and Muller-Landau, 2000; Turnbull et al., 2000). Thus, more abundant species are likely to produce a larger number of seeds than rare species (Salazar et al., 2012), while taller species would disperse seeds to a greater distance from parent plants than shorter species (Thomson et al., 2011). We parameterized our models using mean species abundance and height per PFT (see de Paula et al. 2018). Additionally, we specified that only 20% of the total seed production was dispersed (see Mariano and Christianini, 2016; Salazar et al., 2012). Because dispersed seeds may not find suitable sites for germination and further plant establishment (Herrera et al., 1994; Nathan and Muller-Landau, 2000), or be destroyed by seed predators (Wright et al., 2000), the mean post-dispersal seed mortality in our models was 10% for all individuals (Salazar et al., 2012).

2.4.2 The model accuracy

In order to parameterise our FORMIND models with the basal area (BA) of each PFT in the communities, we used field data from three savanna sites (Table S2). These three sites were chosen because they were the only ones that had two inventories done in different times. We calculated the percentage BA of the eight PFTs (Table S1) and the range of each PFT's BA for the three sites. Once the FORMIND had been parameterized for three sites, we were able to run the FORMIND for the other ones. We run the FORMIND models for 1,000 years (e.g. Dantas et al. 2018). For all sites, we selected parameter values that resulted in BA within the empirical range, indicating the good performance of our models (Fig. S1). In order to test the accuracy of AGB predicted by FORMIND, we used the AGB value reported in the literature as an AGB reference value (Table S3). Then, for each area, we compared the AGB averages provided by FORMIND and standard deviation with the maximum and minimum AGB reference values.

2.4.3 Disturbance scenarios

We created three individual disturbance scenarios: climate change (CC), changes in fire frequency (CF), and defaunation (DE) (Table S4), in all of which there would be no effective policies to control anthropic actions (e.g. Jarvie and Svenning, 2018). In our CC scenario, the length of the dry season was increased from six to seven months and the average maximum daily temperature was increased from 25°C to 27°C (e.g. Torres and Marengo, 2014). We set out the temperature increase at 2°C because it is expected to cause large fires in the future (Silva et al., 2016; P. S. Silva et al., 2019). In the CF scenario, fire frequency was increased to five episodes per 10-yr interval. In both scenarios, the water table decreased relative to the control scenario (e.g. Gosling and Arnell, 2016). In the DE scenario, seed survival of animal-dispersed species was decreased from 10% to 5% due to the decrease of suitable sites (Herrera et al., 1994; Nathan and Muller-Landau, 2000) and post-dispersal seed predation of both animal

and non-animal dispersed species increased from 10% (before defaunation) to 50% (after defaunation) (Wright et al., 2000).

Because our main interest was to understand how the synergistic effects of climate change, changes in fire frequency, and defaunation lead to AGB change of savanna communities, we ran FORMIND with combinations of disturbances, namely climate change and changes in fire frequency (CC.CF), climate change and defaunation (CC.DE), and climate change, changes in fire frequency, and defaunation (CC.CF.DE). The effect of the different disturbance scenarios on the vegetation was assessed by the percentage in the AGB change relative to the C scenario. The AGB change ($agbc$) was calculated as $agbc = [(bI - bn)/bI]$, where, bI was the AGB in the control scenario (C) and bn was the AGB in each disturbance scenario. The $agbc$ could range from 1 (100% AGB increase) to -1 (100% AGB loss).

2.5. Statistical analysis

We compared $agbc$ among the different disturbance scenarios. The comparisons were done among each area of study (eight areas), phytophysiognomies (open savanna, wood savanna, and savanna woodland), and functional strategies (deciduous vs. evergreen, resprouters vs. non-resprouters, and animal vs. non-animal). In these comparisons, the mean $agbc$ was calculated for each disturbance scenario. Because the data did not meet the assumptions of the parametric statistical, we used Games-Howell test, which is recommended when the distribution does not meet the assumption of normality (see Ruxton and Beauchamp, 2008). The 95% confidence interval around the average was calculated by the bootstrap method with 10,000 resamples (Streukens and Leroi-Werelds, 2016).

3. Results

The AGB (ton.ha^{-1}) estimated by FORMIND ranged from 11.74 (open savanna) to 73.27 (savanna woodland) for areas in the baseline scenario (C). Comparisons between the

AGB estimated by FORMIND and AGB reference values reported in studies of the savanna revealed the performance of the FORMIND model was good. Apart from one area, the AGB estimated by FORMIND was within the range (min and max AGB value) of the AGB references (Fig. S2).

3.1 Change in AGB in the areas

Our test revealed a significant change in average AGB for disturbance scenarios in all eight areas (Table 2). However, no significant differences were found in average AGB among all areas in the same disturbance scenario (Table 2). The increase in average AGB between areas ranged from 0.2% to 35%, while the decrease ranged from 21% to 77% (Table 2). In the climate change scenario (CC), there was an average increase in AGB in five areas (Table 2, Fig. 1a, e, f, g, and h) and a decrease in three areas (Table 1, Fig. 1b, c, and d). The CC.FC scenario (climate and fire change combined) produced a decrease in the average AGB in all areas (Table 2, Fig. 1). In the CC.DE scenario (climate change and defaunation-affected), the average AGB increased in half of the areas (Table 2; Fig. 1a, e, g, and h), and decreased in the other half (Table 2; Fig. 1b, c, d, and f). The synergism scenario produced a decrease in average AGB in all areas and it was greater than CC.FC (Table 2; Fig. 1). Overall, the synergistic effects of all types of disturbance (the CC.CF.DE scenario) resulted in the highest decrease (35-77%) in AGB (Table 2).

Table 2. Average rate of AGB (aboveground biomass) change ($agbc$) in four global change scenarios: climate change (CC), climate change and change in fire frequency (CC.CF), climate change, and defaunation (CC.DE), and all factors together (CC.CF.DE). Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. All models were run for 1000 years. The 95% confidence interval, calculated by the bootstrap re-sampling method with 10000 repetitions, is provided. Significant statistical differences between average AGB are indicated by different letters among disturbance scenarios in the same area (uppercase letters), and among areas in the same disturbance scenario (lowercase letters). The letters in the last column correspond to the boxplot of each area in Figure 1.

| scenario | | CC | | CC.CF | | CC.DE | | CC.CF.DE | | Fig. 1 |
|--------------|-----|---------------------|--------------|---------------------|--------------|---------------------|--------------|---------------------|--------------|--------|
| areas | n | $agbc$ | 95% CI | |
| OS1 | 995 | 0.26 ^{Aa} | 0.22, 0.29 | -0.29 ^{Ba} | -0.32, -0.27 | 0.17 ^{Ca} | 0.13, 0.2 | -0.36 ^{Da} | -0.38, -0.34 | a |
| WS1 | 995 | -0.21 ^{Ab} | -0.23, -0.19 | -0.71 ^{Bb} | -0.72, -0.7 | -0.41 ^{Cb} | -0.42, -0.4 | -0.74 ^{Db} | -0.75, -0.74 | b |
| WS2 | 995 | -0.23 ^{Ab} | -0.25, -0.21 | -0.77 ^{Bc} | -0.78, -0.76 | -0.45 ^{Cc} | -0.47, -0.43 | -0.77 ^{Bc} | -0.77, -0.76 | c |
| WS3 | 995 | -0.27 ^{Ac} | -0.29, -0.26 | -0.71 ^{Bb} | -0.72, -0.7 | -0.41 ^{Cb} | -0.43, -0.4 | -0.74 ^{Db} | -0.75, -0.74 | d |
| WS4 | 996 | 0.36 ^{Ad} | 0.33, 0.39 | -0.53 ^{Bd} | -0.56, -0.51 | 0.02 ^{Cd} | -0.003, 0.05 | -0.56 ^{Dd} | -0.58, -0.54 | e |
| WS5 | 996 | 0.21 ^{Aa} | 0.18, 0.24 | -0.51 ^{Bd} | -0.53, -0.48 | 0.04 ^{Cd} | 0.01, 0.07 | -0.58 ^{Dd} | -0.6, -0.56 | f |
| SW1 | 992 | 0.23 ^{Aa} | 0.21, 0.25 | -0.24 ^{Be} | -0.25, -0.22 | 0.06 ^{Cd} | 0.04, 0.07 | -0.39 ^{Da} | -0.41, -0.38 | g |
| SW2 | 989 | 0.35 ^{Ae} | 0.33, 0.37 | -0.21 ^{Be} | -0.23, -0.19 | 0.18 ^{Ca} | 0.16, 0.2 | -0.35 ^{Da} | -0.38, -0.34 | h |

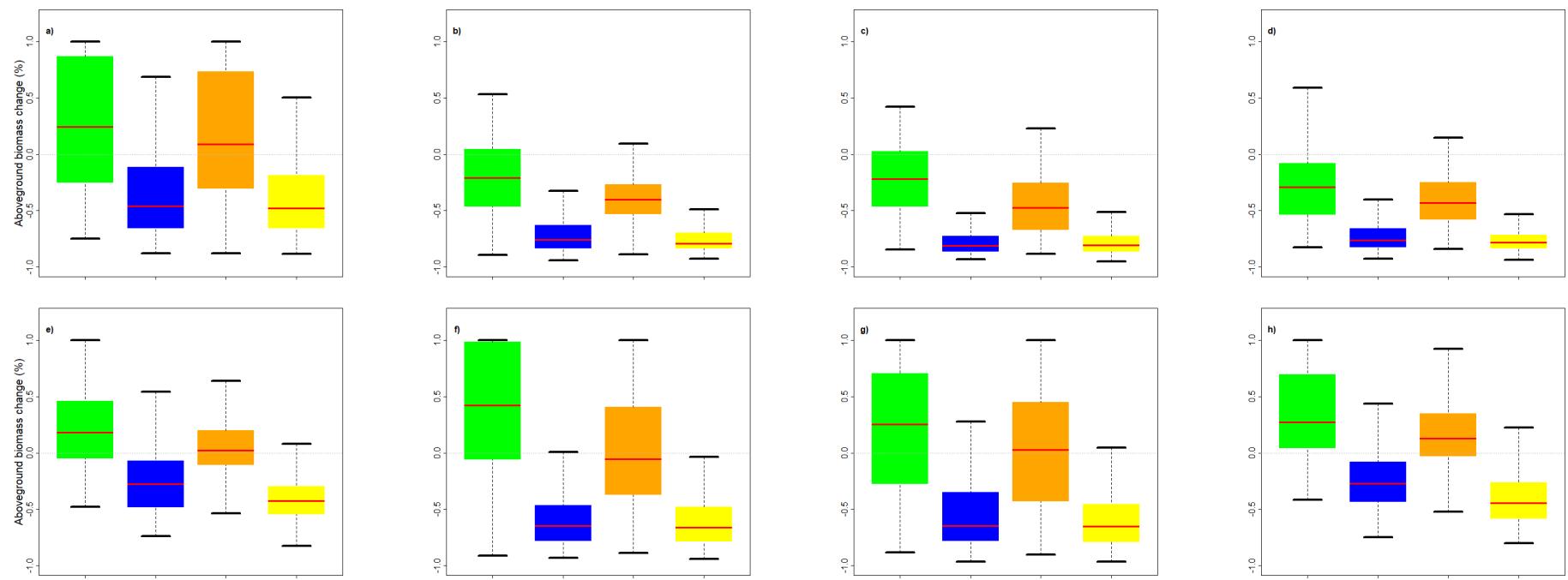


Fig. 1. Changes in aboveground biomass (AGB) in eight study areas (a-h) based on a 1000-year simulation. The colors green (climate change), blue (climate change and change in fire effects), orange (climate change and defaunation), and yellow (all factors together) indicate global change scenarios. Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. The red lines of each boxplot indicate the average AGB in the global change scenarios.

3.2 AGB change within phytophysiognomies

Our test revealed a significant change in the average AGB among the disturbance scenarios for all phytophysiognomies (Table 3). Our test revealed significant AGB losses in woody savanna relative to woodland and open savanna resulting from all disturbance scenarios (Table 3). Average AGB losses were also significantly higher in open savanna than in savanna woodland in the climate and fire change scenario (Table 3). The same overall pattern we found for communities was also observed for the phytophysiognomies (Table 3; Fig. 2). The increase in average AGB ranged from 17% to 26% (open savanna) and from 23% to 38% (savanna woodland), while the decrease ranged from 29% to 36% (open savanna), from 0.2% to 35% (woody savanna), and from 23% to 38% (savanna woodland). Two scenarios (climate change and climate change and defaunation effect) produced an increase in the average AGB for two phytophysiognomies, open savanna and savanna woodland, and a decrease in that of woody savanna (Table 3). Climate change and change in fire frequency (CC.CF), as well as the synergy of all factors (CC.CF.DE), produced a decrease in average AGB for all physiognomies (Table 3; Fig. 2). The CC.CF.DE scenario resulted in the greatest decrease (41.93% and 76.06%) of AGB within the phytophysiognomies.

Table 3. Change in aboveground biomass (AGB) in three physiognomies: open savanna (OS), woody savanna (WS), and savanna woodland (SW). The average rate of AGB change (*agbc*) in four global change scenarios: climate change (CC), climate change and change in fire frequency (CC.CF), climate change, and defaunation (CC.DE), and all factors together (CC.CF.DE). Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. All models were run for 1000 years. The 95% confidence interval, calculated by the bootstrap re-sampling method with 10000 repetitions, is provided. Significant statistical differences between average AGB are indicated by different letters between disturbance scenarios in the same phytophysiognomy (uppercase letters), and between phytophysiognomies in the same disturbance scenario (lowercase letters). The letters in the last column correspond to the boxplot of each area in Figure 2.

| scenario | | CC | | CC.CF | | CC.DE | | CC.CF.DE | | Fig. 2 |
|----------|-------|---------------------|--------------|---------------------|--------------|---------------------|--------------|---------------------|--------------|--------|
| | n | <i>agbc</i> | 95% CI | |
| OS | 995 | 0.26 ^{Aa} | 0.22, 0.29 | -0.29 ^{Ba} | -0.38, -0.29 | 0.17 ^{Ca} | 0.13, 0.20 | -0.36 ^{Da} | -0.38, -0.34 | a |
| WS | 4,975 | -0.09 ^{Ab} | -0.13, -0.08 | -0.71 ^{Bb} | -0.72, -70 | -0.32 ^{Cb} | -0.33, -0.32 | -0.73 ^{Db} | -0.74, -0.72 | b |
| SW | 1,984 | 0.28 ^{Aa} | 0.27, 0.30 | -0.23 ^{Bc} | -0.25, -0.22 | 0.11 ^{Cc} | 0.09, 0.12 | -0.38 ^{Da} | -0.39, -0.37 | c |

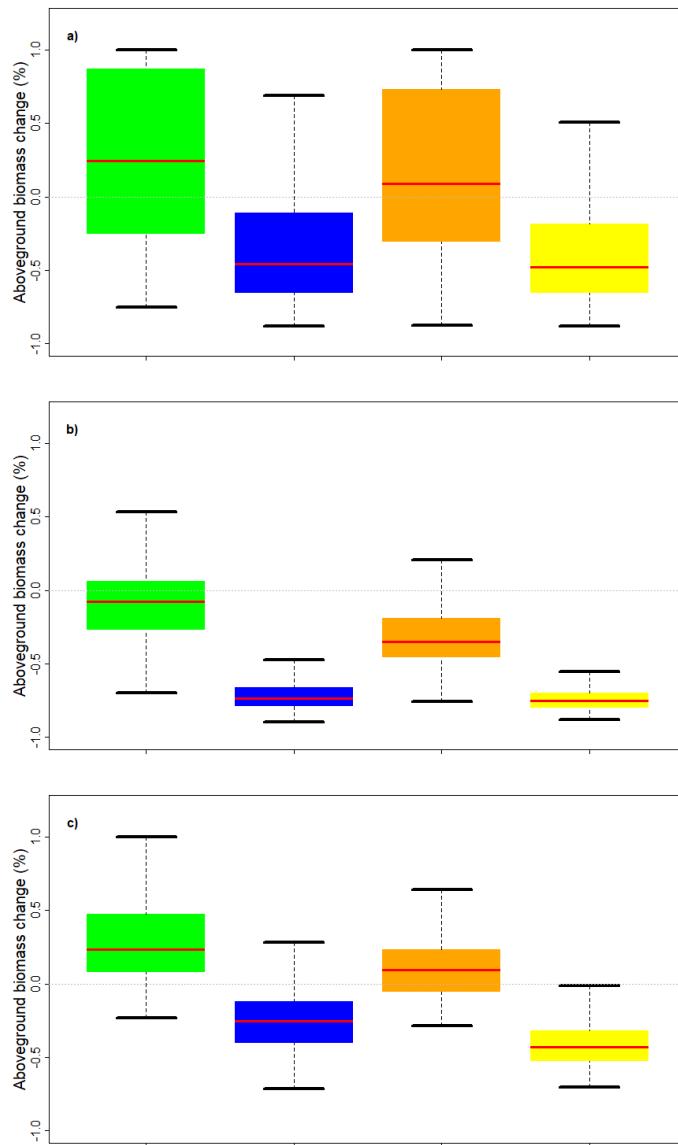


Fig. 2. Changes in aboveground biomass (AGB) in open savanna (a), woody savanna (b), and savanna woodland (c) based on a 1000-year simulation. The colors green (climate change), blue (climate change and change in fire effects), orange (climate change and defaunation), and yellow (all factors together) indicate global change scenarios. Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. The red lines of each boxplot indicate the average AGB in the global change scenarios.

3.3 AGB change and functional strategies

Changes in AGB differed among functional strategies. CC produced an increase in AGB for all functional strategies (Table 3), which ranged from 1% to 9%. All the other scenarios produced a decrease in average AGB for all the strategies, which ranged from 7% to 70% (Table 4). Comparing all functional strategies, the lowest decrease in average AGB was observed in evergreen species while the greatest decrease was observed in those species dispersed by animals and non-resprouters (Table 4; Fig. 3b and c). Deciduous species showed a lower increase and a greater decrease in average AGB than evergreen species (Table 4, Fig. 3a). Similarly, non-resprouting species exhibited a lower increase and a greater decrease in average AGB than resprouter species (Table 4, Fig. 3b). Finally, animal-dispersed species showed a lower increase and greater decrease in average AGB than species that are not dispersed by animals (Table 4, Fig. 3c).

Table 4. Change in aboveground biomass (AGB) in six plant functional types (PFTs): deciduous (DC), evergreen (EV), resprouters (RESP), non-resprouters (NRESP), animal-dispersed (ZOO), and non-animal dispersed (NZOO). The average rate of AGB change (*agbc*) in four global change scenarios: climate change (CC), climate change and change in fire frequency (CC.CF), climate change, and defaunation (CC.DE), and all factors together (CC.CF.DE). Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. All models were run for 1000 years. The 95% confidence interval, calculated by the bootstrap re-sampling method with 10000 repetitions, is provided. Significant statistical differences between average AGB are indicated by different letters between disturbance scenarios in the same PFT (uppercase letters), and between PFTs in the same disturbance scenario (lowercase letters). The letters in the last column correspond to the boxplot of each area in Figure 3.

| | CC | | CC+CF | | CC+DE | | CC+CF+DE | | Fig. 3 |
|------------|-----------|--------------------|---------------|---------------------|---------------|---------------------|-----------------|---------------------|---------------|
| PFT | n | <i>agbc</i> | 95% CI | <i>agbc</i> | 95% CI | <i>agbc</i> | 95% CI | <i>agbc</i> | 95% CI |
| DC | 7,916 | 0.03 ^{Aa} | 0.02, 0.04 | -0.48 ^{Ba} | -0.49, -0.47 | -0.11 ^{Ca} | -0.12, -0.1 | -0.55 ^{Da} | -0.56, -0.55 |
| EV | 7,916 | 0.07 ^{Ab} | 0.06, 0.08 | -0.43 ^{Bb} | -0.44, -0.42 | -0.07 ^{Cb} | -0.08, -0.07 | -0.5 ^{Db} | -0.51, -0.5 |
| NRESP | 7,916 | 0.02 ^{Ac} | 0.01, 0.03 | -0.56 ^{Bc} | -0.57, -0.55 | -0.11 ^{Cc} | -0.12, -0.1 | -0.61 ^{Dc} | -0.62, -0.61 |
| RESP | 7,916 | 0.08 ^{Ad} | 0.07, 0.09 | -0.35 ^{Bd} | -0.36, -0.34 | -0.07 ^{Cd} | -0.08, -0.07 | -0.44 ^{Dd} | -0.45, -0.43 |
| ZOO | 7,916 | 0.09 ^{Ae} | 0.08, 0.1 | -0.51 ^{Be} | -0.52, -0.5 | -0.28 ^{Ce} | -0.29, -0.27 | -0.7 ^{De} | -0.7, -0.69 |
| NZOO | 7,916 | 0.01 ^{Af} | 0, 0.02 | -0.4 ^{Bf} | -0.43, -0.39 | 0.09 ^{Cf} | 0.08, 0.1 | -0.36 ^{Df} | -0.31, -0.3 |

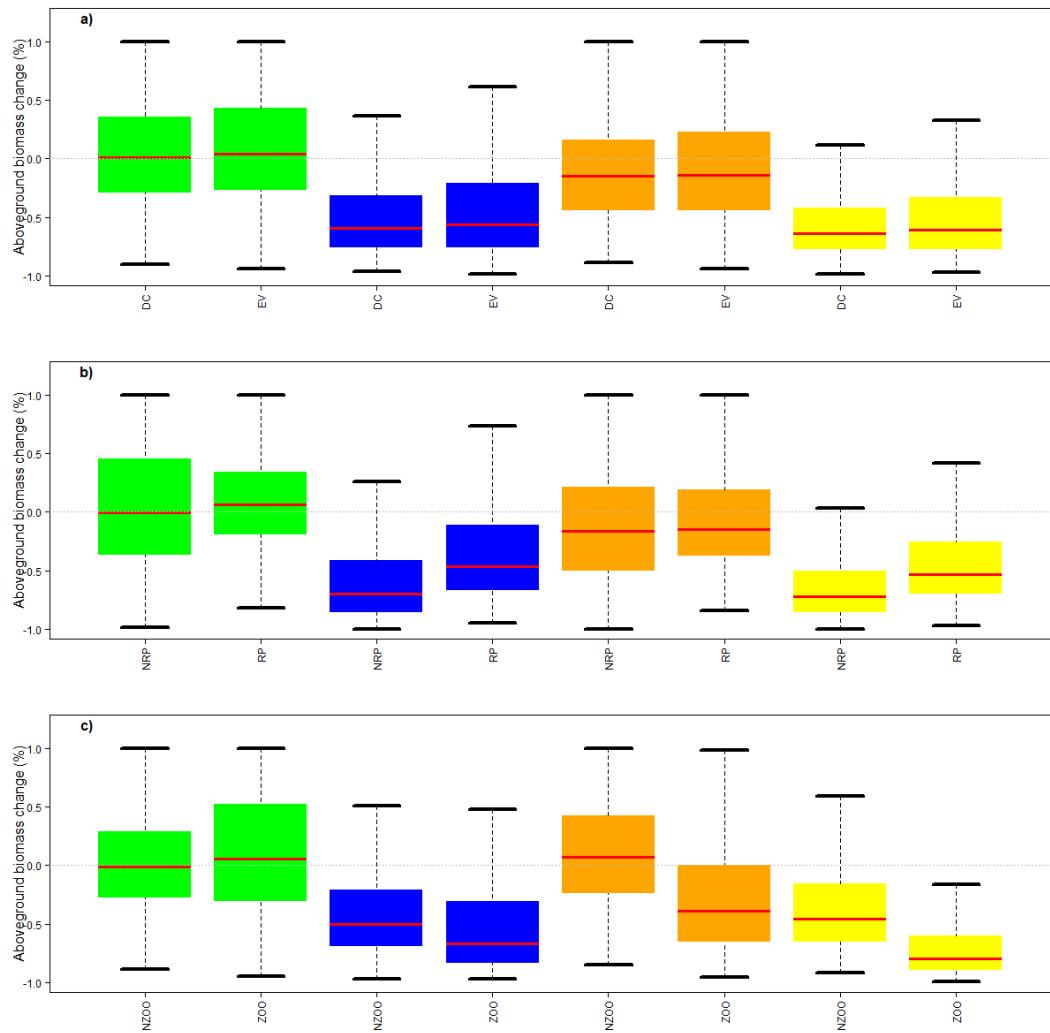


Fig. 3. Change in aboveground biomass (AGB) in six functional groups: deciduous (DC), evergreen (EV), resprouters (RESP), non-resprouters (NRESP), animal dispersed (ZOO), and non-animal dispersed (NZOO), based on a 1000-year simulation. The colors green (climate change), blue (climate change and change in fire effects), orange (climate change and defaunation), and yellow (all factors together) indicate global change scenarios. Positive and negative values indicate gain and loss of AGB relative to the baseline scenario, respectively. The red lines of each boxplot indicate the average AGB in the global change scenarios.

4. Discussion

Our study presents a significantly different trend in AGB over time in response to the synergism of changes in the climate and fire frequency and defaunation compared to the separate effects of these phenomena. When the synergistic effects of these three drivers were considered, AGB fell in all the areas of our study, ranging from 35% to 77%. As has been suggested (Sankaran, 2019), changes in savanna vegetation result not only from the occurrence of the dry season but also from the multiple drivers that work together, such as change in fire intensity and the loss of animal dispersers. To the best of our knowledge, this is the first study to consider the synergistic effects of these drivers on change in AGB on a long-term scale (1000 years) in Brazilian savannas. Our study, therefore, can contribute to closing the knowledge gap regarding longer-term atrophic effects in these savannas (Arruda et al., 2018; Gomes et al., 2018).

Different physiognomies responded differently to disturbance. Woody savanna was the most affected physiognomy, presenting the greatest AGB loss in all scenarios. Climate change and defaunation (CC.DE) decreased the AGB in woody savanna, while AGB increased in open savanna and savanna woodland. Open savanna in CC.DE conditions showed a significant increase relative to savanna woodland. Thus, the loss of seed dispersers appears to have less impact on the open savanna. Unlike open savanna, which has a dominant shrub component, the arboreal component dominates woody savannas and savanna woodland (Coutinho, 1978; Furley, 1992). Animal dispersal is significantly more frequent in the arboreal component than in the shrub component (Batalha and Mantovani, 2000; Mantovani and Martins, 1988; Pirani et al., 2009), so it makes sense that open savanna was the least affected physiognomy when defaunation was added to our model. Conversely, when fire was considered in our model, the greatest loss of AGB was experienced in the following order: woody savanna, open savanna, and woodland savanna. In this case, savanna woodland was the least affected. Savanna

woodland has a denser structure (very much like woods) than the other savanna types. These denser savannas normally lose less moisture than open savannas and woody savannas, consequently, fire disturbance affects them differently (Pereira Júnior et al., 2014).

Evergreens had a larger increase (CC) and a less decrease (other scenarios) in AGB than deciduous species. The difference in AGB loss of deciduous species was 5% more than evergreens. Specific leaf mass (SLM) was one of the contrasting parameters between evergreen (high SLM) and deciduous species (low SLM) in our model. SLM is the main predictor of sclerophyllly (Neves et al., 2010), which is an evolutionary strategy for managing resources in habitats with limited water (Lohbeck et al., 2013). Large SLM contributes to the growth rate in favorable seasons (Eamus, 1999; Hartley and Jones, 1997), with a consequent delay in ABG loss in a disturbance scenario. Our studies, therefore, support the hypothesis that sclerophyllly is a trait providing savanna species with some adaptation to changing environmental conditions (Fonseca et al., 2018; Guerra and Scremen-Dias, 2018; Lebrija-Trejos et al., 2011, 2010).

Resprouting species had greater AGB than non-resprouting species in both pre- and post-disturbance scenarios. Change in the fire regimen resulted in a loss of AGB that was 20% greater in non-resprouters compared to resprouters. Our results give support to the idea that non-resprouters are more vulnerable than resprouters to change in the fire regime (Salazar and Goldstein, 2014). Studies based on the seed bank have shown that fire plays a key role in the maintenance of the savanna's plant diversity (Abreu et al., 2017; Andrade and Miranda, 2014). However, because many species are fire-sensitive, changes in natural fire regimes are reported to cause changes in plant community structure (Salazar and Goldstein, 2014), which is in agreement with our results. We recommend that fire management should be carefully conducted, keeping fire frequency as close as possible to the natural frequency, at around four-year intervals (Coutinho, 1990; Furley et al., 2008; Pivello, 2011; Pivello and Coutinho, 1992).

The higher AGB demonstrated by animal-dispersed species in the control model was reversed after disturbance. In our study, there were PFTs dispersed by animals that lost 60% of their AGB. This represented a 30% greater AGB loss than the observed in groups that were not dispersed by animals. In our models, when the population of dispersers decreases, the chances that dispersed seeds arrive at safe sites decrease too (Herrera et al., 1994; Nathan and Muller-Landau, 2000). Thus, an extinction vortex is triggered that begins with a reduction in seed dispersers as a result of anthropic processes, leads to a decrease in plant species, and again a further decrease in savanna animals. These results reinforce the need to consider mutual interactions in savanna management as a strategy to maintain savanna ecosystem services. However, our findings must be interpreted with caution because our model did not distinguish between the functional role of vertebrates and invertebrates and, as we know, ants play an important role in the dispersal of the seeds of savannas tree species (Christianini et al., 2012; Christianini and Oliveira, 2010, 2009). In addition, responses to anthropic disturbances may differ between vertebrates and invertebrates. Therefore, further studies with more focus on different animal groups are required.

4.1 Implications of our findings

Our model investigated the response of AGB to a rise in temperature of 2° C. When this effect of climate change was considered in isolation, there was no loss of biomass in any of the areas, phytophysiognomies, or PFTs. However, when fires became more frequent, AGB losses were evident in all study areas, phytophysiognomies, and PFTs. Recently, a systematic increase in the risk of extreme fire events has been reported if global temperature increases by 2 °C in daily values (Silva et al., 2016). According to these authors, an increase of more than 1.5 °C in global temperature will result in increasingly frequent fires in the savanna. Our results show that these changes can cause huge losses in AGB and consequently in the savannas' carbon storage capacity. Therefore, it is wise that warming should be kept at a maximum of

1.5 °C by the end of the century (Silva et al., 2019) to minimize the effects of fire on the savanna AGB.

Most carbon in the savanna is stored in the belowground biomass (Bustamante et al., 2012). However, the biomass of branches, trunks, and leaves is estimated to be 50% carbon (Batlle-Bayer et al., 2010). Thus, AGB also plays an important role in atmospheric carbon storage in the savanna (Paiva et al., 2011; Roquette, 2018; Scolforo et al., 2015). When the synergism of the three factors was considered in our model, AGB losses in the ecosystems were 38%, 74%, and 39% relative to a baseline scenario, in the open savanna, woody savanna, and savanna woodland, respectively. Such AGB loss would result in higher greenhouse gas emissions. AGB losses and carbon emission in the Cerrado have mainly been attributed to the effects of habitat loss and fire (Bustamante et al., 2012; Carvalho et al., 2014). According to our model, the loss of seed dispersers also plays an important role in the loss of AGB and, consequently, in greenhouse gas emissions. The most important savanna in South America is found in Brazil, which is a signatory to global greenhouse gas reduction targets. The main strategies to achieve greenhouse gas reduction targets are focused on reducing deforestation, which is justified because deforestation is one of the largest sources of greenhouse gas emissions. However, non-deforested areas may be susceptible to fire and hunting. Thus, incorporating the analysis and intervention of different factors should be considered future challenges in the study of savanna dynamics in South America.

5. Conclusions

The most obvious finding to emerge from our study is that climate, fire, and defaunation act synergistically leading to the loss of AGB in savannas. Our study also showed that responses were different between phytobiognomies and plant functional groups. The response of physiognomies and functional groups depends on the type of disturbance

considered. In general, woody savannas were the most seriously affected among phytophysiognomies. Deciduous, non-resprouting, animal-dispersed species were the most susceptible to disturbance. This information can be used to target interventions aimed at the conservation of plants and ecosystem services in the savanna.

CONSIDERAÇÕES FINAIS

Três resultados importantes emergem desta tese. Primeiro, o déficit florístico é maior em alguns gêneros, famílias e ecorregiões das savanas da América do Sul (SAS). Segundo, mais de 40.76% das 2.203 espécies de árvores encontradas na SAS foram classificadas nas sete formas de raridade de Rabinowitz; e 74.89% dessas 2.203 espécies tinham pelo menos uma ocorrência na rede de áreas protegidas (APs). Finalmente, a mudança climática, a mudança no regime do fogo e a defaunação tiveram um efeito sinérgico na performance de grupos funcionais de plantas; em alguns casos levando a uma perda de 70% de biomassa vegetal.

O déficit florístico é maior em alguns táxons e ecorregiões da SAS. Diferente do que era esperado, esse déficit florístico é grande tanto nos gêneros e famílias mais diversos quanto nos gêneros e famílias menos diversos. Dentre os táxons mais diversos da SAS com grande déficit florístico estavam, por exemplo, as famílias Myrtaceae, Fabaceae e Malpighiaceae, e gêneros como *Myrcia*, *Eugenia* e *Psidium*. Dentre os táxons menos diversos da SAS com grande déficit florístico estavam famílias como Verbenaceae, Sapindaceae e Polygalaceae, e gêneros como *Ficus*, *Ilex* e *Tapirira*. Espacialmente, o maior déficit florístico foi observado no sentido Cerrado-Savana de Beni-Llanos e savanas da Amazônia. No Cerrado, o déficit florístico foi principalmente observado nas zonas de transições Cerrado-Amazônia e Cerrado-Caatinga. Nos Llanos e nas Savana da Amazônia, o déficit florístico parece ser localizado principalmente nas regiões mais ao norte. Sugerimos que os táxons e as regiões onde o maior déficit florístico foi observado devem ser foco para pesquisas futuras na SAS.

De modo geral, nossos resultados parecem indicar que a rede de APs da SAS tem um bom desempenho, pois, 74.89% de todas as espécies têm alguma ocorrência dentro daquelas APs. Esse resultado seria bastante otimista se não fosse o fato de que muitas das espécies que ocorrem dentro de APs apresentaram maior proporção de registros fora. De acordo com nossos resultados, 59.7% das espécies verificadas dentro de APs tinham cerca 75% dos seus registros

de ocorrência fora daquelas APs. Esses casos envolviam desde espécies comuns como *Byrsonima clausseniana* A.Juss. (~16% do total de indivíduos dentro de áreas protegidas) até espécies raras como *Campomanesia xanthocarpa* (Mart.) O. Berg. (~8% do total de indivíduos dentro de áreas protegidas). No caso de *C. xanthocarpa*, esse percentual de indivíduos dentro das APs correspondia a um dos 12 registros observados para aquela espécie. Casos como o de *C. xanthocarpa* sugerem que a rede de APs das SAS precisa ser ampliada. Embora pareça contraditório, nossos resultados revelaram que a razão entre o número de registros dentro de APs e o número total de registros por célula de grade é baixo. Também mostramos que quanto maior é a razão entre os registros dentro de APs e os registros totais, maior é a razão entre o número de espécies que ocorrem em APs e o total de espécies por uma célula de grade. Isso sugere que, se nosso esforço amostral dentro das APs aumentar, nosso conhecimento sobre a lacuna de conservação de espécies pode mudar. Assim, nossos resultados sugerem que, mais estudos dentro das APs são necessários para melhor avaliação do status de conservação das espécies da SAS.

De acordo com nossos achados, os três fatores antrópicos analisados aqui (mudança climática e do regime de incêndios e defaunação) têm um efeito sinérgico na diminuição da performance de grupos funcionais de plantas. Atualmente, tanto as estratégias de planejamento de áreas protegidas como as estratégias de manejo dentro das áreas protegidas já existentes estão focadas na ação desses fatores separadamente. Acreditamos que as ações estratégicas de planejamento e manejo dentro do contexto de mudanças globais devem levar em conta o sinergismo de diferentes fatores antrópicos.

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APÊNDICES

Supplementary material chapter 2

Table 1. Keywords used to search for papers published. Different combinations were done with the words in each column.

| Keywords 1 | Keywords 2 | Keywords 3 |
|--|--|---|
| ‘cerrado’ ‘cerradão’ | ‘Stricto sensu’ ‘rupestre’ | ‘vegetation’ |
| ‘savanna’ ‘Guiana savanna’ ‘Llanos’ Llanos of moxos’ ‘amazonia savanna’ ‘caatinga savanna’ | ‘dense’ ‘grassland’ ‘forested’ ‘tree’ ‘arboreal’ ‘hyper seasonal’ ‘steppe’ ‘woodland’ ‘parkland’ ‘eutrophic’ | ‘phytosociological’ ‘floristic’ ‘structure’ ‘community’ |

Table 2. Results of the regression analysis between relative flora shortfall (rfs) and estimated relative flora shortfall by different interpolation techniques (it) for floristic and phytosociological surveys.

| surveys | it | R-squared | Estimate | Std. Error | t value | Pr(> t) |
|-------------------|------------------|-----------|-----------|------------|---------|------------|
| floristic | Idw | 0.9113 | 1.045649 | 0.027563 | 37.937 | <2e-16 *** |
| | Spline | 0.8201 | 0.955156 | 0.037804 | 25.266 | <2e-16 *** |
| | Natural Neighbor | 0.8585 | 1.0231136 | 0.0351091 | 29.141 | <2e-16 *** |
| | Kriging | 0.5927 | 1.201880 | 0.084213 | 14.272 | <2e-16 *** |
| phytosociological | Idw | 0.9113 | 1.045649 | 0.027563 | 37.937 | <2e-16 *** |
| | Spline | 0.875 | 0.9608528 | 0.0269187 | 35.695 | <2e-16 *** |
| | Natural Neighbor | 0.8875 | 1.054924 | 0.027845 | 37.886 | <2e-16 *** |
| | Kriging | 0.6771 | 1.16106 | 0.05943 | 19.538 | <2e-16 *** |

Table 3. Papers reporting phytosociological or floristic surveys in the Neotropical savanna and they respective references.

| Literature | Reference |
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Table 4. Number of absolut flora shortfall (afs), binomials, total records (N) and relative shortfall (rfs = afs/N) found in each paper. The tables are sorted by year in a decrescent way. Each row in the table represents a survey for floristic and phytosociological studies separately.

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|-------|-----------|-----|-----|----------|-------|----------------|-----|----|----------|
| 1970 | Floristic | 10 | 26 | 0.384615 | 1982 | Phytosociology | 5 | 22 | 0.227273 |
| 1971 | Floristic | 4 | 26 | 0.153846 | 1983 | Phytosociology | 9 | 33 | 0.272727 |
| 1980 | Floristic | 14 | 120 | 0.116667 | 1983 | Phytosociology | 1 | 54 | 0.018519 |
| 1983 | Floristic | 4 | 106 | 0.037736 | 1986 | Phytosociology | 14 | 41 | 0.341463 |
| 1984 | Floristic | 2 | 14 | 0.142857 | 1986 | Phytosociology | 2 | 27 | 0.074074 |
| 1984 | Floristic | 44 | 115 | 0.382609 | 1987 | Phytosociology | 8 | 36 | 0.222222 |
| 1985 | Floristic | 6 | 55 | 0.109091 | 1993 | Phytosociology | 7 | 66 | 0.106061 |
| 1985 | Floristic | 3 | 35 | 0.085714 | 1993 | Phytosociology | 6 | 71 | 0.084507 |
| 1985 | Floristic | 15 | 81 | 0.185185 | 1993 | Phytosociology | 3 | 57 | 0.052632 |
| 1986 | Floristic | 3 | 146 | 0.020548 | 1993 | Phytosociology | 4 | 68 | 0.058824 |
| 1986 | Floristic | 2 | 93 | 0.021505 | 1993 | Phytosociology | 3 | 58 | 0.051724 |
| 1986 | Floristic | 4 | 121 | 0.033058 | 1993 | Phytosociology | 9 | 68 | 0.132353 |
| 1986 | Floristic | 1 | 112 | 0.008929 | 1994 | Phytosociology | 1 | 23 | 0.043478 |
| 1987 | Floristic | 3 | 31 | 0.096774 | 1994 | Phytosociology | 5 | 30 | 0.166667 |
| 1988 | Floristic | 5 | 51 | 0.098039 | 1994 | Phytosociology | 1 | 21 | 0.047619 |
| 1988 | Floristic | 11 | 79 | 0.139241 | 1994 | Phytosociology | 2 | 25 | 0.08 |
| 1988 | Floristic | 1 | 54 | 0.018519 | 1994 | Phytosociology | 1 | 24 | 0.041667 |
| 1988 | Floristic | 8 | 101 | 0.079208 | 1994 | Phytosociology | 17 | 69 | 0.246377 |
| 1989 | Floristic | 7 | 34 | 0.205882 | 1994 | Phytosociology | 6 | 36 | 0.166667 |
| 1989 | Floristic | 1 | 21 | 0.047619 | 1994 | Phytosociology | 4 | 37 | 0.108108 |
| 1989 | Floristic | 9 | 67 | 0.134328 | 1994 | Phytosociology | 2 | 26 | 0.076923 |
| 1989 | Floristic | 5 | 27 | 0.185185 | 1994 | Phytosociology | 3 | 27 | 0.111111 |
| 1989 | Floristic | 6 | 121 | 0.049587 | 1994 | Phytosociology | 10 | 68 | 0.147059 |
| 1990 | Floristic | 11 | 51 | 0.215686 | 1995 | Phytosociology | 9 | 58 | 0.155172 |
| 1990 | Floristic | 1 | 50 | 0.02 | 1996 | Phytosociology | 6 | 41 | 0.146341 |
| 1992 | Floristic | 2 | 27 | 0.074074 | 1997 | Phytosociology | 14 | 41 | 0.341463 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| 1992 | Floristic | 3 | 34 | 0.088235 | 1997 | Phytosociology | 3 | 41 | 0.073171 |
| 1993 | Floristic | 1 | 19 | 0.052632 | 1997 | Phytosociology | 3 | 46 | 0.065217 |
| 1995 | Floristic | 4 | 61 | 0.065574 | 1997 | Phytosociology | 1 | 33 | 0.030303 |
| 1996 | Floristic | 1 | 151 | 0.006623 | 1997 | Phytosociology | 2 | 40 | 0.05 |
| 1997 | Floristic | 1 | 29 | 0.034483 | 1997 | Phytosociology | 1 | 34 | 0.029412 |
| 1997 | Floristic | 2 | 15 | 0.133333 | 1997 | Phytosociology | 1 | 41 | 0.02439 |
| 1997 | Floristic | 7 | 28 | 0.25 | 1997 | Phytosociology | 4 | 42 | 0.095238 |
| 1997 | Floristic | 4 | 22 | 0.181818 | 1997 | Phytosociology | 1 | 39 | 0.025641 |
| 1997 | Floristic | 2 | 9 | 0.222222 | 1997 | Phytosociology | 2 | 39 | 0.051282 |
| 1997 | Floristic | 1 | 22 | 0.045455 | 1997 | Phytosociology | 1 | 38 | 0.026316 |
| 1997 | Floristic | 2 | 18 | 0.111111 | 1997 | Phytosociology | 4 | 43 | 0.093023 |
| 1997 | Floristic | 1 | 7 | 0.142857 | 1997 | Phytosociology | 2 | 40 | 0.05 |
| 1997 | Floristic | 3 | 11 | 0.272727 | 1997 | Phytosociology | 1 | 35 | 0.028571 |
| 1997 | Floristic | 3 | 152 | 0.019737 | 1997 | Phytosociology | 4 | 45 | 0.088889 |
| 2000 | Floristic | 2 | 28 | 0.071429 | 1997 | Phytosociology | 3 | 43 | 0.069767 |
| 2001 | Floristic | 1 | 90 | 0.011111 | 1997 | Phytosociology | 1 | 45 | 0.022222 |
| 2001 | Floristic | 3 | 73 | 0.041096 | 1997 | Phytosociology | 1 | 25 | 0.04 |
| 2001 | Floristic | 1 | 90 | 0.011111 | 1997 | Phytosociology | 1 | 11 | 0.090909 |
| 2001 | Floristic | 2 | 85 | 0.023529 | 1997 | Phytosociology | 1 | 5 | 0.2 |
| 2001 | Floristic | 5 | 73 | 0.068493 | 1997 | Phytosociology | 1 | 10 | 0.1 |
| 2001 | Floristic | 1 | 77 | 0.012987 | 1997 | Phytosociology | 2 | 11 | 0.181818 |
| 2001 | Floristic | 4 | 94 | 0.042553 | 1997 | Phytosociology | 1 | 6 | 0.166667 |
| 2001 | Floristic | 3 | 86 | 0.034884 | 1998 | Phytosociology | 2 | 18 | 0.111111 |
| 2001 | Floristic | 2 | 88 | 0.022727 | 1998 | Phytosociology | 2 | 32 | 0.0625 |
| 2001 | Floristic | 1 | 87 | 0.011494 | 2000 | Phytosociology | 1 | 158 | 0.006329 |
| 2001 | Floristic | 1 | 74 | 0.013514 | 2000 | Phytosociology | 6 | 33 | 0.181818 |
| 2001 | Floristic | 3 | 43 | 0.069767 | 2000 | Phytosociology | 23 | 92 | 0.25 |
| 2001 | Floristic | 2 | 63 | 0.031746 | 2000 | Phytosociology | 21 | 74 | 0.283784 |
| 2001 | Floristic | 1 | 47 | 0.021277 | 2000 | Phytosociology | 14 | 65 | 0.215385 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| 2001 | Floristic | 1 | 44 | 0.022727 | 2000 | Phytosociology | 8 | 52 | 0.153846 |
| 2001 | Floristic | 1 | 57 | 0.017544 | 2000 | Phytosociology | 5 | 22 | 0.227273 |
| 2001 | Floristic | 1 | 53 | 0.018868 | 2001 | Phytosociology | 6 | 64 | 0.09375 |
| 2001 | Floristic | 4 | 65 | 0.061538 | 2001 | Phytosociology | 1 | 67 | 0.014925 |
| 2001 | Floristic | 2 | 61 | 0.032787 | 2001 | Phytosociology | 3 | 67 | 0.044776 |
| 2001 | Floristic | 1 | 67 | 0.014925 | 2001 | Phytosociology | 1 | 17 | 0.058824 |
| 2001 | Floristic | 1 | 44 | 0.022727 | 2001 | Phytosociology | 11 | 40 | 0.275 |
| 2001 | Floristic | 2 | 57 | 0.035088 | 2001 | Phytosociology | 16 | 42 | 0.380952 |
| 2001 | Floristic | 1 | 64 | 0.015625 | 2001 | Phytosociology | 3 | 93 | 0.032258 |
| 2001 | Floristic | 1 | 52 | 0.019231 | 2001 | Phytosociology | 3 | 76 | 0.039474 |
| 2001 | Floristic | 1 | 68 | 0.014706 | 2002 | Phytosociology | 2 | 63 | 0.031746 |
| 2001 | Floristic | 2 | 63 | 0.031746 | 2002 | Phytosociology | 8 | 59 | 0.135593 |
| 2001 | Floristic | 3 | 72 | 0.041667 | 2002 | Phytosociology | 9 | 58 | 0.155172 |
| 2001 | Floristic | 5 | 68 | 0.073529 | 2002 | Phytosociology | 11 | 72 | 0.152778 |
| 2001 | Floristic | 1 | 29 | 0.034483 | 2002 | Phytosociology | 13 | 69 | 0.188406 |
| 2001 | Floristic | 2 | 75 | 0.026667 | 2002 | Phytosociology | 1 | 44 | 0.022727 |
| 2001 | Floristic | 1 | 58 | 0.017241 | 2002 | Phytosociology | 3 | 38 | 0.078947 |
| 2001 | Floristic | 1 | 70 | 0.014286 | 2002 | Phytosociology | 3 | 42 | 0.071429 |
| 2001 | Floristic | 2 | 66 | 0.030303 | 2002 | Phytosociology | 3 | 73 | 0.041096 |
| 2001 | Floristic | 2 | 69 | 0.028986 | 2002 | Phytosociology | 23 | 61 | 0.377049 |
| 2001 | Floristic | 3 | 59 | 0.050847 | 2002 | Phytosociology | 10 | 27 | 0.37037 |
| 2001 | Floristic | 1 | 58 | 0.017241 | 2002 | Phytosociology | 7 | 59 | 0.118644 |
| 2001 | Floristic | 4 | 98 | 0.040816 | 2002 | Phytosociology | 7 | 52 | 0.134615 |
| 2001 | Floristic | 3 | 59 | 0.050847 | 2002 | Phytosociology | 17 | 86 | 0.197674 |
| 2001 | Floristic | 3 | 47 | 0.06383 | 2002 | Phytosociology | 17 | 72 | 0.236111 |
| 2001 | Floristic | 4 | 55 | 0.072727 | 2002 | Phytosociology | 9 | 67 | 0.134328 |
| 2001 | Floristic | 3 | 26 | 0.115385 | 2003 | Phytosociology | 4 | 41 | 0.097561 |
| 2001 | Floristic | 2 | 35 | 0.057143 | 2003 | Phytosociology | 17 | 51 | 0.333333 |
| 2001 | Floristic | 2 | 26 | 0.076923 | 2003 | Phytosociology | 18 | 60 | 0.3 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| 2001 | Floristic | 2 | 41 | 0.04878 | 2003 | Phytosociology | 44 | 100 | 0.44 |
| 2001 | Floristic | 2 | 63 | 0.031746 | 2003 | Phytosociology | 14 | 58 | 0.241379 |
| 2001 | Floristic | 1 | 49 | 0.020408 | 2003 | Phytosociology | 2 | 87 | 0.022989 |
| 2001 | Floristic | 5 | 76 | 0.065789 | 2003 | Phytosociology | 10 | 88 | 0.113636 |
| 2001 | Floristic | 1 | 77 | 0.012987 | 2003 | Phytosociology | 6 | 63 | 0.095238 |
| 2001 | Floristic | 3 | 54 | 0.055556 | 2003 | Phytosociology | 5 | 65 | 0.076923 |
| 2001 | Floristic | 1 | 24 | 0.041667 | 2003 | Phytosociology | 12 | 48 | 0.25 |
| 2001 | Floristic | 1 | 81 | 0.012346 | 2003 | Phytosociology | 23 | 76 | 0.302632 |
| 2001 | Floristic | 1 | 74 | 0.013514 | 2003 | Phytosociology | 11 | 59 | 0.186441 |
| 2001 | Floristic | 1 | 54 | 0.018519 | 2003 | Phytosociology | 24 | 54 | 0.444444 |
| 2001 | Floristic | 3 | 68 | 0.044118 | 2003 | Phytosociology | 15 | 42 | 0.357143 |
| 2001 | Floristic | 2 | 84 | 0.02381 | 2003 | Phytosociology | 1 | 74 | 0.013514 |
| 2001 | Floristic | 4 | 74 | 0.054054 | 2004 | Phytosociology | 2 | 18 | 0.111111 |
| 2001 | Floristic | 1 | 54 | 0.018519 | 2004 | Phytosociology | 23 | 92 | 0.25 |
| 2001 | Floristic | 1 | 74 | 0.013514 | 2004 | Phytosociology | 28 | 107 | 0.261682 |
| 2001 | Floristic | 1 | 84 | 0.011905 | 2004 | Phytosociology | 14 | 42 | 0.333333 |
| 2001 | Floristic | 3 | 80 | 0.0375 | 2004 | Phytosociology | 32 | 63 | 0.507937 |
| 2001 | Floristic | 1 | 76 | 0.013158 | 2004 | Phytosociology | 31 | 65 | 0.476923 |
| 2001 | Floristic | 2 | 83 | 0.024096 | 2004 | Phytosociology | 1 | 57 | 0.017544 |
| 2001 | Floristic | 2 | 90 | 0.022222 | 2005 | Phytosociology | 5 | 12 | 0.416667 |
| 2001 | Floristic | 1 | 72 | 0.013889 | 2005 | Phytosociology | 12 | 58 | 0.206897 |
| 2001 | Floristic | 1 | 83 | 0.012048 | 2005 | Phytosociology | 4 | 50 | 0.08 |
| 2001 | Floristic | 2 | 80 | 0.025 | 2005 | Phytosociology | 20 | 118 | 0.169492 |
| 2001 | Floristic | 1 | 82 | 0.012195 | 2005 | Phytosociology | 10 | 46 | 0.217391 |
| 2001 | Floristic | 2 | 90 | 0.022222 | 2005 | Phytosociology | 8 | 60 | 0.133333 |
| 2001 | Floristic | 1 | 91 | 0.010989 | 2005 | Phytosociology | 8 | 73 | 0.109589 |
| 2001 | Floristic | 2 | 55 | 0.036364 | 2005 | Phytosociology | 24 | 131 | 0.183206 |
| 2001 | Floristic | 1 | 82 | 0.012195 | 2005 | Phytosociology | 1 | 77 | 0.012987 |
| 2001 | Floristic | 2 | 72 | 0.027778 | 2005 | Phytosociology | 1 | 77 | 0.012987 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| 2001 | Floristic | 1 | 60 | 0.016667 | 2005 | Phytosociology | 3 | 15 | 0.2 |
| 2001 | Floristic | 3 | 80 | 0.0375 | 2005 | Phytosociology | 10 | 55 | 0.181818 |
| 2001 | Floristic | 3 | 67 | 0.044776 | 2005 | Phytosociology | 13 | 78 | 0.166667 |
| 2001 | Floristic | 2 | 91 | 0.021978 | 2005 | Phytosociology | 13 | 62 | 0.209677 |
| 2001 | Floristic | 3 | 78 | 0.038462 | 2005 | Phytosociology | 5 | 55 | 0.090909 |
| 2002 | Floristic | 28 | 85 | 0.329412 | 2005 | Phytosociology | 13 | 66 | 0.19697 |
| 2004 | Floristic | 22 | 114 | 0.192982 | 2005 | Phytosociology | 9 | 52 | 0.173077 |
| 2005 | Floristic | 3 | 38 | 0.078947 | 2005 | Phytosociology | 11 | 49 | 0.22449 |
| 2006 | Floristic | 2 | 12 | 0.166667 | 2005 | Phytosociology | 18 | 66 | 0.272727 |
| 2006 | Floristic | 6 | 91 | 0.065934 | 2005 | Phytosociology | 7 | 30 | 0.233333 |
| 2006 | Floristic | 18 | 86 | 0.209302 | 2005 | Phytosociology | 1 | 27 | 0.037037 |
| 2006 | Floristic | 3 | 63 | 0.047619 | 2005 | Phytosociology | 3 | 71 | 0.042254 |
| 2006 | Floristic | 1 | 130 | 0.007692 | 2006 | Phytosociology | 1 | 56 | 0.017857 |
| 2007 | Floristic | 27 | 54 | 0.5 | 2006 | Phytosociology | 3 | 50 | 0.06 |
| 2007 | Floristic | 11 | 28 | 0.392857 | 2006 | Phytosociology | 4 | 49 | 0.081633 |
| 2007 | Floristic | 5 | 37 | 0.135135 | 2006 | Phytosociology | 22 | 69 | 0.318841 |
| 2007 | Floristic | 5 | 30 | 0.166667 | 2006 | Phytosociology | 51 | 106 | 0.481132 |
| 2007 | Floristic | 3 | 26 | 0.115385 | 2006 | Phytosociology | 14 | 27 | 0.518519 |
| 2007 | Floristic | 6 | 29 | 0.206897 | 2006 | Phytosociology | 32 | 59 | 0.542373 |
| 2007 | Floristic | 7 | 44 | 0.159091 | 2006 | Phytosociology | 20 | 39 | 0.512821 |
| 2007 | Floristic | 2 | 15 | 0.133333 | 2006 | Phytosociology | 19 | 43 | 0.44186 |
| 2008 | Floristic | 10 | 86 | 0.116279 | 2007 | Phytosociology | 3 | 42 | 0.071429 |
| 2008 | Floristic | 18 | 132 | 0.136364 | 2007 | Phytosociology | 14 | 34 | 0.411765 |
| 2008 | Floristic | 4 | 21 | 0.190476 | 2007 | Phytosociology | 2 | 38 | 0.052632 |
| 2008 | Floristic | 11 | 33 | 0.333333 | 2007 | Phytosociology | 1 | 56 | 0.017857 |
| 2008 | Floristic | 2 | 83 | 0.024096 | 2007 | Phytosociology | 14 | 39 | 0.358974 |
| 2008 | Floristic | 2 | 83 | 0.024096 | 2007 | Phytosociology | 11 | 35 | 0.314286 |
| 2008 | Floristic | 1 | 44 | 0.022727 | 2007 | Phytosociology | 26 | 72 | 0.361111 |
| 2008 | Floristic | 2 | 50 | 0.04 | 2007 | Phytosociology | 9 | 92 | 0.097826 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| 2009 | Floristic | 5 | 44 | 0.113636 | 2007 | Phytosociology | 2 | 63 | 0.031746 |
| 2009 | Floristic | 7 | 48 | 0.145833 | 2007 | Phytosociology | 2 | 22 | 0.090909 |
| 2012 | Floristic | 2 | 35 | 0.057143 | 2007 | Phytosociology | 9 | 44 | 0.204545 |
| 2012 | Floristic | 1 | 9 | 0.111111 | 2007 | Phytosociology | 5 | 32 | 0.15625 |
| 2012 | Floristic | 1 | 63 | 0.015873 | 2007 | Phytosociology | 8 | 42 | 0.190476 |
| 2013 | Floristic | 10 | 109 | 0.091743 | 2007 | Phytosociology | 4 | 47 | 0.085106 |
| 2015 | Floristic | 4 | 70 | 0.057143 | 2007 | Phytosociology | 4 | 50 | 0.08 |
| 2015 | Floristic | 2 | 50 | 0.04 | 2007 | Phytosociology | 10 | 35 | 0.285714 |
| 2015 | Floristic | 2 | 41 | 0.04878 | 2007 | Phytosociology | 11 | 62 | 0.177419 |
| NA | Floristic | 2 | 59 | 0.033898 | 2008 | Phytosociology | 31 | 67 | 0.462687 |
| NA | Floristic | 5 | 84 | 0.059524 | 2008 | Phytosociology | 4 | 52 | 0.076923 |
| NA | Floristic | 2 | 61 | 0.032787 | 2008 | Phytosociology | 1 | 95 | 0.010526 |
| NA | Floristic | 2 | 64 | 0.03125 | 2008 | Phytosociology | 1 | 69 | 0.014493 |
| NA | Floristic | 3 | 105 | 0.028571 | 2008 | Phytosociology | 1 | 101 | 0.009901 |
| NA | Floristic | 2 | 78 | 0.025641 | 2008 | Phytosociology | 8 | 90 | 0.088889 |
| NA | Floristic | 2 | 85 | 0.023529 | 2008 | Phytosociology | 6 | 93 | 0.064516 |
| NA | Floristic | 2 | 58 | 0.034483 | 2008 | Phytosociology | 17 | 69 | 0.246377 |
| NA | Floristic | 2 | 60 | 0.033333 | 2008 | Phytosociology | 13 | 68 | 0.191176 |
| NA | Floristic | 4 | 80 | 0.05 | 2008 | Phytosociology | 10 | 35 | 0.285714 |
| | | | | | 2008 | Phytosociology | 23 | 51 | 0.45098 |
| | | | | | 2008 | Phytosociology | 14 | 59 | 0.237288 |
| | | | | | 2008 | Phytosociology | 6 | 49 | 0.122449 |
| | | | | | 2008 | Phytosociology | 15 | 55 | 0.272727 |
| | | | | | 2008 | Phytosociology | 20 | 61 | 0.327869 |
| | | | | | 2009 | Phytosociology | 6 | 53 | 0.113208 |
| | | | | | 2009 | Phytosociology | 9 | 44 | 0.204545 |
| | | | | | 2009 | Phytosociology | 14 | 58 | 0.241379 |
| | | | | | 2009 | Phytosociology | 3 | 43 | 0.069767 |
| | | | | | 2009 | Phytosociology | 19 | 67 | 0.283582 |

| Years | Study | afs | N | rfs | Years | Study | afs | N | rfs |
|--------------|--------------|------------|----------|------------|--------------|----------------|------------|----------|------------|
| | | | | | 2011 | Phytosociology | 5 | 59 | 0.084746 |
| | | | | | 2011 | Phytosociology | 6 | 98 | 0.061224 |
| | | | | | 2011 | Phytosociology | 7 | 82 | 0.085366 |
| | | | | | 2011 | Phytosociology | 14 | 52 | 0.269231 |
| | | | | | 2012 | Phytosociology | 3 | 48 | 0.0625 |
| | | | | | 2012 | Phytosociology | 11 | 117 | 0.094017 |
| | | | | | 2012 | Phytosociology | 4 | 96 | 0.041667 |
| | | | | | 2012 | Phytosociology | 29 | 91 | 0.318681 |
| | | | | | 2012 | Phytosociology | 1 | 50 | 0.02 |
| | | | | | 2012 | Phytosociology | 1 | 54 | 0.018519 |
| | | | | | 2012 | Phytosociology | 2 | 71 | 0.028169 |
| | | | | | 2012 | Phytosociology | 1 | 76 | 0.013158 |
| | | | | | 2012 | Phytosociology | 4 | 54 | 0.074074 |
| | | | | | 2013 | Phytosociology | 4 | 92 | 0.043478 |
| | | | | | 2013 | Phytosociology | 2 | 80 | 0.025 |
| | | | | | 2014 | Phytosociology | 2 | 97 | 0.020619 |
| | | | | | 2014 | Phytosociology | 9 | 50 | 0.18 |
| | | | | | 2014 | Phytosociology | 11 | 35 | 0.314286 |
| | | | | | 2015 | Phytosociology | 7 | 58 | 0.12069 |
| | | | | | 2015 | Phytosociology | 13 | 73 | 0.178082 |
| | | | | | 2015 | Phytosociology | 5 | 91 | 0.054945 |
| | | | | | 2015 | Phytosociology | 3 | 90 | 0.033333 |
| | | | | | 2015 | Phytosociology | 2 | 76 | 0.026316 |

Table 5. The number of surveys (surveys) across ecoregions according to Olson et al. (2001) for floristic (flora) and phytosociological (phyto).

| Study | Ecoregion Olson et al. 2001 | surveys |
|-------|--------------------------------|---------|
| Flora | Alto Paraná Atlantic forests | 3 |
| Flora | Araucaria moist forests | 1 |
| Flora | Atlantic Coast restingas | 2 |
| Flora | Atlantic dry forests | 1 |
| Flora | Caatinga | 4 |
| Flora | Cerrado | 104 |
| Flora | Guiana savanna | 10 |
| Flora | Gurupa varzeá | 1 |
| Flora | Llanos | 2 |
| Flora | Madeira-Tapajós moist forests | 11 |
| Flora | Maranhão Babacu forests | 4 |
| Flora | Mato Grosso seasonal forests | 9 |
| Flora | Pantanal | 2 |
| Flora | Pernambuco interior forests | 1 |
| Flora | Purus madeira | 1 |
| Flora | Uatumá-Trombetas moist forests | 5 |
| Phyto | Amazonas-Orinoco | 2 |
| Phyto | Araucaria moist forest | 2 |
| Phyto | Atlantic dry forest | 2 |
| Phyto | Caatinga | 26 |
| Phyto | Cerrado | 97 |
| Phyto | Chiquitano Dry-forest | 1 |
| Phyto | Guiana savanna | 5 |
| Phyto | Llanos | 2 |
| Phyto | Madeira Tapajos moist forest | 3 |
| Phyto | Maranhão Babacu | 48 |
| Phyto | Mato Grosso seasonal forest | 1 |
| Phyto | Pantanal | 1 |
| Phyto | Pernambuco interior forest | 3 |
| Phyto | Purus madeira | 1 |

Table 6. Parameter estimates of the radon intercept and slope SMA model applied to the description of the relative flora shortfall as a function of the species richness of floristic (n=161) and phytosociological (n=194) surveys. The confidence interval (CI) and the p-value at a 5% level of significance are presented.

| | y Intercept β_0 | | | Slope β_1 | | | |
|-------------------|---|-------------------------|-----------------|-----------------------------------|----------------|----------------------|--|
| model | Estimate | CI (2.5 ±97.5 %) | Estimate | CI (2.5 ±97.5 %) | p-value | R² | |
| floristic | -7.52 | -9.40± -5.90 | 0.18 | 0.15±0.21 | 0.006 | 0.04 | |
| phytosociological | -10.68 | -13.43±-8.28 | 0.34 | -8.28±0.30 | 0.009 | 0.12 | |

Table 7. Parameter estimates of the radon intercept and slope GLM model (*quasi-Poisson* distribution and logit link) applied to the description of the relative flora shortfall as a function of the time (years) of the floristic (n=161) and phytosociological (n=194) surveys. The confidence interval (CI) and the p-value at a 5% level of significance are presented. The standard error (SE) and the Pr(>|t|) at a 5% level of significance.

| | y Intercept β_0 | | | | Slope β_1 | | |
|-------------------|-----------------------|-----------|--------------------|--|-----------------|-----------|--------------------|
| model | Estimate | SE | Pr(> t) | | Estimate | SE | Pr(> t) |
| floristic | 39.14716 | 24.13353 | 0.1069 | | -0.02082 | 0.01208 | 0.0867 |
| phytosociological | -24.32945 | 21.93874 | 0.269 | | 0.01131 | 0.01095 | 0.303 |

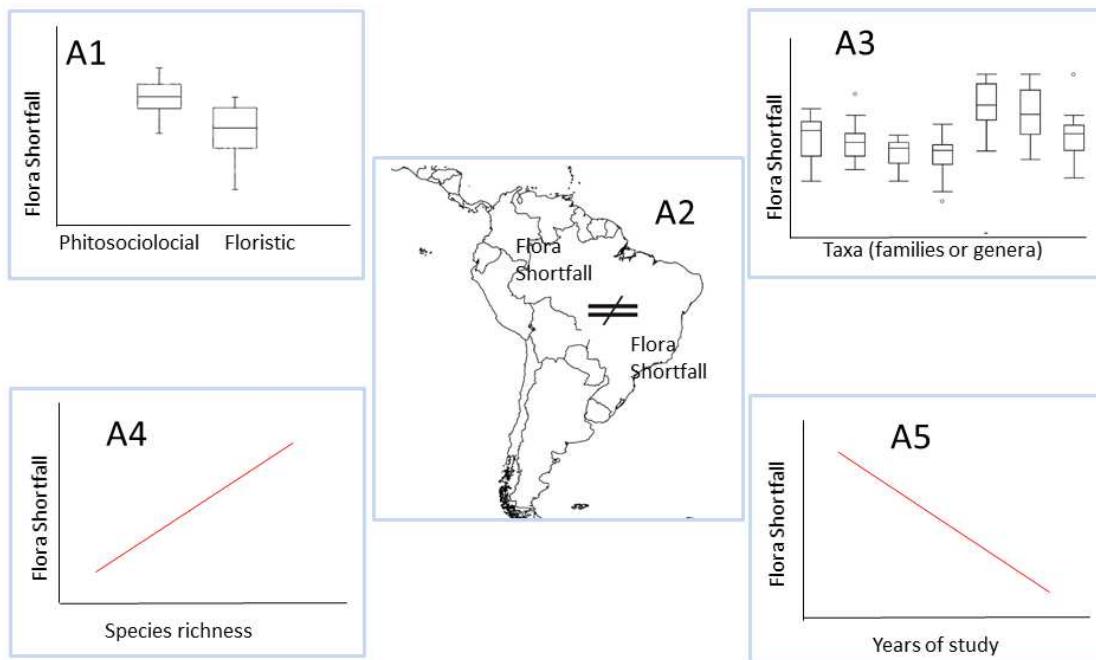


Fig. 1. Scheme assumed to classify flora shortfall. The number in each box corresponds to the question number in the introduction (for example question 1 in the introduction is equal to A1 in the box).

Supplementary material chapter 3

Additional information.

To quantify the forms of rarity in the ESAS, we used the NeoTropSavanna database, as we were interested in verifying farther geographic range intervals, the ecoregion number, and the number of grid cells that species occupy.

Geographic range - To quantify the geographic range, we created a matrix with the points of occurrence of all species. We then used the function *frequency* of the MASS package in R (R CoreTeam, 2019) to create a matrix of species occurrence and latitudinal belts.

Ecoregion number - To quantify the species affinity for the habitat, we used a layer of the ecoregions in matrix form (Olson et al., 2001). We used the *raster dataset* tool of the Data Management toolbox in ArcGIS to convert the polygonal ecoregions (Olson et al., 2001) to a raster form with a 0.005 cell size. We created a vector layer of the savanna's rare species and used the extraction tool of the spatial analyst (ESRI, 2011) to extract information about the ecoregions where each species occurred. Then, to extract information on where species occur, we used the *extraction* function in ArcGIS (ESRI, 2011) and generated one matrix with all species and ecoregions. Then, we quantified the number of ecoregions in which species occurred.

Population size - We used Hawth's Analysis Tools (ESRI, 2011) to create a grid of cells in the ESAS polygon. We used the function *Create Mosaic Dataset* on Arcgis to create a vector grid of the SAS and recorded species occurrence in 331 of the total grid cells. Species with one or two individuals sampled per grid cell were considered rare or as having a scarce population. We used the *union* function of the Overlay tool to combine species occurrence and grid cells and generated a data matrix of species and the number of grid cells they occurred in. Then, the number of occurrences that each species has in each grid was considered the local population.

Network protected area - To check the protection level of SAS species, we consulted the World Database on Protected Areas (WDPA), an up-to-date database of protected areas in the world (UNEP-WCMC and IUCN, 2019). We created a vector layer with all SAS species (rare and common) and their geographical location, then used the extraction tool function to extract the values of the cells into two matrix layers (ESRI, 2011).

Table 1. Keywords used to search for papers published. Different combinations were done with the words in each column.

| Keywords 1 | Keywords 2 | Keywords 3 |
|--------------------------|------------------------------|-------------------------|
| ‘cerrado’ ‘cerradão’ | ‘Stricto sensu’ ‘rupestre’ | ‘vegetation’ |
| ‘savanna’ ‘Guiana | ‘dense’ ‘grassland’ | ‘phytosociological’ |
| savanna’ ‘Llanos’ Llanos | ‘forested’ ‘tree’ ‘arboreal’ | ‘floristic’ ‘structure’ |
| of moxos’ ‘amazonia | ‘hyper seasonal’ ‘steppe’ | ‘community’ |
| savanna’ ‘caatinga | ‘woodland’ ‘parkland’ | |
| savanna’ | ‘eutrophic’ | |

Table 2. Literature reporting phytosociological or floristic surveys (n=104) in the Neotropical savanna and they respective references.

| Literature | Reference |
|------------|---|
| Annals | Ribeiro, J. F., & Haridasan, M. (1984). Comparação fitossociológica de um cerrado denso e um cerradão em solos distróficos no Distrito Federal. In Congresso nacional de botanica (Vol. 35, pp. 342-347). Manaus: Sociedade de Botânica do Brasil. |
| Annals | Fonseca, E. M., Ferreira, M.A., Nunes, J.R.S., Pinho, N.G.C.; Ferraz, L., Macedo, M., Guarim Neto, G. (2004). Aspectos fitossociológicos de uma comunidade de Carvoal (<i>Callisthene fasciculata</i>) no pantanal de Mato Grosso, Brasil. Simpósio sobre Recursos Naturais e Socioeconômicos do Pantanal, 229-243. |
| Annals | Rodrigues, W. A. (1971). Plantas dos Campos do Rio Branco (Território de Roraima). In III (Ie Terceiro) Simposio sobre o Cerrado. |
| Annals | Castro, A. A. J. F. (1984) Vegetação e flora da Estação Ecológica de Uruçuí-Una (resultados preliminares). In: Annals of 34º National Congress of Botany, Porto Alegre, 2, 251-261. |
| Article | Ribeiro-Silva, S., Medeiros, M. D., Gomes, B. M., Seixas, E. N. C., & Silva, M. A. P. (2012). Angiosperms from the Araripe national forest, Ceará, brazil. Check List, 8(4), 744-751. |
| Article | Ratter, J. A., Bridgewater, S., & Ribeiro, J. F. (2001). Espécies lenhosas da fitofisionomia cerrado sentido amplo em 170 localidades do bioma cerrado. Boletim do Herbário Ezequias Paulo Heringer, 7(1), 5-112. |
| Article | Sinimbu, G. (2008). Fitossociologia em Cerrado Sentido Restrito na Floresta Nacional de Brasília-DF. Revista Brasileira de Biociências, 5(S2), pg-1183. |
| Article | Felfili, J.M., et al. (1993). Análise Comparativa da Florística e Fitossociologia da Vegetação Arbórea do Cerrado sensu stricto na Chapada Pratinha, DF, Brasil. Acta Botanica Brasilica, 6(2): 27-46. |
| Article | Andrade, L. A., Felfili, J. M., & Violatti, L. (2002). Phytosociology of an area of "cerrado denso" at the RECOR-IBGE, Brasília-DF, Brazil. Acta Botanica Brasilica, 16(2), 225-240. |
| Article | Ribeiro, J. F., Silva, J. C. S., & Batmanian, G. J. (1985). Fitossociologia de tipos fisionômicos de cerrado em Planaltina-DF. Revista Brasileira de Botânica, 8(2), 131-142. |
| Article | Araujo Neto, M. D., Furley, P. A., Haridasan, M., & Johnson, C. E. (1986). The murundus of the cerrado region of Central Brazil. Journal of Tropical Ecology, 2(01), 17-35. |
| Article | Moura, I. I. O., Klein, V. L. G., Felfili, J. M., & Ferreira, H. D. (2007). Fitossociologia da comunidade lenhosa de uma área de cerrado rupestre no Parque Estadual dos Pireneus, Pirenópolis, Goiás. Revista de Biologia Neotropical, 4(2), 18. |
| Article | Borges, H. B. N., & Shepherd, G. J. (2005). Flora e estrutura do estrato lenhoso numa comunidade de Cerrado em Santo Antônio do Leverger, MT, Brasil. Revista Brasileira de Botânica, 28(1), 61-74. |

| Literature | Reference |
|------------|--|
| Article | Felfili, J. M., Nogueira, P.E., Silva, Silva Júnior, M. C. D., Marimon, B. S., & Delitti, W. B. C. (2002). Composição florística e fitossociologia do cerrado sentido restrito no município de Água Boa-MT. <i>Acta Botanica Brasilica</i> , 16(1), 103-112. |
| Article | Marimon, B.S., Lima, E.D.S., Duarte, T.G., Chieregatto, L.C., & Ratter, J. (2006). Observations on the vegetation of Northeastern Mato Grosso, Brazil. IV. Na analysis of the Cerrado–Amazonian forest ecotone. <i>Edinburgh Journal of Botany</i> , 63(2-3), 323-341. |
| Article | Nascimento, M. T., & Saddi, N. (1992). Structure and floristic composition in an area of cerrado in Cuiabá-MT, Brazil. <i>Revista Brasileira Botânica</i> , 15(1), 47-55. |
| Article | Guarim, V. L. M. S., Moraes, E. C. C., Prance, G. T., & Ratter, J. A. (2000). Inventory of a mesotrophic Callisthene cerradão in the Pantanal of Mato Grosso, Brazil. <i>Edinburgh Journal of Botany</i> , 57(03), 429-436. |
| Article | Marimon-Junior, B., & Haridasan, M. (2005). Comparação da vegetação arbórea e características edáficas de um cerradão e um cerrado sensu stricto em áreas adjacentes sobre solo distrófico no leste de Mato Grosso, Brasil. <i>Acta Botanica Brasilica</i> , 19(4), 913-926. |
| Article | Prance, G. T., & Schaller, G. B. (1982). Preliminary study of some vegetation types of the Pantanal, Mato Grosso, Brazil. <i>Brittonia</i> , 34(2), 228-251. |
| Article | Oliveira-Filho, A. T., Martins, F.R. (1986). Distribuicao, caracterizacao e composicao floristica das formacoes vegetais da regiao da Salgadeira, na Chapada dos Guimaraes (MT). <i>Revista Brasileira de Botânica</i> , 9(2), 2007-223. |
| Article | Costa, I. R. D., & Araújo, F. S. D. (2007). Organização comunitária de um encrave de cerrado sensu stricto no bioma Caatinga, chapada do Araripe, Barbalha, Ceará. <i>Acta Botanica Brasilica</i> , 21(2), 281-291. |
| Article | Medeiros, M. B., Walter, B. M. T., & Silva, G. P. (2008). Fitossociologia do cerrado stricto sensu no município de Carolina, MA, Brasil. <i>Cerne</i> , 14(4), 285-294. |
| Article | Imaña-Encinas, J., & Paula, J. E. D. (2003). Análise da vegetação de cerrado no município de Santa Quitéria-Maranhão. <i>Brasil Florestal</i> , 78, 33-42. |
| Article | Ribeiro, L. F., & Tabarelli, M. (2002). A structural gradient in cerrado vegetation of Brazil: changes in woody plant density, species richness, life history and plant composition. <i>Journal of tropical ecology</i> , 18(05), 775-794. |
| Article | Vidotto, E., Pessenda, L.C.R., Ribeiro, A.S., Freitas, H.A., Bendassolli, J.A. (2007). Dinâmica do ecotônico floresta-campo no sul do estado do Amazonas no Holoceno, através de estudos isotópicos e fitossociológicos. <i>Acta Amazônica</i> , 37(3), 385-400. |
| Article | Sanaiotti, T. M., Bridgewater, S., & Ratter, J. A. (1997). A floristic study of the savanna vegetation of the state of Amapá, Brazil, and suggestions for its conservation. <i>Boletim do Museu Paraense Emilio Goeldi, Série Botânica</i> , 13(1), 3-29. |

| Literature | Reference |
|------------|--|
| Article | Miranda, I. S. (1993). Estrutura do estrato arbóreo do cerrado amazônico em Alter-do-Chão, Pará, Brasil. <i>Revista Brasileira de Botânica</i> , 16(2), 143-150. |
| Article | Bastos, M. D. N. D. C. (1984). Levantamento florístico dos campos do Estado do Pará. I. Campo de Joanes (Ilha de Marajó). <i>Boletim Museu Paraense Emilio Goeldi, Série Botânica</i> , 1(1/2), 67-86. |
| Article | Miranda, I. S., Almeida, S. S., & Dantas, P. J. (2006). Florística e estrutura de comunidades arbóreas em cerrados de Rondônia, Brasil. <i>Acta Amazônica</i> , 36(4), 419-430. |
| Article | Saporetti Jr, A. W., Meira Neto, J. A. A., & Almado, R. D. P. (2003). Fitossociologia de cerrado sensu stricto no município de Abaeté-MG. <i>Revista Árvore</i> , 27(3), 413-419. |
| Article | Campos, E. P. D., Duarte, T. G., Neri, A. V., Silva, A. F. D., Meira-Neto, J. A. A., & Valente, G. E. (2006). Composição florística de um trecho de cerradão e cerrado sensu stricto e sua relação com o solo na Floresta Nacional (FLONA) de Paraopeba, MG, Brasil. <i>Revista Árvore</i> , 30(3), 471-479. |
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| Article | Cardoso, E., Moreno, M. I. C., & Guimarães, A. J. M. (2002). Estudo fitossociológico em área de cerrado sensu stricto na Estação de Pesquisa e Desenvolvimento Ambiental Galheiro-Perdizes, MG. <i>Caminhos de Geografia</i> , 3(5). |
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| Article | Brandão, M., Laca Buendia, J. P., Araujo, M. G., & Ferreira, F. B. D. (1995). Município de Uberaba, MG: cobertura vegetal e composição florística. <i>Daphne</i> , 5(1), 19-39. |
| Article | Costa, A. A., & Araújo, G. M. D. (2001). Comparação da vegetação arbórea de cerradão e de cerrado na Reserva do Panga, Uberlândia, Minas Gerais. <i>Acta botanica brasiliaca</i> . |
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| Article | Ratter, J. A., Leitao Filho, H., Argent, G., Gibbs, P. E., Semir, J., Shepherd, G., & Tamashiro, J. (1988). Floristic composition and community structure of a southern cerrado area in Brazil. Notes from the Royal Botanic Garden, Edinburgh. |
| Article | Durigan, G., Bacic, M. C., Franco, G. A. D. C., & Siqueira, M. D. (1999). Inventario floristico de Cerrado na Estacao Ecologica de Assis, SP. Inventory of cerrado in the Assis Ecological Station.) <i>Hoehnea</i> , 26(2), 149-172. |
| Article | Brando, P. M., & Durigan, G. (2005). Changes in cerrado vegetation after disturbance by frost (São Paulo State, Brazil). <i>Plant Ecology</i> , 175(2), 205-215. |

| Literature | Reference |
|------------|---|
| Article | Cesar, O., Pagano, S. N., Leitao Filho, H. D. F., Monteiro, R., da Silva, O. A., Marinis, G. D., & Shepherd, G. J. (1988). Estrutura fitossociologica do estrato arboreo de uma area de vegetacao de cerrado no municipio de Corumbatal. <i>Naturalia</i> , 13, 91-101. |
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Table 3. For each species the table has a total number of latitudes belt that a species has occurred in our database (lat), the species that were below (1) and after breaking point (3), and the total number of ecoregions where species were found (ecoregion). Tree species (s) of the South American savanna (n=2,203). Rabinowitz's forms (form), the value of protection ratio (r) and the status of protection (status) for each species are presented. The values of r range from 0 (no species occurrence within protected areas) to 1 (all species occurrence inside protected area).

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Abarema barbouriana</i> | (Standl.)Barneby & J.W.Grimes | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Abarema barnebyana</i> | Iganci & M.P.Morim | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Abarema campestris</i> | (Benth.)Barneby & J.W.Grimes | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Abarema cochleata</i> | (Willd.)Barneby & J.W.Grimes | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Abarema ferruginea</i> | (Benth.)Pittier | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Abarema filamentosa</i> | (Benth.)Pittier | 9 | 3 | 5 | common | 0.3 | partially protected |
| <i>Abarema jupunba</i> | (Willd.)Britton & Killip | 7 | 3 | 6 | common | 0.444444 | partially protected |
| <i>Abatia americana</i> | (Gardner) Eichler | 2 | 1 | 2 | form5 | 0.4 | partially protected |
| <i>Abuta grandifolia</i> | (Mart.) Sandwith | 24 | 3 | 25 | common | 0.284553 | partially protected |
| <i>Acalypha stenoloba</i> | MÃƒÂfÃ¼ll.Arg. | 2 | 1 | 1 | form7 | 1 | protected |
| <i>Acosmium bijugum</i> | (Vogel)Yakovlev | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Acosmium cardenasii</i> | H.S.Irwin & Arroyo | 8 | 3 | 5 | common | 0.272727 | partially protected |
| <i>Acosmium dasycarpum</i> | (Vogel)Yakovlev | 23 | 3 | 16 | common | 0.126285 | partially protected |
| <i>Acosmium nitens</i> | (Vogel)Yakovlev | 15 | 3 | 12 | common | 0.255814 | partially protected |
| <i>Acritopappus catolesensis</i> | D.J.N.Hind & Bautista | 5 | 3 | 3 | common | 0.222222 | partially protected |
| <i>Acritopappus confertus</i> | (Gardner) R.M.King & H.Rob. | 9 | 3 | 7 | common | 0.358974 | partially protected |
| <i>Acritopappus connatifolius</i> | (Soar.Nunes) R.M.King & H.Rob. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Acritopappus harleyi</i> | R.M.King & H.Rob. | 3 | 3 | 3 | form4 | 0.3 | partially protected |
| <i>Acritopappus heterolepis</i> | (Baker) R.M.King & H.Rob. | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Acritopappus irwinii</i> | R.M.King & H.Rob. | 3 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Acritopappus pereirae</i> | Bautista, S.Ortiz & Rodr.Oubi | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Acritopappus pintoi</i> | Bautista & D.J.N.Hind | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Acritopappus subomentosus</i> | R.M.King & H.Rob. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Acritopappus teixeirae</i> | R.M.King & H.Rob. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Acrocomia aculeata</i> | (Jacq.) Lodd. ex Mart. | 29 | 3 | 20 | common | 0.128134 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Acrocomia glaucescens</i> | Lorenzi | 4 | 3 | 1 | form2 | 0 | unprotected |
| <i>Actinostemon schomburgkii</i> | (Klotzsch) Hochr. | 7 | 3 | 6 | common | 0.5 | partially protected |
| <i>Adelia membranifolia</i> | (MÃƒÂ¼ll.Arg.) Chodat & Hassl. | 2 | 1 | 1 | form7 | 0.2 | partially protected |
| <i>Adenaria floribunda</i> | Kunth | 15 | 3 | 9 | common | 0.111111 | partially protected |
| <i>Aegiphila bracteolosa</i> | Moldenke | 9 | 1 | 9 | form1 | 0.5 | partially protected |
| <i>Aegiphila integrifolia</i> | (Jacq.) B.D.Jacks. | 32 | 3 | 22 | common | 0.199029 | partially protected |
| <i>Aegiphila laxiflora</i> | Benth. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Aegiphila luschnathii</i> | Schauer | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Aegiphila mollis</i> | Kunth | 7 | 3 | 8 | common | 0.111111 | partially protected |
| <i>Aegiphila pernambucensis</i> | Moldenke | 2 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Aegiphila verticillata</i> | Vell. | 28 | 3 | 27 | common | 0.167176 | partially protected |
| <i>Agarista boliviensis</i> | (Sleumer) Judd | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Agarista chapadensis</i> | (Kin.-Gouv.) Judd | 4 | 3 | 3 | common | 0.5 | partially protected |
| <i>Agarista coriifolia</i> | (Thunb.) Hook. ex Nied. | 8 | 3 | 6 | common | 0.25 | partially protected |
| <i>Agarista duckei</i> | (Huber) Judd | 5 | 3 | 4 | common | 0.5 | partially protected |
| <i>Agarista glaberrima</i> | (Sleumer) Judd | 5 | 3 | 3 | common | 0.227273 | partially protected |
| <i>Agarista oleifolia</i> | (Cham.) G.Don | 8 | 3 | 4 | common | 0.190476 | partially protected |
| <i>Agarista pulchra</i> | G.Don | 4 | 3 | 5 | common | 0.357143 | partially protected |
| <i>Agarista revoluta</i> | (Spreng.) Hook. ex Nied. | 5 | 3 | 4 | common | 0.333333 | partially protected |
| <i>Agonandra brasiliensis</i> | Miers ex Benth. | 29 | 3 | 25 | common | 0.125616 | partially protected |
| <i>Aiouea macedoana</i> | Vattimo | 5 | 3 | 3 | common | 0 | unprotected |
| <i>Aiouea piauhyensis</i> | (Meisn.) Mez | 15 | 3 | 7 | common | 0.2 | partially protected |
| <i>Aiouea saligna</i> | Meisn. | 15 | 3 | 7 | common | 0.2 | partially protected |
| <i>Aiouea trinervis</i> | Meisn. | 16 | 3 | 12 | common | 0.2 | partially protected |
| <i>Albizia guachapele</i> | (Kunth)Dugand | 6 | 3 | 2 | common | 0.045455 | partially protected |
| <i>Albizia inundata</i> | (Mart.)Barneby & J.W.Grimes | 3 | 3 | 2 | form4 | 0 | unprotected |
| <i>Albizia niopoides</i> | (Benth.)Burkart | 15 | 3 | 11 | common | 0.074074 | partially protected |
| <i>Albizia pedicellaris</i> | (Dc.)L.Rico | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Albizia saman</i> | (Jacq.)Merr. | 2 | 3 | 2 | form4 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Albizia subdimidiata</i> | (Splitg.) Barneby & J.W.Grimes | 15 | 3 | 17 | common | 0.571429 | partially protected |
| <i>Alchornea castaneifolia</i> | (Humb. & Bonpl. ex Willd.) A.Juss. | 18 | 3 | 15 | common | 0.210526 | partially protected |
| <i>Alchornea discolor</i> | Poepp. | 19 | 3 | 10 | common | 0.21875 | partially protected |
| <i>Alchornea glandulosa</i> | Poepp. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Alchornea latifolia</i> | Sw. | 5 | 1 | 3 | form1 | 0.333333 | partially protected |
| <i>Alchornea triplinervia</i> | (Spreng.) MÃƒÂ¼ll.Arg. | 5 | 3 | 4 | common | 0.3 | partially protected |
| <i>Algernonia kuhlmannii</i> | (Emmerich) G.L.Webster | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Alibertia bertieriifolia</i> | K.Schum. | 16 | 3 | 15 | common | 0.173913 | partially protected |
| <i>Alibertia duckei</i> | Standl. | 5 | 1 | 5 | form1 | 1 | protected |
| <i>Alibertia edulis</i> | (Rich.) A.Rich. ex DC. | 34 | 3 | 30 | common | 0.167986 | partially protected |
| <i>Alibertia latifolia</i> | (Benth.) K.Schum. | 13 | 3 | 14 | common | 0.272727 | partially protected |
| <i>Alibertia pilosa</i> | K.Krause | 5 | 1 | 6 | form1 | 0 | unprotected |
| <i>Alibertia rotunda</i> | (Cham.) K.Schum. | 6 | 3 | 3 | common | 0.345455 | partially protected |
| <i>Allagoptera caudescens</i> | (Mart.) Kuntze | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Allamanda blanchetii</i> | A.DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Allophylus edulis</i> | (A.St.-Hil., A.Juss. & Cambess.) Radlk. | 4 | 3 | 3 | common | 0 | unprotected |
| <i>Allophylus quercifolius</i> | (Mart.) Radlk. | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Allophylus racemosus</i> | Sw. | 13 | 3 | 9 | common | 0.458333 | partially protected |
| <i>Aloysia virgata</i> | (Ruiz & Pav.) Juss. | 7 | 3 | 6 | common | 0.133333 | partially protected |
| <i>Amaioua guianensis</i> | Aubl. | 27 | 3 | 26 | common | 0.214286 | partially protected |
| <i>Amaioua intermedia</i> | Mart. ex Schult. & Schult.f. | 10 | 3 | 3 | common | 0.357143 | partially protected |
| <i>Amanoa glaucophylla</i> | MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Amazonia campestris</i> | (Aubl.) Moldenke | 3 | 1 | 2 | form5 | 0.2 | partially protected |
| <i>Amburana cearensis</i> | (Allemao)A.C.Sm. | 5 | 3 | 5 | common | 0.055556 | partially protected |
| <i>Ampelocera edentula</i> | Kuhlm. | 20 | 3 | 19 | common | 0.473684 | partially protected |
| <i>Anacardium occidentale</i> | L. | 30 | 3 | 25 | common | 0.180328 | partially protected |
| <i>Anadenanthera colubrina</i> | (Vell.)Brenan | 19 | 3 | 13 | common | 0.068376 | partially protected |
| <i>Anadenanthera peregrina</i> | (L.)Speg. | 30 | 3 | 21 | common | 0.120301 | partially protected |
| <i>Andira anthelmia</i> | (Vell.) Benth. | 2 | 1 | 2 | form5 | 0.375 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------|----------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Andira cordata</i> | R.T.Penn. & H.C.Lima | 13 | 3 | 5 | common | 0.15942 | partially protected |
| <i>Andira cujabensis</i> | Benth. | 19 | 3 | 11 | common | 0.149329 | partially protected |
| <i>Andira fraxinifolia</i> | Benth. | 18 | 3 | 11 | common | 0.184783 | partially protected |
| <i>Andira humilis</i> | Mart. ex Benth. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Andira inermis</i> | (Wright)DC. | 17 | 3 | 10 | common | 0.238095 | partially protected |
| <i>Andira legalis</i> | (Vell.) Toledo | 3 | 1 | 2 | form5 | 0.714286 | partially protected |
| <i>Andira nitida</i> | Benth. | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Andira vermicula</i> | Benth. | 21 | 3 | 14 | common | 0.127764 | partially protected |
| <i>Anemopaegma velutinum</i> | Mart. ex DC. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Aniba firmula</i> | (Nees & Mart.) Mez | 2 | 1 | 2 | form5 | 0.8 | partially protected |
| <i>Aniba heringeri</i> | Vattimo | 3 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Anisacanthus trilobus</i> | Lindau | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Annona acutiflora</i> | Mart. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Annona aurantiaca</i> | Barb.Rodr. | 10 | 3 | 3 | common | 0.215686 | partially protected |
| <i>Annona cacans</i> | Warm. | 4 | 3 | 2 | common | 0.153846 | partially protected |
| <i>Annona coriacea</i> | Mart. | 23 | 3 | 17 | common | 0.135327 | partially protected |
| <i>Annona crassiflora</i> | Mart. | 24 | 3 | 15 | common | 0.132196 | partially protected |
| <i>Annona cuspidata</i> | (Mart.) H.Rainer | 16 | 3 | 11 | common | 0.142857 | partially protected |
| <i>Annona dioica</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Annona emarginata</i> | (Schltdl.) H.Rainer | 14 | 3 | 9 | common | 0.16129 | partially protected |
| <i>Annona exsucca</i> | DC. | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Annona glabra</i> | L. | 11 | 3 | 6 | common | 0.448276 | partially protected |
| <i>Annona hypoglauca</i> | Mart. | 19 | 3 | 22 | common | 0.111111 | partially protected |
| <i>Annona jahnii</i> | Saff. | 10 | 3 | 9 | common | 0.072727 | partially protected |
| <i>Annona leptopetala</i> | (R.E.Fr.) H.Rainer | 11 | 3 | 6 | common | 0.178571 | partially protected |
| <i>Annona montana</i> | Macfad. | 24 | 3 | 19 | common | 0.163636 | partially protected |
| <i>Annona nutans</i> | (R.E.Fr.) R.E.Fr. | 8 | 3 | 5 | common | 0.277778 | partially protected |
| <i>Annona paludosa</i> | Aubl. | 20 | 3 | 13 | common | 0.298246 | partially protected |
| <i>Annona pickelii</i> | (Diels) H.Rainer | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Annona sylvatica</i> | A.St.-Hil. | 10 | 3 | 4 | common | 0.125 | partially protected |
| <i>Anteremanthus hatschbachii</i> | H.Rob. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Antonia ovata</i> | Pohl | 26 | 3 | 21 | common | 0.185015 | partially protected |
| <i>Aparisthium cordatum</i> | (A.Juss.) Baill. | 23 | 3 | 25 | common | 0.396226 | partially protected |
| <i>Apeiba tibourbou</i> | Aubl. | 25 | 3 | 28 | common | 0.136364 | partially protected |
| <i>Apterokarpos gardneri</i> | (Engl.) Rizzini | 2 | 3 | 2 | form4 | 0.333333 | partially protected |
| <i>Apuleia leiocarpa</i> | (Vogel) J.F.Macbr. | 28 | 3 | 29 | common | 0.192513 | partially protected |
| <i>Archytaea triflora</i> | Mart. | 5 | 3 | 4 | common | 0.555556 | partially protected |
| <i>Arrojadoa rhodantha</i> | (GÃƒÂƒrke) Britton & Rose | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Arthrocereus melanurus</i> | (K.Schum.) Diers, P.Br. & Esteves | 8 | 1 | 9 | form1 | 0 | unprotected |
| <i>Aspidosperma cuspa</i> | (Kunth) S.F.Blake ex Pittier | 20 | 3 | 13 | common | 0.116751 | partially protected |
| <i>Aspidosperma cylindrocarpon</i> | MÃƒÂƒll.Arg. | 16 | 3 | 10 | common | 0.103448 | partially protected |
| <i>Aspidosperma discolor</i> | A.DC. | 21 | 3 | 15 | common | 0.224719 | partially protected |
| <i>Aspidosperma dispermum</i> | MÃƒÂƒll.Arg. | 4 | 3 | 3 | common | 0.454545 | partially protected |
| <i>Aspidosperma macrocarpon</i> | Mart. | 26 | 3 | 26 | common | 0.142427 | partially protected |
| <i>Aspidosperma multiflorum</i> | A.DC. | 16 | 3 | 10 | common | 0.183333 | partially protected |
| <i>Aspidosperma nobile</i> | MÃƒÂƒll.Arg. | 16 | 3 | 10 | common | 0.154362 | partially protected |
| <i>Aspidosperma polyneuron</i> | MÃƒÂƒll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Aspidosperma pyrifolium</i> | Mart. | 18 | 3 | 11 | common | 0.088496 | partially protected |
| <i>Aspidosperma spruceanum</i> | Benth. ex MÃƒÂƒll.Arg. | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Aspidosperma subincanum</i> | Mart. ex A.DC. | 22 | 3 | 15 | common | 0.161348 | partially protected |
| <i>Aspidosperma tomentosum</i> | Mart. | 22 | 3 | 16 | common | 0.1141 | partially protected |
| <i>Astrocaryum campestre</i> | Mart. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Astrocaryum echinatum</i> | Barb.Rodr. | 8 | 3 | 3 | common | 0.21875 | partially protected |
| <i>Astrocaryum ulei</i> | Burret | 10 | 3 | 7 | common | 0.428571 | partially protected |
| <i>Astrocaryum vulgare</i> | Mart. | 22 | 3 | 16 | common | 0.322917 | partially protected |
| <i>Astronium fraxinifolium</i> | Schott | 25 | 3 | 18 | common | 0.136895 | partially protected |
| <i>Astronium graveolens</i> | Jacq. | 21 | 3 | 14 | common | 0.432432 | partially protected |
| <i>Attalea brasiliensis</i> | Glassman | 4 | 3 | 2 | common | 0.3 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|---------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Attalea butyracea</i> | (Mutis ex L.f.) Wess.Boer | 10 | 3 | 8 | common | 0.333333 | partially protected |
| <i>Attalea compta</i> | Mart. | 3 | 3 | 2 | form4 | 0.083333 | partially protected |
| <i>Attalea funifera</i> | Mart. | 4 | 3 | 4 | common | 0.2 | partially protected |
| <i>Attalea maripa</i> | (Aubl.) Mart. | 19 | 3 | 21 | common | 0.44898 | partially protected |
| <i>Attalea phalerata</i> | Mart. ex Spreng. | 26 | 3 | 21 | common | 0.194245 | partially protected |
| <i>Attalea speciosa</i> | Mart. ex Spreng. | 21 | 3 | 20 | common | 0.292135 | partially protected |
| <i>Aureliana fasciculata</i> | (Vell.) Sendtn. | 3 | 1 | 2 | form5 | 1 | protected |
| <i>Austrocritonia angulicaulis</i> | (Sch.Bip. ex Baker) R.M.King & H.Rob. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Austroeupatorium inulaefolium</i> | (Kunth) R.M.King & H.Rob. | 7 | 3 | 4 | common | 0.4375 | partially protected |
| <i>Averrhoidium gardnerianum</i> | Baill. | 5 | 3 | 2 | common | 0.4 | partially protected |
| <i>Baccharis calvescens</i> | DC. | 6 | 3 | 4 | common | 0.25 | partially protected |
| <i>Baccharis dracunculifolia</i> | DC. | 12 | 3 | 8 | common | 0.150442 | partially protected |
| <i>Baccharis intermixta</i> | Gardner | 7 | 3 | 5 | common | 0.272727 | partially protected |
| <i>Baccharis lychnophora</i> | Gardner | 3 | 3 | 3 | form4 | 0.384615 | partially protected |
| <i>Baccharis microdonta</i> | DC. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Baccharis oblongifolia</i> | (Ruiz & Pav.) Pers. | 9 | 3 | 6 | common | 0.365385 | partially protected |
| <i>Baccharis retusa</i> | DC. | 11 | 3 | 5 | common | 0.278481 | partially protected |
| <i>Baccharis rufidula</i> | (Spreng.) Joch.MÃƒÂfÃ¼ll. | 3 | 3 | 2 | form4 | 0.4 | partially protected |
| <i>Baccharis singularis</i> | (Vell.) G.M.Barroso | 4 | 3 | 3 | common | 0.4 | partially protected |
| <i>Baccharis vulneraria</i> | Baker | 9 | 3 | 5 | common | 0.346154 | partially protected |
| <i>Bactris campestris</i> | Poepp. | 3 | 1 | 4 | form5 | 0.5 | partially protected |
| <i>Bactris concinna</i> | Mart. | 12 | 3 | 8 | common | 0 | unprotected |
| <i>Bactris ferruginea</i> | Burret | 6 | 3 | 5 | common | 0 | unprotected |
| <i>Bactris glaucescens</i> | Drude | 8 | 3 | 5 | common | 0.1 | partially protected |
| <i>Bactris horridispatha</i> | Noblick ex A.J.Hend. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Bactris major</i> | Jacq. | 22 | 3 | 26 | common | 0.409091 | partially protected |
| <i>Bactris riparia</i> | Mart. | 14 | 3 | 12 | common | 0.411765 | partially protected |
| <i>Bactris vulgaris</i> | Barb.Rodr. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Balfourodendron molle</i> | (Miq.) Pirani | 3 | 3 | 2 | form4 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Banara arguta</i> | Briq. | 16 | 3 | 14 | common | 0.125 | partially protected |
| <i>Banara guianensis</i> | Aubl. | 18 | 3 | 17 | common | 0.5625 | partially protected |
| <i>Banara nitida</i> | Spruce ex Benth. | 8 | 1 | 9 | form1 | 0 | unprotected |
| <i>Banisteriopsis arborea</i> | B.Gates | 2 | 1 | 3 | form5 | 0.2 | partially protected |
| <i>Banisteriopsis latifolia</i> | (A.Juss.) B.Gates | 14 | 3 | 5 | common | 0.230088 | partially protected |
| <i>Barnebya harleyi</i> | W.R.Anderson & B.Gates | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Bastardiodia densiflora</i> | (Hook. & Arn.) Hassl. | 8 | 3 | 3 | common | 0.142857 | partially protected |
| <i>Bauhinia aculeata</i> | L. | 4 | 3 | 4 | common | 0 | unprotected |
| <i>Bauhinia acuruana</i> | Moric. | 13 | 3 | 5 | common | 0.142857 | partially protected |
| <i>Bauhinia bauhinoides</i> | (Mart.)J.F.Macbr. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Bauhinia bombaciflora</i> | Ducke | 13 | 3 | 7 | common | 0.038462 | partially protected |
| <i>Bauhinia cheilantha</i> | (Bong.)Steud. | 5 | 3 | 3 | common | 0.090909 | partially protected |
| <i>Bauhinia corniculata</i> | Benth. | 10 | 3 | 11 | common | 0.272727 | partially protected |
| <i>Bauhinia cupulata</i> | Benth. | 16 | 3 | 8 | common | 0.148515 | partially protected |
| <i>Bauhinia dubia</i> | G.Don | 10 | 3 | 9 | common | 0.162162 | partially protected |
| <i>Bauhinia forficata</i> | Link | 12 | 3 | 9 | common | 0.096154 | partially protected |
| <i>Bauhinia longicuspis</i> | Benth. | 3 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Bauhinia longifolia</i> | (Bong.)Steud. | 6 | 3 | 5 | common | 0.058824 | partially protected |
| <i>Bauhinia longipedicellata</i> | Ducke | 3 | 3 | 3 | form4 | 1 | protected |
| <i>Bauhinia membranacea</i> | Benth. | 8 | 3 | 4 | common | 0.142857 | partially protected |
| <i>Bauhinia mollis</i> | (Bong.) D.Dietr. | 15 | 3 | 9 | common | 0.133333 | partially protected |
| <i>Bauhinia pentandra</i> | (Bong.)Steud. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Bauhinia platypetala</i> | Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Bauhinia pulchella</i> | Benth. | 20 | 3 | 10 | common | 0.179641 | partially protected |
| <i>Bauhinia rufa</i> | (Bong.)Steud. | 20 | 3 | 15 | common | 0.137157 | partially protected |
| <i>Bauhinia subclavata</i> | Benth. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Bauhinia unguis</i> | L. | 34 | 3 | 24 | common | 0.155216 | partially protected |
| <i>Bejaria sprucei</i> | Meisn. | 5 | 3 | 4 | common | 0.6 | partially protected |
| <i>Bellucia acutata</i> | Pilg. | 19 | 3 | 14 | common | 0.230769 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Bellucia grossularioides</i> | (L.) Triana | 22 | 3 | 24 | common | 0.191489 | partially protected |
| <i>Berberis laurina</i> | Billb. | 1 | 1 | 2 | form5 | 0.8 | partially protected |
| <i>Bixa arborea</i> | Huber | 19 | 3 | 14 | common | 0.333333 | partially protected |
| <i>Blepharandra hypoleuca</i> | (Benth.) Griseb. | 2 | 3 | 2 | form4 | 0.555556 | partially protected |
| <i>Blepharocalyx salicifolius</i> | (Kunth) O.Berg | 19 | 3 | 12 | common | 0.182819 | partially protected |
| <i>Bocageopsis mattogrossensis</i> | (R.E.Fr.) R.E.Fr. | 15 | 3 | 8 | common | 0.2 | partially protected |
| <i>Bocageopsis multiflora</i> | (Mart.) R.E.Fr. | 18 | 3 | 18 | common | 0.285714 | partially protected |
| <i>Bocoa ratteri</i> | H.E. Ireland | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Bonnetia paniculata</i> | Spruce ex Benth. | 13 | 3 | 13 | common | 0.7 | partially protected |
| <i>Bonnetia sessilis</i> | Benth. | 7 | 3 | 9 | common | 0.5 | partially protected |
| <i>Bonnetia stricta</i> | (Nees) Nees & Mart. | 7 | 3 | 6 | common | 0.3125 | partially protected |
| <i>Bonyunia antoniifolia</i> | Progel | 13 | 3 | 9 | common | 0.692308 | partially protected |
| <i>Bonyunia minor</i> | N.E.Br. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Bonyunia superba</i> | R.H.Schomb. ex Progel | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Bougainvillea glabra</i> | Choisy | 8 | 3 | 2 | common | 0.272727 | partially protected |
| <i>Bourreria boliviensis</i> | Gottschling & J.S.Mill. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Bowdichia virgilioides</i> | Kunth | 34 | 3 | 37 | common | 0.154523 | partially protected |
| <i>Brasiliopuntia brasiliensis</i> | (Willd.) A.Berger | 8 | 3 | 5 | common | 0.166667 | partially protected |
| <i>Bredemeyera floribunda</i> | Willd. | 26 | 3 | 23 | common | 0.174033 | partially protected |
| <i>Bredemeyera hebeclada</i> | (DC.) J.F.B. Pastore | 3 | 1 | 4 | form5 | 0.333333 | partially protected |
| <i>Bredemeyera laurifolia</i> | Klotzsch ex A.W.Benn. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Bredemeyera myrtifolia</i> | A.W.Benn. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Brosimum alicastrum</i> | Sw. | 16 | 3 | 14 | common | 0.227273 | partially protected |
| <i>Brosimum gaudichaudii</i> | TrÃƒÂ©cul | 24 | 3 | 22 | common | 0.136126 | partially protected |
| <i>Brosimum glaucum</i> | Taub. | 2 | 1 | 2 | form5 | 0.8 | partially protected |
| <i>Buchenavia oxycarpa</i> | (Mart.) Eichler | 19 | 3 | 21 | common | 0.181818 | partially protected |
| <i>Buchenavia tetraphylla</i> | (Aubl.) R.A.Howard | 26 | 3 | 31 | common | 0.300971 | partially protected |
| <i>Buchenavia tomentosa</i> | Eichler | 25 | 3 | 25 | common | 0.115714 | partially protected |
| <i>Buddleja stachyoides</i> | Cham. & Schldl. | 4 | 3 | 4 | common | 0.416667 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|-------------------------|-----|------------|-----------|----------|---------------------|--------|
| <i>Bunchosia argentea</i> | (Jacq.) DC. | 6 | 3 | 3 common | 0.666667 | partially protected | |
| <i>Bunchosia mollis</i> | Benth. | 5 | 3 | 2 common | 0.086957 | partially protected | |
| <i>Bunchosia odorata</i> | (Jacq.) Juss. | 1 | 1 | 1 form7 | | 0 unprotected | |
| <i>Bunchosia paraguariensis</i> | Nied. | 7 | 3 | 4 common | 0.416667 | partially protected | |
| <i>Butia capitata</i> | (Mart.) Becc. | 9 | 3 | 4 common | 0.119718 | partially protected | |
| <i>Butia paraguayensis</i> | (Barb.Rodr.) L.H.Bailey | 8 | 3 | 4 common | 0.107527 | partially protected | |
| <i>Butia purpurascens</i> | Glassman | 4 | 3 | 2 common | 0.096774 | partially protected | |
| <i>Byrsonima affinis</i> | W.R.Anderson | 11 | 3 | 5 common | 0.081818 | partially protected | |
| <i>Byrsonima arthropoda</i> | A.Juss. | 16 | 3 | 10 common | 0.37037 | partially protected | |
| <i>Byrsonima basiloba</i> | A.Juss. | 1 | 1 | 2 form5 | | 0 unprotected | |
| <i>Byrsonima christianeae</i> | W.R.Anderson | 3 | 1 | 3 form5 | | 1 protected | |
| <i>Byrsonima clausseniana</i> | A.Juss. | 9 | 3 | 5 common | 0.166667 | partially protected | |
| <i>Byrsonima coccobifolia</i> | Kunth | 33 | 3 | 27 common | 0.138009 | partially protected | |
| <i>Byrsonima concinna</i> | Benth. | 10 | 3 | 9 common | 0.636364 | partially protected | |
| <i>Byrsonima coniophylla</i> | A.Juss. | 10 | 1 | 6 form1 | | 1 protected | |
| <i>Byrsonima correifolia</i> | A.Juss. | 15 | 3 | 7 common | 0.178404 | partially protected | |
| <i>Byrsonima crassifolia</i> | (L.) Kunth | 34 | 3 | 34 common | 0.220447 | partially protected | |
| <i>Byrsonima crispa</i> | A.Juss. | 20 | 3 | 21 common | 0.315789 | partially protected | |
| <i>Byrsonima cydoniifolia</i> | A.Juss. | 19 | 3 | 17 common | 0.203931 | partially protected | |
| <i>Byrsonima fanshawei</i> | W.R.Anderson | 2 | 1 | 1 form7 | 0.5 | partially protected | |
| <i>Byrsonima fonsecae</i> | W.R.Anderson | 2 | 1 | 2 form5 | | 0 unprotected | |
| <i>Byrsonima gardneriana</i> | A.Juss. | 14 | 3 | 11 common | 0.396226 | partially protected | |
| <i>Byrsonima hypoleuca</i> | Turcz. | 1 | 1 | 1 form7 | | 1 protected | |
| <i>Byrsonima incarnata</i> | Sandwith | 7 | 3 | 9 common | | 1 protected | |
| <i>Byrsonima intermedia</i> | A.Juss. | 20 | 3 | 13 common | 0.11617 | partially protected | |
| <i>Byrsonima lancifolia</i> | A.Juss. | 13 | 3 | 7 common | 0.24 | partially protected | |
| <i>Byrsonima laxiflora</i> | Griseb. | 10 | 3 | 3 common | 0.413043 | partially protected | |
| <i>Byrsonima linguifera</i> | Cuatrec. | 10 | 3 | 6 common | 0.046512 | partially protected | |
| <i>Byrsonima martiana</i> | A.Juss. | 2 | 1 | 1 form7 | 0.5 | partially protected | |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|-----------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Byrsonima nitidissima</i> | Kunth | 3 | 3 | 1 | form6 | 0.25 | partially protected |
| <i>Byrsonima pachyphylla</i> | A.Juss. | 20 | 3 | 13 | common | 0.138288 | partially protected |
| <i>Byrsonima punctulata</i> | A.Juss. | 17 | 3 | 11 | common | 0.4 | partially protected |
| <i>Byrsonima rotunda</i> | Griseb. | 12 | 3 | 4 | common | 0.2 | partially protected |
| <i>Byrsonima salzmanniana</i> | A.Juss. | 4 | 3 | 3 | common | 0.2 | partially protected |
| <i>Byrsonima schomburgkiana</i> | Benth. | 9 | 3 | 8 | common | 0.392857 | partially protected |
| <i>Byrsonima sericea</i> | DC. | 20 | 3 | 21 | common | 0.253333 | partially protected |
| <i>Byrsonima spicata</i> | (Cav.) Rich. ex Kunth | 13 | 3 | 10 | common | 0.470588 | partially protected |
| <i>Byrsonima spinensis</i> | W.R.Anderson | 5 | 3 | 2 | common | 0.428571 | partially protected |
| <i>Byrsonima stannardii</i> | W.R.Anderson | 4 | 3 | 3 | common | 0.3 | partially protected |
| <i>Byrsonima stipulacea</i> | A.Juss. | 16 | 3 | 16 | common | 0.526316 | partially protected |
| <i>Byrsonima umbellata</i> | Mart. ex A.Juss. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Byrsonima vacciniifolia</i> | A.Juss. | 12 | 3 | 6 | common | 0.160714 | partially protected |
| <i>Byrsonima variabilis</i> | A.Juss. | 18 | 3 | 8 | common | 0.214286 | partially protected |
| <i>Byrsonima verbascifolia</i> | (L.) DC. | 34 | 3 | 27 | common | 0.154808 | partially protected |
| <i>Cabralea canjerana</i> | (Vell.) Mart. | 20 | 3 | 12 | common | 0.19403 | partially protected |
| <i>Caesalpinia coriaria</i> | (Jacq.) Willd. | 4 | 3 | 2 | common | 0.055556 | partially protected |
| <i>Calliandra asplenioides</i> | (Nees)Renvoize | 10 | 3 | 5 | common | 0.268293 | partially protected |
| <i>Calliandra bella</i> | (Spreng.)Benth. | 3 | 3 | 3 | form4 | 0.375 | partially protected |
| <i>Calliandra brevipes</i> | Benth. | 5 | 3 | 5 | common | 0.454545 | partially protected |
| <i>Calliandra calycina</i> | Benth. | 4 | 3 | 3 | common | 0.444444 | partially protected |
| <i>Calliandra carrascana</i> | Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Calliandra elegans</i> | Renvoize | 3 | 3 | 3 | form4 | 0.285714 | partially protected |
| <i>Calliandra erubescens</i> | Renvoize | 4 | 3 | 3 | common | 0.285714 | partially protected |
| <i>Calliandra fernandesii</i> | Barneby | 3 | 1 | 1 | form7 | 0.4 | partially protected |
| <i>Calliandra foliolosa</i> | Benth. | 10 | 3 | 6 | common | 0.23913 | partially protected |
| <i>Calliandra glomerulata</i> | H.Karst. | 3 | 3 | 1 | form6 | 0.125 | partially protected |
| <i>Calliandra harrisii</i> | (Lindl.)Benth. | 3 | 3 | 3 | form4 | 0.4 | partially protected |
| <i>Calliandra laxa</i> | (Willd.) Benth. | 16 | 3 | 18 | common | 0.485714 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|---------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Calliandra lintea</i> | Barneby | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Calliandra macrocalyx</i> | Harms | 3 | 3 | 4 | form4 | 0 | unprotected |
| <i>Calliandra parviflora</i> | Benth. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Calliandra renvoizeana</i> | Barneby | 3 | 3 | 2 | form4 | 0.4 | partially protected |
| <i>Calliandra surinamensis</i> | Benth. | 12 | 3 | 16 | common | 0.428571 | partially protected |
| <i>Calliandra trinervia</i> | Benth. | 12 | 3 | 9 | common | 0.2 | partially protected |
| <i>Calliandra umbellifera</i> | Benth. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Callianthe rufinerva</i> | (A. St.-Hil.) Donnell | 1 | 1 | 3 | form5 | 0.545455 | partially protected |
| <i>Callisthene fasciculata</i> | Mart. | 20 | 3 | 13 | common | 0.110656 | partially protected |
| <i>Callisthene major</i> | Mart. | 18 | 3 | 11 | common | 0.186335 | partially protected |
| <i>Callisthene microphylla</i> | Warm. | 14 | 3 | 8 | common | 0.265487 | partially protected |
| <i>Callisthene minor</i> | Mart. | 11 | 3 | 4 | common | 0.105263 | partially protected |
| <i>Callisthene mollissima</i> | Warm. | 10 | 3 | 2 | common | 0.105263 | partially protected |
| <i>Calophyllum brasiliense</i> | Cambess. | 28 | 3 | 29 | common | 0.276042 | partially protected |
| <i>Calycolpus goetheanus</i> | (Mart. ex DC.) O.Berg | 11 | 3 | 12 | common | 0.657143 | partially protected |
| <i>Calycolpus legrandii</i> | Mattos | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Calycolpus revolutus</i> | (Schauer) O.Berg | 9 | 3 | 7 | common | 0.571429 | partially protected |
| <i>Calycophyllum multiflorum</i> | Griseb. | 9 | 3 | 7 | common | 0.153846 | partially protected |
| <i>Calyptranthes brasiliensis</i> | Spreng. | 5 | 3 | 4 | common | 0.384615 | partially protected |
| <i>Calyptranthes concinna</i> | DC. | 2 | 1 | 3 | form5 | 0 | unprotected |
| <i>Calyptranthes fasciculata</i> | O.Berg | 18 | 3 | 14 | common | 0.444444 | partially protected |
| <i>Calyptranthes lucida</i> | Mart. ex DC. | 22 | 3 | 16 | common | 0.258065 | partially protected |
| <i>Calyptranthes ovalifolia</i> | Cambess. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Calyptranthes pulchella</i> | DC. | 12 | 3 | 11 | common | 0.272727 | partially protected |
| <i>Camarea affinis</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Campomanesia aromatica</i> | (Aubl.) Griseb. | 13 | 3 | 13 | common | 0.30303 | partially protected |
| <i>Campomanesia dichotoma</i> | (O.Berg) Mattos | 9 | 3 | 7 | common | 0.225806 | partially protected |
| <i>Campomanesia eugenoides</i> | (Cambess.) D.Legrand ex Landrum | 9 | 3 | 4 | common | 0.175439 | partially protected |
| <i>Campomanesia guaviroba</i> | (DC.) Kiaersk. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|-------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Campomanesia ilhoensis</i> | Mattos | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Campomanesia pubescens</i> | (Mart. ex DC.) O.Berg | 8 | 3 | 5 | common | 0.222222 | partially protected |
| <i>Campomanesia rufa</i> | (O.Berg) Nied. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Campomanesia sessiliflora</i> | (O.Berg) Mattos | 14 | 3 | 9 | common | 0.091483 | partially protected |
| <i>Campomanesia velutina</i> | (Cambess.) O.Berg | 21 | 3 | 8 | common | 0.136126 | partially protected |
| <i>Campomanesia xanthocarpa</i> | (Mart.) O.Berg | 8 | 3 | 3 | common | 0.083333 | partially protected |
| <i>Capirona decorticans</i> | Spruce | 21 | 3 | 17 | common | 0.434783 | partially protected |
| <i>Capparidastrum frondosum</i> | (Jacq.) Cornejo & Iltis | 13 | 3 | 15 | common | 0.411765 | partially protected |
| <i>Carapa densifolia</i> | Mart. | 23 | 3 | 27 | common | 0.306122 | partially protected |
| <i>Carapa llanorum</i> | Cuatrec. | 11 | 3 | 9 | common | 0.114286 | partially protected |
| <i>Carapa savannarum</i> | Kubitzki | 21 | 3 | 19 | common | 0.235294 | partially protected |
| <i>Carapa tereticaulis</i> | Tul. | 4 | 1 | 5 | form1 | 1 | protected |
| <i>Cardiopetalum calophyllum</i> | Schltdl. | 21 | 3 | 19 | common | 0.208791 | partially protected |
| <i>Cariniana domestica</i> | (Mart.) Miers | 15 | 3 | 13 | common | 0.4 | partially protected |
| <i>Caryocar brasiliense</i> | A.St.-Hil. | 18 | 3 | 14 | common | 0.112782 | partially protected |
| <i>Caryocar coriaceum</i> | Wittm. | 13 | 3 | 6 | common | 0.156134 | partially protected |
| <i>Casearia aculeata</i> | Jacq. | 25 | 3 | 23 | common | 0.220779 | partially protected |
| <i>Casearia arborea</i> | (Rich.) Urb. | 27 | 3 | 25 | common | 0.283951 | partially protected |
| <i>Casearia arguta</i> | Kunth | 2 | 3 | 1 | form6 | 0 | unprotected |
| <i>Casearia commersoniana</i> | Cambess. | 27 | 3 | 40 | common | 0.297674 | partially protected |
| <i>Casearia decandra</i> | Jacq. | 25 | 3 | 22 | common | 0.207547 | partially protected |
| <i>Casearia eichleriana</i> | Sleumer | 9 | 3 | 5 | common | 0.4 | partially protected |
| <i>Casearia gossypiosperma</i> | Briq. | 19 | 3 | 16 | common | 0.15894 | partially protected |
| <i>Casearia grandiflora</i> | Cambess. | 32 | 3 | 34 | common | 0.237643 | partially protected |
| <i>Casearia guianensis</i> | (Aubl.) Urb. | 15 | 3 | 11 | common | 0.5 | partially protected |
| <i>Casearia lasiophylla</i> | Eichler | 8 | 3 | 3 | common | 0.1875 | partially protected |
| <i>Casearia mestrensis</i> | Sleumer | 2 | 1 | 3 | form5 | 0.666667 | partially protected |
| <i>Casearia mollis</i> | Kunth | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Casearia praecox</i> | Griseb. | 2 | 3 | 1 | form6 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|----------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Casearia rufescens</i> | Cambess. | 3 | 3 | 3 | form4 | 0.193548 | partially protected |
| <i>Casearia rupestris</i> | Eichler | 14 | 3 | 9 | common | 0.155405 | partially protected |
| <i>Casearia selloana</i> | Eichler | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Casearia spinescens</i> | (Sw.) Griseb. | 10 | 3 | 10 | common | 0.235294 | partially protected |
| <i>Casearia sylvestris</i> | Sw. | 34 | 3 | 46 | common | 0.148885 | partially protected |
| <i>Casearia tenuipilosa</i> | Sleumer | 8 | 1 | 6 | form1 | | 1 protected |
| <i>Casearia ulmifolia</i> | Vahl ex Vent. | 20 | 3 | 16 | common | 0.138889 | partially protected |
| <i>Casearia zizyphoides</i> | Kunth | 2 | 3 | 2 | form4 | 0.142857 | partially protected |
| <i>Cassia fastuosa</i> | Willd. ex Benth. | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Cassipourea guianensis</i> | Aubl. | 19 | 3 | 18 | common | 0.393939 | partially protected |
| <i>Cathedra rubricaulis</i> | Miers | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Catostemma lemense</i> | Sanoja | 1 | 3 | 1 | form6 | | 1 protected |
| <i>Cecropia concolor</i> | Willd. | 21 | 3 | 18 | common | 0.230769 | partially protected |
| <i>Cecropia distachya</i> | Huber | 17 | 3 | 20 | common | 0.368421 | partially protected |
| <i>Cecropia latiloba</i> | Miq. | 19 | 3 | 20 | common | 0.258065 | partially protected |
| <i>Cecropia obtusa</i> | TrÃƒÂ©cul | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cecropia pachystachya</i> | TrÃƒÂ©cul | 14 | 3 | 10 | common | 0.172414 | partially protected |
| <i>Cecropia saxatilis</i> | Snehl. | 10 | 3 | 6 | common | 0.25 | partially protected |
| <i>Cedrela fissilis</i> | Vell. | 23 | 3 | 17 | common | 0.192308 | partially protected |
| <i>Ceiba glaziovii</i> | (Kuntze) K.Schum. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ceiba pubiflora</i> | (A.St.-Hil.) K.Schum. | 14 | 3 | 6 | common | 0.157895 | partially protected |
| <i>Celtis iguanaea</i> | (Jacq.) Sarg. | 32 | 3 | 27 | common | 0.186508 | partially protected |
| <i>Cenostigma macrophyllum</i> | Tul. | 16 | 3 | 11 | common | 0.149798 | partially protected |
| <i>Cereus albicaulis</i> | (Britton & Rose) Luetzelb. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Cereus fernambucensis</i> | Lem. | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Cereus hexagonus</i> | (L.) Mill. | 4 | 3 | 3 | common | 0.833333 | partially protected |
| <i>Cereus hildmannianus</i> | K.Schum. | 7 | 3 | 4 | common | 0.277778 | partially protected |
| <i>Cereus jamacaru</i> | DC. | 15 | 3 | 10 | common | 0.164384 | partially protected |
| <i>Cestrum alternifolium</i> | (Jacq.) O.E.Schulz | 1 | 3 | 1 | form6 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|-------------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Cestrum laevigatum</i> | Schltdl. | 2 | 1 | 3 | form5 | 0 | unprotected |
| <i>Cestrum mariquitense</i> | Kunth | 17 | 3 | 11 | common | 0.296296 | partially protected |
| <i>Cestrum obovatum</i> | Sendtn. | 8 | 3 | 4 | common | 0.458333 | partially protected |
| <i>Cestrum pedicellatum</i> | Sendtn. | 8 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Cestrum reflexum</i> | Sendtn. | 12 | 3 | 7 | common | 0.25 | partially protected |
| <i>Cestrum retrofractum</i> | Dunal | 5 | 3 | 5 | common | 0.285714 | partially protected |
| <i>Cestrum strigilatum</i> | Ruiz & Pav. | 14 | 3 | 9 | common | 0.225806 | partially protected |
| <i>Chaetocarpus echinocarpus</i> | (Baill.) Ducke | 29 | 3 | 20 | common | 0.341176 | partially protected |
| <i>Chaetocarpus myrsinites</i> | Baill. | 5 | 3 | 4 | common | 0.307692 | partially protected |
| <i>Chaetocarpus schomburgkianus</i> | (Kuntze) Pax & K.Hoffm. | 13 | 3 | 13 | common | 0.5 | partially protected |
| <i>Chamaecrista adenophylla</i> | (Taub.) H.S.Irwin & Barneby | 1 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Chamaecrista bahiae</i> | (H.S.Irwin) H.S.Irwin & Barneby | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Chamaecrista blanchetii</i> | (Benth.) Conc. et al. | 6 | 3 | 4 | common | 0.428571 | partially protected |
| <i>Chamaecrista brachystachya</i> | Conc., L.P. Queiroz & G.P. Lewis | 6 | 3 | 5 | common | 0 | unprotected |
| <i>Chamaecrista bracteolata</i> | (Vogel) H.S.Irwin & Barneby | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Chamaecrista celiae</i> | (H.S.Irwin & Barneby) H.S.Irwin & Barneby | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Chamaecrista chaetostegia</i> | (H.S.Irwin & Barneby) H.S.Irwin & Barneby | 4 | 1 | 2 | form1 | 0.75 | partially protected |
| <i>Chamaecrista confertiformis</i> | (H.S. Irwin & Barneby) Conc., L.P. Queiroz & G.P. | 4 | 1 | 2 | form1 | 0 | unprotected |
| <i>Chamaecrista cytisoides</i> | (Collad.) H.S.Irwin & Barneby | 11 | 3 | 6 | common | 0.2 | partially protected |
| <i>Chamaecrista decora</i> | (H.S. Irwin & Barneby) Conc., L.P. Queiroz & G.P. | 7 | 1 | 3 | form5 | 0 | unprotected |
| <i>Chamaecrista dentata</i> | (Vogel) H.S.Irwin & Barneby | 3 | 3 | 4 | form4 | 0.333333 | partially protected |
| <i>Chamaecrista dumalis</i> | (Hoehne) H.S.Irwin & Barneby | 7 | 3 | 3 | common | 0.2 | partially protected |
| <i>Chamaecrista eitenorum</i> | (H.S.Irwin & Barneby) H.S.Irwin & Barneby | 12 | 3 | 6 | common | 0.166667 | partially protected |
| <i>Chamaecrista ensiformis</i> | (Vell.) H.S.Irwin & Barneby | 9 | 3 | 6 | common | 0.368421 | partially protected |
| <i>Chamaecrista flexuosa</i> | (L.) Greene | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Chamaecrista fulgida</i> | Barneby | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Chamaecrista glaziovii</i> | (Harms) H.S.Irwin & Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Chamaecrista glischrodes</i> | H.S.Irwin & Barneby | 2 | 1 | 3 | form5 | 0.666667 | partially protected |
| <i>Chamaecrista hedsaroides</i> | (Vogel)H.S.Irwin & Barneby | 3 | 3 | 3 | form4 | 0.272727 | partially protected |
| <i>Chamaecrista machaerifolia</i> | (Benth.)H.S.Irwin & Barneby | 1 | 3 | 1 | common | 0.352941 | partially protected |
| <i>Chamaecrista multiseta</i> | (Benth.)H.S.Irwin & Barneby | 8 | 1 | 3 | form1 | 0.121212 | partially protected |
| <i>Chamaecrista nuda</i> | (H.S.Irwin & Barneby) H.S.Irwin & Barneby | 9 | 3 | 6 | form6 | 1 | protected |
| <i>Chamaecrista orbiculata</i> | (Benth.)H.S.Irwin & Barneby | 1 | 1 | 1 | form1 | 0.130081 | partially protected |
| <i>Chamaecrista pachyclada</i> | (Harms)H.S.Irwin & Barneby | 1 | 1 | 1 | form1 | 0.4 | partially protected |
| <i>Chamaecrista phyllostachya</i> | (Benth.)H.S.Irwin & Barneby | 13 | 1 | 7 | form1 | 0.285714 | partially protected |
| <i>Chamaecrista pilicarpa</i> | (Harms)H.S.Irwin & Barneby | 4 | 1 | 2 | form7 | 1 | protected |
| <i>Chamaecrista roraimae</i> | (Benth.)Gleason | 8 | 3 | 6 | common | 0.25 | partially protected |
| <i>Chamaecrista sincorana</i> | (Harms)H.S.Irwin & Barneby | 2 | 3 | 2 | form4 | 0.428571 | partially protected |
| <i>Chamaecrista spinulosa</i> | (H.S.Irwin & Barneby) H.S.Irwin & Barneby | 2 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Chamaecrista ulmea</i> | H.S.Irwin & Barneby | 2 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Chamaecrista zygophylloides</i> | (Taub.) H.S.Irwin & Barneby | 8 | 3 | 5 | common | 0.266667 | partially protected |
| <i>Chaunochiton angustifolium</i> | Sleumer | 13 | 3 | 14 | common | 0.086957 | partially protected |
| <i>Chaunochiton kappleri</i> | (Sagot ex Engl.) Ducke | 8 | 3 | 6 | common | 0.1 | partially protected |
| <i>Cheiloclinium cognatum</i> | (Miers) A.C.Sm. | 25 | 3 | 25 | common | 0.243421 | partially protected |
| <i>Chelyocarpus chuco</i> | (Mart.) H.E.Moore | 5 | 3 | 4 | common | 0.181818 | partially protected |
| <i>Chiococca nitida</i> | Benth. | 8 | 3 | 10 | common | 0.5 | partially protected |
| <i>Chionanthus crassifolius</i> | (Mart.) P.S.Green | 6 | 3 | 4 | common | 0.230769 | partially protected |
| <i>Chionanthus filiformis</i> | (Vell.) P.S.Green | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Chloroleucon acacioides</i> | (Ducke)Barneby & J.W.Grimes | 9 | 3 | 9 | common | 0.470588 | partially protected |
| <i>Chloroleucon foliolosum</i> | (Benth.)G.P.Lewis | 2 | 1 | 1 | form7 | 1 | protected |
| <i>Chloroleucon mangense</i> | (Jacq.)Britton & Rose | 3 | 3 | 1 | form6 | 0.111111 | partially protected |
| <i>Chloroleucon tenuiflorum</i> | (Benth.)Barneby & J.W.Grimes | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Chomelia anisomeris</i> | MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Chomelia obtusa</i> | Cham. & Schltdl. | 23 | 3 | 18 | common | 0.179641 | partially protected |
| <i>Chomelia paniculata</i> | (Bartl. ex DC.) Steyermark | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Chomelia parviflora</i> | (MÃƒÂ¼ll.Arg.) MÃƒÂ¼ll.Arg. | 11 | 3 | 5 | common | 0.2 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|---------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Chomelia pohliana</i> | MÃƒÂ½ll.Arg. | 11 | 3 | 3 | common | 0 | unprotected |
| <i>Chomelia pubescens</i> | Cham. & Schltdl. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Chomelia spinosa</i> | Jacq. | 5 | 3 | 6 | common | 0.25 | partially protected |
| <i>Chomelia tenuiflora</i> | Benth. | 14 | 3 | 15 | common | 0.5 | partially protected |
| <i>Christiana africana</i> | DC. | 13 | 3 | 13 | common | 0.2 | partially protected |
| <i>Christiana mennegae</i> | (Jans.-Jac. & Westra) Kubitzki | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Chromolaena odorata</i> | (L.) R.M.King & H.Rob. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Chronopappus bifrons</i> | (DC. ex Pers.) DC. | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Chrysobalanus icaco</i> | L. | 12 | 3 | 9 | common | 0.333333 | partially protected |
| <i>Chrysolaena platensis</i> | (Spreng.) H.Rob. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Chrysophyllum arenarium</i> | AllemafÃ±o | 4 | 3 | 3 | common | 0.375 | partially protected |
| <i>Chrysophyllum gonocarpum</i> | (Mart. & Eichler ex Miq.) Engl. | 4 | 3 | 4 | common | 0 | unprotected |
| <i>Chrysophyllum januariense</i> | Eichler | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Chrysophyllum marginatum</i> | (Hook. & Arn.) Radlk. | 15 | 3 | 12 | common | 0.148148 | partially protected |
| <i>Chrysophyllum rufum</i> | Mart. | 2 | 1 | 3 | form5 | 0 | unprotected |
| <i>Cinnamomum erythropus</i> | (Nees & Mart.) Kosterm. | 4 | 3 | 3 | common | 0.44 | partially protected |
| <i>Cinnamomum haussknechtii</i> | (Mez) Kosterm. | 8 | 3 | 3 | common | 0.26087 | partially protected |
| <i>Cinnamomum sellowianum</i> | (Nees & Mart.) Kosterm. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cinnamomum tomentulosum</i> | Kosterm. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cinnamomum triplinerve</i> | (Ruiz & Pav.) Kosterm. | 11 | 3 | 9 | common | 0.75 | partially protected |
| <i>Clethra scabra</i> | Pers. | 9 | 3 | 7 | common | 0.346939 | partially protected |
| <i>Clusia amabilis</i> | Maguire | 2 | 3 | 1 | form6 | 0.666667 | partially protected |
| <i>Clusia brachystyla</i> | Maguire | 4 | 3 | 3 | common | 0.571429 | partially protected |
| <i>Clusia burchellii</i> | Engl. | 4 | 3 | 1 | form2 | 0.3 | partially protected |
| <i>Clusia columnaris</i> | Engl. | 22 | 3 | 25 | common | 0.421053 | partially protected |
| <i>Clusia crassifolia</i> | Planch. & Triana | 3 | 3 | 3 | form4 | 1 | protected |
| <i>Clusia criuva</i> | Cambess. | 11 | 3 | 7 | common | 0.293333 | partially protected |
| <i>Clusia dardanoi</i> | G.Mariz & Maguire | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Clusia diamantina</i> | Bittrich | 1 | 1 | 2 | form5 | 0.333333 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|-------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Clusia fockeana</i> | Miq. | 8 | 3 | 8 | common | 0.714286 | partially protected |
| <i>Clusia fragrans</i> | Gardner | 6 | 3 | 4 | common | 0.2 | partially protected |
| <i>Clusia gardneri</i> | Planch. & Triana | 4 | 3 | 1 | form2 | 0.6 | partially protected |
| <i>Clusia grandiflora</i> | Splitg. | 5 | 3 | 4 | common | 0.5 | partially protected |
| <i>Clusia hilariana</i> | Schltdl. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Clusia hoffmannseggiana</i> | Schltdl. | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Clusia huberi</i> | Pipoly | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Clusia melchiori</i> | Gleason | 4 | 3 | 3 | common | 0.3 | partially protected |
| <i>Clusia nemorosa</i> | G.Mey. | 23 | 3 | 25 | common | 0.293333 | partially protected |
| <i>Clusia obdeltifolia</i> | Bittrich | 3 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Clusia obovata</i> | (Spruce ex Planch. & Triana) Pipoly | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Clusia panapanari</i> | (Aubl.) Choisy | 15 | 3 | 16 | common | 0.352941 | partially protected |
| <i>Clusia paralicola</i> | G.Mariz | 5 | 3 | 4 | common | 0.2 | partially protected |
| <i>Clusia pernambucensis</i> | G.Mariz | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Clusia polysepala</i> | Engl. | 6 | 1 | 5 | form1 | 0.5 | partially protected |
| <i>Clusia pusilla</i> | Steyermark | 5 | 3 | 4 | common | 0.636364 | partially protected |
| <i>Clusia riedeliana</i> | Engl. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Clusia savannarum</i> | Maguire | 2 | 3 | 2 | form4 | 1 | protected |
| <i>Clusia schomburgkiana</i> | (Planch. & Triana) Benth. ex Engl. | 5 | 3 | 7 | common | 0.5 | partially protected |
| <i>Clusia sellowiana</i> | Schltdl. | 2 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Clusia tabulamontana</i> | Maguire | 3 | 3 | 2 | form4 | 0.666667 | partially protected |
| <i>Clusia weddelliana</i> | Planch. & Triana | 16 | 3 | 8 | common | 0.333333 | partially protected |
| <i>Cnidoscolus bahianus</i> | (Ule) Pax & K.Hoffm. | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Cnidoscolus vitifolius</i> | (Mill.) Pohl | 15 | 3 | 6 | common | 0.183099 | partially protected |
| <i>Coccoloba alnifolia</i> | Casar. | 10 | 3 | 8 | common | 0.148148 | partially protected |
| <i>Coccoloba caracasana</i> | Meisn. | 5 | 3 | 2 | common | 0.05 | partially protected |
| <i>Coccoloba cujabensis</i> | Wedd. | 8 | 3 | 4 | common | 0.173913 | partially protected |
| <i>Coccoloba declinata</i> | (Vell.) Mart. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Coccoloba densifrons</i> | Mart. ex Meisn. | 3 | 3 | 3 | form4 | 0.25 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Coccoloba excelsa</i> | Benth. | 19 | 3 | 20 | common | 0.5 | partially protected |
| <i>Coccoloba laevis</i> | Casar. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Coccoloba latifolia</i> | Poir. | 9 | 3 | 8 | common | 0.307692 | partially protected |
| <i>Coccoloba llewelynii</i> | R.A.Howard | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Coccoloba lucidula</i> | Benth. | 15 | 3 | 11 | common | 0.35 | partially protected |
| <i>Coccoloba marginata</i> | Benth. | 20 | 3 | 18 | common | 0.148148 | partially protected |
| <i>Coccoloba mollis</i> | Casar. | 29 | 3 | 23 | common | 0.127563 | partially protected |
| <i>Coccoloba obtusifolia</i> | Jacq. | 7 | 3 | 7 | common | 0.125 | partially protected |
| <i>Coccoloba orinocana</i> | R.A.Howard | 4 | 3 | 1 | form2 | 0.142857 | partially protected |
| <i>Coccoloba paraguariensis</i> | Lindau | 3 | 3 | 4 | form4 | 0.25 | partially protected |
| <i>Coccoloba ramosissima</i> | Wedd. | 12 | 3 | 10 | common | 0.297297 | partially protected |
| <i>Coccoloba salicifolia</i> | Wedd. | 3 | 3 | 3 | form4 | 0.214286 | partially protected |
| <i>Coccoloba savannarum</i> | Standl. | 3 | 3 | 3 | form4 | 0.2 | partially protected |
| <i>Cochlospermum orinocense</i> | (Kunth) Steud. | 21 | 3 | 22 | common | 0.227273 | partially protected |
| <i>Cochlospermum regium</i> | (Schrank) Pilg. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Cochlospermum vitifolium</i> | (Willd.) Spreng. | 28 | 3 | 33 | common | 0.227723 | partially protected |
| <i>Colubrina glandulosa</i> | G.Perkins | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Combretum duarteanum</i> | Cambess. | 22 | 3 | 10 | common | 0.141304 | partially protected |
| <i>Combretum lanceolatum</i> | Pohl ex Eichler | 25 | 3 | 21 | common | 0.238636 | partially protected |
| <i>Combretum leprosum</i> | Mart. | 22 | 3 | 17 | common | 0.14717 | partially protected |
| <i>Combretum mellifluum</i> | Eichler | 18 | 3 | 10 | common | 0.190789 | partially protected |
| <i>Combretum monetaria</i> | Mart. | 12 | 3 | 4 | common | 0.266667 | partially protected |
| <i>Combretum rotundifolium</i> | Rich. | 21 | 3 | 26 | common | 0.133333 | partially protected |
| <i>Commiphora leptophloeos</i> | (Mart.) J.B.Gillett | 9 | 3 | 9 | common | 0.263158 | partially protected |
| <i>Compsoneura ulei</i> | Warb. ex Pilg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Conchocarpus longifolius</i> | (A.St.-Hil.) Kallunki & Pirani | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Connarus detersus</i> | Planch. | 2 | 3 | 2 | form4 | 0.571429 | partially protected |
| <i>Connarus incomptus</i> | Planch. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Connarus perrottetii</i> | (DC.) Planch. | 23 | 3 | 19 | common | 0.390625 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|-------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Connarus suberosus</i> | Planch. | 22 | 3 | 14 | common | 0.120165 | partially protected |
| <i>Connarus venezuelanus</i> | Baill. | 5 | 3 | 4 | common | 0 | unprotected |
| <i>Conostegia speciosa</i> | Naudin | 1 | 3 | 2 | form4 | 0 | unprotected |
| <i>Copaifera arenicola</i> | (Ducke) J.Costa & L.P.Queiroz | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Copaifera coriacea</i> | Mart. | 7 | 3 | 4 | common | 0.555556 | partially protected |
| <i>Copaifera duckei</i> | Dwyer | 15 | 3 | 12 | common | 0.310345 | partially protected |
| <i>Copaifera elliptica</i> | Mart. | 5 | 3 | 5 | common | 0.155556 | partially protected |
| <i>Copaifera langsdorffii</i> | Desf. | 23 | 3 | 19 | common | 0.11733 | partially protected |
| <i>Copaifera luetzelburgii</i> | Harms | 12 | 3 | 6 | common | 0.10625 | partially protected |
| <i>Copaifera magnifolia</i> | Dwyer | 8 | 3 | 1 | form2 | 0.225806 | partially protected |
| <i>Copaifera malmei</i> | Harms | 13 | 3 | 5 | common | 0.090323 | partially protected |
| <i>Copaifera martii</i> | Hayne | 15 | 3 | 11 | common | 0.34375 | partially protected |
| <i>Copaifera oblongifolia</i> | Mart. ex Hayne | 20 | 3 | 11 | common | 0.167059 | partially protected |
| <i>Copaifera pubiflora</i> | Benth. | 7 | 3 | 3 | common | 0.104167 | partially protected |
| <i>Copaifera sabulicola</i> | J.A.S.Costa & L.P. Queiroz | 8 | 3 | 4 | common | 0.15625 | partially protected |
| <i>Copernicia alba</i> | Morong | 4 | 3 | 4 | common | 0.2 | partially protected |
| <i>Copernicia prunifera</i> | (Mill.) H.E.Moore | 9 | 3 | 5 | common | 0.222222 | partially protected |
| <i>Cordia bicolor</i> | A.DC. | 18 | 3 | 23 | common | 0.305556 | partially protected |
| <i>Cordia curassavica</i> | (Jacq.) Roem. & Schult. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Cordia exaltata</i> | Lam. | 16 | 3 | 18 | common | 0.454545 | partially protected |
| <i>Cordia glabrata</i> | (Mart.) A.DC. | 19 | 3 | 14 | common | 0.165049 | partially protected |
| <i>Cordia insignis</i> | Cham. | 8 | 3 | 6 | common | 0.156627 | partially protected |
| <i>Cordia leucocephala</i> | Moric. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cordia nodosa</i> | Lam. | 21 | 3 | 27 | common | 0.307692 | partially protected |
| <i>Cordia polyccephala</i> | (Lam.) I.M.Johnst. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cordia rufescens</i> | A.DC. | 13 | 3 | 6 | common | 0.09 | partially protected |
| <i>Cordia scabrifolia</i> | A.DC. | 16 | 3 | 17 | common | 0.357143 | partially protected |
| <i>Cordia sellowiana</i> | Cham. | 27 | 3 | 20 | common | 0.181818 | partially protected |
| <i>Cordia superba</i> | Cham. | 16 | 3 | 9 | common | 0.078431 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Cordia trichotoma</i> | (Vell.) ArrÃƒÂ¡fÃ¡b. ex Steud. | 20 | 3 | 10 | common | 0.073529 | partially protected |
| <i>Cordiera concolor</i> | (Cham.) Kuntze | 18 | 3 | 12 | common | 0.102941 | partially protected |
| <i>Cordiera elliptica</i> | (Cham.) Kuntze | 21 | 3 | 10 | common | 0.174168 | partially protected |
| <i>Cordiera macrophylla</i> | (K.Schum.) Kuntze | 15 | 3 | 8 | common | 0.174419 | partially protected |
| <i>Cordiera myrciifolia</i> | (K.Schum.) Perss. & Delprete | 18 | 3 | 10 | common | 0.275 | partially protected |
| <i>Cordiera rigida</i> | (K.Schum.) Kuntze | 18 | 3 | 7 | common | 0.134503 | partially protected |
| <i>Cordiera sessilis</i> | (Vell.) Kuntze | 22 | 3 | 17 | common | 0.134111 | partially protected |
| <i>Couepia belemii</i> | Prance | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Couepia grandiflora</i> | (Mart. & Zucc.) Benth. ex Hook.f. | 22 | 3 | 13 | common | 0.145 | partially protected |
| <i>Couepia multiflora</i> | Benth. | 3 | 3 | 3 | form4 | 0.153846 | partially protected |
| <i>Couepia ovalifolia</i> | (Schott) Benth. ex Hook.f. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Couepia schottii</i> | Fritsch | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Couepia uiti</i> | (Mart. & Zucc.) Benth. ex Hook.f. | 14 | 3 | 7 | common | 0.212121 | partially protected |
| <i>Couepia venosa</i> | Prance | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Couma rigida</i> | MÃƒÂ¢fÃ¼ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Coursetia ferruginea</i> | (Kunth)Lavin | 6 | 3 | 2 | common | 0.055556 | partially protected |
| <i>Coussarea hydrangeifolia</i> | (Benth.) Benth. & Hook.f. ex MÃƒÂ¢fÃ¼ll.Arg. | 23 | 3 | 15 | common | 0.135231 | partially protected |
| <i>Coussarea obscura</i> | MÃƒÂ¢fÃ¼ll.Arg. | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Coussarea paniculata</i> | (Vahl) Standl. | 14 | 3 | 13 | common | 0.363636 | partially protected |
| <i>Coussarea platyphylla</i> | MÃƒÂ¢fÃ¼ll.Arg. | 8 | 3 | 4 | common | 0.375 | partially protected |
| <i>Coussarea violacea</i> | Aubl. | 3 | 1 | 2 | form5 | 1 | protected |
| <i>Coutarea hexandra</i> | (Jacq.) K.Schum. | 33 | 3 | 25 | common | 0.193277 | partially protected |
| <i>Crateva tapia</i> | L. | 23 | 3 | 27 | common | 0.173077 | partially protected |
| <i>Cratylia argentea</i> | (Desv.)Kuntze | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cratylia mollis</i> | Benth. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Crescentia amazonica</i> | Ducke | 4 | 1 | 5 | form1 | 0 | unprotected |
| <i>Crotalaria holosericea</i> | Nees & Mart. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Croton argyrophyllus</i> | Kunth | 9 | 3 | 8 | common | 0.181818 | partially protected |
| <i>Croton blanchetianus</i> | Baill. | 6 | 3 | 3 | common | 0.263158 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Croton campestris</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Croton celtidifolius</i> | Baill. | 3 | 3 | 3 | form4 | 0.25 | partially protected |
| <i>Croton draconoides</i> | MÃƒÂ¼ll.Arg. | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Croton essequiboensis</i> | Klotzsch | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Croton floribundus</i> | Spreng. | 13 | 3 | 4 | common | 0.172414 | partially protected |
| <i>Croton gossypiifolius</i> | Vahl | 11 | 3 | 11 | common | 0.225 | partially protected |
| <i>Croton heliotropiifolius</i> | Kunth | 8 | 3 | 6 | common | 0.133333 | partially protected |
| <i>Croton matourensis</i> | Aubl. | 20 | 3 | 24 | common | 0.384615 | partially protected |
| <i>Croton nepetifolius</i> | Baill. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Croton orinocensis</i> | MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Croton palanostigma</i> | Klotzsch | 15 | 3 | 12 | common | 0.181818 | partially protected |
| <i>Croton pedicellatus</i> | Kunth | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Croton sonderianus</i> | MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Croton tricolor</i> | Klotzsch ex Baill. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Croton urticifolius</i> | Lam. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Croton yavitenensis</i> | Croizat | 7 | 3 | 4 | common | 0 | unprotected |
| <i>Cupania bracteosa</i> | Radlk. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Cupania emarginata</i> | Cambess. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Cupania impressinervia</i> | Acev.-Rodr. | 9 | 3 | 7 | common | 0.318182 | partially protected |
| <i>Cupania ludwigii</i> | Somner & Ferrucci | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Cupania paniculata</i> | Cambess. | 9 | 3 | 5 | common | 0.207792 | partially protected |
| <i>Cupania platycarpa</i> | Radlk. | 7 | 3 | 5 | common | 0.166667 | partially protected |
| <i>Cupania racemosa</i> | (Vell.) Radlk. | 5 | 3 | 5 | common | 0.090909 | partially protected |
| <i>Cupania rubiginosa</i> | (Poir.) Radlk. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Cupania rugosa</i> | Radlk. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Cupania vernalis</i> | Cambess. | 19 | 3 | 12 | common | 0.169014 | partially protected |
| <i>Cuphea antisiphilitica</i> | Kunth | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cuphea campestris</i> | Mart. ex Koehne | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Cuphea ericoides</i> | Cham. & Schldl. | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Curatella americana</i> | L. | 33 | 3 | 33 | common | 0.144351 | partially protected |
| <i>Cyathea microdonta</i> | (Desv.) Domin | 28 | 3 | 21 | common | 0.313725 | partially protected |
| <i>Cyathea villosa</i> | Humb. & Bonpl. ex Willd. | 8 | 3 | 6 | common | 0.363636 | partially protected |
| <i>Cybianthus detergens</i> | Mart. | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Cybianthus fulvopulverulentus</i> | (Mez) G.Agostini | 11 | 3 | 13 | common | 0.428571 | partially protected |
| <i>Cybianthus gardneri</i> | (A.DC.) G.Agostini | 13 | 3 | 4 | common | 0.283019 | partially protected |
| <i>Cybianthus goyazensis</i> | Mez | 3 | 3 | 1 | form6 | 0.25 | partially protected |
| <i>Cybianthus minutiflorus</i> | Mez | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Cybianthus quelchii</i> | (N.E.Br.) G.Agostini | 3 | 3 | 3 | form4 | 1 | protected |
| <i>Cybianthus resinosus</i> | Mez | 7 | 3 | 6 | common | 0.666667 | partially protected |
| <i>Cybianthus venezuelanus</i> | Mez | 13 | 3 | 8 | common | 0.333333 | partially protected |
| <i>Cybistax antisiphilitica</i> | (Mart.) Mart. | 25 | 3 | 21 | common | 0.136119 | partially protected |
| <i>Cynometra bauhiniifolia</i> | Benth. | 19 | 3 | 17 | common | 0.178571 | partially protected |
| <i>Cynophalla flexuosa</i> | (L.) J.Presl | 23 | 3 | 20 | common | 0.211268 | partially protected |
| <i>Cynophalla hastata</i> | (Jacq.) J.Presl | 12 | 3 | 8 | common | 0.285714 | partially protected |
| <i>Cynophalla retusa</i> | (Griseb.) Cornejo & Iltis | 10 | 3 | 7 | common | 0.294118 | partially protected |
| <i>Cyrilla racemiflora</i> | L. | 11 | 3 | 12 | common | 0.6 | partially protected |
| <i>Cyrtocarpa velutinifolia</i> | (R.S.Cowan) J.D.Mitch. & Daly | 4 | 3 | 2 | common | 0.083333 | partially protected |
| <i>Cyrtocymura scorpioides</i> | (Lam.) H.Rob. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Dacryodes glabra</i> | (Steyer.) Cuatrec. | 5 | 3 | 4 | common | 0.25 | partially protected |
| <i>Dacryodes microcarpa</i> | Cuatrec. | 16 | 3 | 16 | common | 0.25 | partially protected |
| <i>Dacryodes roraimensis</i> | Cuatrec. | 3 | 1 | 2 | form5 | 0.75 | partially protected |
| <i>Dahlstedtia araripensis</i> | (Benth.) M.J. Silva & A.M.G. Azevedo | 5 | 3 | 5 | common | 0.333333 | partially protected |
| <i>Dalbergia acuta</i> | Benth. | 8 | 3 | 3 | common | 0.185185 | partially protected |
| <i>Dalbergia brasiliensis</i> | Vogel | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Dalbergia cearensis</i> | Ducke | 6 | 3 | 5 | common | 0.4 | partially protected |
| <i>Dalbergia densiflora</i> | (Benth.) Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Dalbergia glaucescens</i> | (Benth.) Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Dalbergia miscolobium</i> | Benth. | 23 | 3 | 14 | common | 0.119093 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Dalbergia spruceana</i> | (Benth.) Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Daphnopsis fasciculata</i> | (Meisn.) Nevling | 12 | 3 | 6 | common | 0.175926 | partially protected |
| <i>Daphnopsis filipedunculata</i> | Nevling & Barringer | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Daphnopsis longipedunculata</i> | Gilg ex Domke | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Davilla elliptica</i> | A.St.-Hil. | 23 | 3 | 17 | common | 0.133501 | partially protected |
| <i>Davilla grandiflora</i> | A.St.-Hil. & Tul. | 20 | 3 | 13 | common | 0.146341 | partially protected |
| <i>Davilla nitida</i> | (Vahl) Kubitzki | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Dendropanax cuneatus</i> | (DC.) Decne. & Planch. | 19 | 3 | 10 | common | 0.214286 | partially protected |
| <i>Dialium guianense</i> | (Aubl.) Sandwith | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Dialypetalanthus fuscescens</i> | Kuhlm. | 13 | 3 | 10 | common | 0.5 | partially protected |
| <i>Dilodendron bipinnatum</i> | Radlk. | 15 | 3 | 13 | common | 0.127701 | partially protected |
| <i>Dimorphandra gardneriana</i> | Tul. | 16 | 3 | 13 | common | 0.17126 | partially protected |
| <i>Dimorphandra mollis</i> | Benth. | 20 | 3 | 13 | common | 0.124038 | partially protected |
| <i>Diodella apiculata</i> | (Willd. ex Roem. & Schult.) Delporte | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Diospyros coccolobifolia</i> | Mart. ex Miq. | 15 | 3 | 6 | common | 0.156379 | partially protected |
| <i>Diospyros guianensis</i> | (Aubl.) GÃƒÂ¶cke | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Diospyros hispida</i> | A.DC. | 23 | 3 | 17 | common | 0.141579 | partially protected |
| <i>Diospyros inconstans</i> | Jacq. | 19 | 3 | 15 | common | 0.149533 | partially protected |
| <i>Diospyros sericea</i> | A.DC. | 18 | 3 | 14 | common | 0.174129 | partially protected |
| <i>Diospyros tetrandra</i> | Hiern | 6 | 1 | 8 | form1 | 0 | unprotected |
| <i>Diplotropis brasiliensis</i> | (Tul.) Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Dipteryx alata</i> | Vogel | 22 | 3 | 15 | common | 0.1261 | partially protected |
| <i>Dipteryx lacunifera</i> | Ducke | 5 | 1 | 3 | form1 | 0 | unprotected |
| <i>Diptychandra aurantiaca</i> | Tul. | 20 | 3 | 17 | common | 0.119835 | partially protected |
| <i>Dodonaea viscosa</i> | (L.) Jacq. | 10 | 3 | 10 | common | 0.3 | partially protected |
| <i>Drimys brasiliensis</i> | Miers | 9 | 3 | 5 | common | 0.363636 | partially protected |
| <i>Duguetia furfuracea</i> | (A.St.-Hil.) Saff. | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Duguetia gardneriana</i> | Mart. | 6 | 3 | 5 | common | 0.375 | partially protected |
| <i>Duguetia lanceolata</i> | A.St.-Hil. | 9 | 3 | 5 | common | 0.108108 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|---------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Duguetia spixiana</i> | Mart. | 17 | 3 | 12 | common | 0.4375 | partially protected |
| <i>Duguetia stelechantha</i> | (Diels) R.E.Fr. | 16 | 3 | 12 | common | 0 | unprotected |
| <i>Duguetia yeshidan</i> | Sandwith | 9 | 3 | 8 | common | 0.6 | partially protected |
| <i>Duroia duckei</i> | Huber | 16 | 3 | 17 | common | 0.25 | partially protected |
| <i>Duroia micrantha</i> | (Ladbr.) Zarucchi & J.H.Kirkbr. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ecclinusa dumetorum</i> | (Baehni) T.D.Penn. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ecclinusa ramiflora</i> | Mart. | 21 | 3 | 20 | common | 0.433333 | partially protected |
| <i>Elaeoluma schomburgkiana</i> | (Miq.) Baill. | 14 | 1 | 13 | form1 | 0.5 | partially protected |
| <i>Emmotum acuminatum</i> | (Benth.) Miers | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Emmotum affine</i> | Miers | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Emmotum fagifolium</i> | Desv. ex Ham. | 3 | 1 | 3 | form5 | 0.6 | partially protected |
| <i>Emmotum harleyi</i> | R.Duno | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Emmotum nitens</i> | (Benth.) Miers | 23 | 3 | 19 | common | 0.15738 | partially protected |
| <i>Endlicheria paniculata</i> | (Spreng.) J.F.Macbr. | 20 | 3 | 12 | common | 0.277778 | partially protected |
| <i>Enterolobium contortisiliquum</i> | (Vell.) Morong | 22 | 3 | 19 | common | 0.102273 | partially protected |
| <i>Enterolobium gummiferum</i> | (Mart.) J.F.Macbr. | 22 | 3 | 13 | common | 0.118902 | partially protected |
| <i>Eperua bijuga</i> | Mart. ex Benth. | 8 | 3 | 7 | common | 0.833333 | partially protected |
| <i>Ephedranthus parviflorus</i> | S.Moore | 16 | 3 | 10 | common | 0.404762 | partially protected |
| <i>Ephedranthus pisocarpus</i> | R.E.Fr. | 4 | 3 | 3 | common | 0.235294 | partially protected |
| <i>Eremanthus arboreus</i> | (Gardner) MacLeish | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Eremanthus argenteus</i> | MacLeish & Schumacher | 3 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Eremanthus brasiliensis</i> | (Gardner) MacLeish | 3 | 3 | 1 | form6 | 0.125 | partially protected |
| <i>Eremanthus capitatus</i> | (Spreng.) MacLeish | 11 | 3 | 9 | common | 0.1875 | partially protected |
| <i>Eremanthus cinctus</i> | Baker | 8 | 3 | 3 | common | 0.069767 | partially protected |
| <i>Eremanthus crotonoides</i> | (DC.) Sch.Bip. | 3 | 3 | 3 | form4 | 0.387097 | partially protected |
| <i>Eremanthus elaeagnus</i> | (Mart. ex DC.) Sch.Bip. | 8 | 3 | 4 | common | 0.103448 | partially protected |
| <i>Eremanthus erythropappus</i> | (DC.) MacLeish | 8 | 3 | 4 | common | 0.146552 | partially protected |
| <i>Eremanthus glomerulatus</i> | Less. | 12 | 3 | 6 | common | 0.165266 | partially protected |
| <i>Eremanthus goyazensis</i> | (Gardner) Sch.Bip. | 12 | 3 | 4 | common | 0.288288 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Eremanthus incanus</i> | (Less.) Less. | 12 | 3 | 6 | common | 0.135802 | partially protected |
| <i>Eremanthus leucodendron</i> | Mattf. | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Eremanthus mattogrossensis</i> | Kuntze | 15 | 3 | 6 | common | 0.166667 | partially protected |
| <i>Eremanthus pohlii</i> | (Baker) MacLeish | 3 | 3 | 2 | form4 | 0.466667 | partially protected |
| <i>Eremanthus polycephalus</i> | (DC.) MacLeish | 7 | 3 | 4 | common | 0.157895 | partially protected |
| <i>Eremanthus rondoniensis</i> | MacLeish & H.Schumach. | 2 | 3 | 3 | form4 | 0.428571 | partially protected |
| <i>Eremanthus uniflorus</i> | MacLeish & H.Schumach. | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Eremanthus veadeiroensis</i> | H.Rob. | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Eriope exaltata</i> | Harley | 3 | 3 | 2 | form4 | 0.571429 | partially protected |
| <i>Eriope latifolia</i> | (Mart. ex Benth.) Harley | 7 | 3 | 4 | common | 0.25 | partially protected |
| <i>Eriotheca gracilipes</i> | (K.Schum.) A.Robyns | 20 | 3 | 11 | common | 0.109785 | partially protected |
| <i>Eriotheca macrophylla</i> | (K.Schum.) A.Robyns | 7 | 1 | 4 | form1 | 0 | unprotected |
| <i>Eriotheca parvifolia</i> | (Mart. & Zucc.) A.Robyns | 10 | 3 | 2 | common | 0.2 | partially protected |
| <i>Eriotheca pubescens</i> | (Mart. & Zucc.) Schott & Endl. | 14 | 3 | 9 | common | 0.131635 | partially protected |
| <i>Eriotheca roseorum</i> | (Cuatrec.) A.Robyns | 7 | 3 | 4 | common | 0.066667 | partially protected |
| <i>Erythrina dominguezii</i> | Hassl. | 11 | 3 | 6 | common | 0.321429 | partially protected |
| <i>Erythrina fusca</i> | Lour. | 8 | 3 | 6 | common | 0.230769 | partially protected |
| <i>Erythrociton fallax</i> | Kallunki | 6 | 1 | 5 | form1 | 0.166667 | partially protected |
| <i>Erythroxylum affine</i> | A.St.-Hil. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Erythroxylum ambiguum</i> | Peyr. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Erythroxylum andrei</i> | Plowman | 3 | 3 | 4 | form4 | 0 | unprotected |
| <i>Erythroxylum anguifugum</i> | Mart. | 22 | 3 | 14 | common | 0.205882 | partially protected |
| <i>Erythroxylum arrojadoi</i> | O.E.Schulz | 5 | 3 | 2 | common | 0.142857 | partially protected |
| <i>Erythroxylum ayrtonianum</i> | Loiola & M.F.Sales | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Erythroxylum barbatum</i> | O.E.Schulz | 9 | 3 | 7 | common | 0.5625 | partially protected |
| <i>Erythroxylum bezerrae</i> | Plowman | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Erythroxylum buxus</i> | Peyr. | 3 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Erythroxylum citrifolium</i> | A.St.-Hil. | 27 | 3 | 29 | common | 0.185185 | partially protected |
| <i>Erythroxylum columbinum</i> | Mart. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Erythroxylum cuneifolium</i> | (Mart.) O.E.Schulz | 14 | 3 | 13 | common | 0.181818 | partially protected |
| <i>Erythroxylum daphnites</i> | Mart. | 19 | 3 | 17 | common | 0.189055 | partially protected |
| <i>Erythroxylum deciduum</i> | A.St.-Hil. | 23 | 3 | 16 | common | 0.1703 | partially protected |
| <i>Erythroxylum engleri</i> | O.E.Schulz | 15 | 3 | 5 | common | 0.144509 | partially protected |
| <i>Erythroxylum foetidum</i> | Plowman | 2 | 3 | 1 | form6 | 0 | unprotected |
| <i>Erythroxylum hamigerum</i> | O.E.Schulz | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Erythroxylum havanense</i> | Jacq. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Erythroxylum impressum</i> | O.E.Schulz | 4 | 3 | 1 | form2 | 0.090909 | partially protected |
| <i>Erythroxylum laetevirens</i> | O.E.Schulz | 6 | 3 | 3 | common | 0.25 | partially protected |
| <i>Erythroxylum ligustrinum</i> | DC. | 8 | 3 | 7 | common | 0.578947 | partially protected |
| <i>Erythroxylum maracasense</i> | Plowman | 3 | 1 | 2 | form5 | 0.4 | partially protected |
| <i>Erythroxylum mucronatum</i> | Benth. | 20 | 3 | 19 | common | 0.444444 | partially protected |
| <i>Erythroxylum oreophilum</i> | (O.E.Schulz ex Pilg.) Steyermark & Maguire | 2 | 3 | 2 | form4 | 0.25 | partially protected |
| <i>Erythroxylum orinocense</i> | Kunth | 5 | 3 | 4 | common | 0.090909 | partially protected |
| <i>Erythroxylum passerinum</i> | Mart. | 10 | 3 | 8 | common | 0.230769 | partially protected |
| <i>Erythroxylum pelleterianum</i> | A.St.-Hil. | 11 | 3 | 3 | common | 0.104167 | partially protected |
| <i>Erythroxylum revolutum</i> | Mart. | 5 | 3 | 2 | common | 0.166667 | partially protected |
| <i>Erythroxylum rimosum</i> | O.E.Schulz | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Erythroxylum roraimae</i> | Klotzsch ex O.E.Schulz | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Erythroxylum rufum</i> | Cav. | 29 | 3 | 23 | common | 0.169231 | partially protected |
| <i>Erythroxylum santosii</i> | Plowman | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Erythroxylum schomburgkii</i> | Peyr. | 3 | 3 | 2 | form4 | 0.727273 | partially protected |
| <i>Erythroxylum squamatum</i> | Sw. | 23 | 3 | 19 | common | 0.347826 | partially protected |
| <i>Erythroxylum steyermarkii</i> | Plowman | 3 | 3 | 1 | form6 | 0.125 | partially protected |
| <i>Erythroxylum suberosum</i> | A.St.-Hil. | 33 | 3 | 33 | common | 0.163711 | partially protected |
| <i>Erythroxylum subracemosum</i> | Turcz. | 23 | 3 | 13 | common | 0.210526 | partially protected |
| <i>Erythroxylum subrotundum</i> | A.St.-Hil. | 9 | 3 | 5 | common | 0.35 | partially protected |
| <i>Erythroxylum tortuosum</i> | Mart. | 19 | 3 | 12 | common | 0.139503 | partially protected |
| <i>Erythroxylum ulei</i> | O.E.Schulz | 7 | 3 | 5 | common | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|-------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Erythroxylum vaccinifolium</i> | Mart. | 13 | 3 | 6 | common | 0.26 | partially protected |
| <i>Erythroxylum vernicosum</i> | O.E.Schulz | 9 | 3 | 8 | common | 0.333333 | partially protected |
| <i>Escallonia chlorophylla</i> | Cham. & Schldl. | 1 | 1 | 2 | form5 | 1 | protected |
| <i>Escallonia hispida</i> | (Vell.) Sleumer | 1 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Escallonia reticulata</i> | Sleumer | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Eschweilera nana</i> | (O.Berg) Miers | 20 | 3 | 8 | common | 0.11583 | partially protected |
| <i>Eschweilera ovata</i> | (Cambess.) Mart. ex Miers | 19 | 3 | 19 | common | 0.4375 | partially protected |
| <i>Eschweilera subcordata</i> | S.A.Mori | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eschweilera wachenheimii</i> | (Benoist) Sandwith | 7 | 3 | 7 | common | 1 | protected |
| <i>Esenbeckia febrifuga</i> | (A.St.-Hil.) A.Juss. ex Mart. | 13 | 3 | 2 | common | 0.2 | partially protected |
| <i>Esenbeckia grandiflora</i> | Mart. | 17 | 3 | 13 | common | 0.213115 | partially protected |
| <i>Esenbeckia irwiniana</i> | Kaastra | 3 | 1 | 3 | form5 | 0.5 | partially protected |
| <i>Esenbeckia oligantha</i> | Kaastra | 8 | 3 | 3 | common | 0.5 | partially protected |
| <i>Euceraea nitida</i> | Mart. | 5 | 3 | 8 | common | 1 | protected |
| <i>Eugenia astringens</i> | Cambess. | 8 | 3 | 6 | common | 0.416667 | partially protected |
| <i>Eugenia aurata</i> | O.Berg | 20 | 3 | 15 | common | 0.151002 | partially protected |
| <i>Eugenia azeda</i> | Sobral | 1 | 3 | 2 | form4 | 0 | unprotected |
| <i>Eugenia azurensis</i> | O.Berg | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia bahiensis</i> | DC. | 2 | 3 | 1 | form6 | 0.25 | partially protected |
| <i>Eugenia biflora</i> | (L.) DC. | 26 | 3 | 23 | common | 0.253846 | partially protected |
| <i>Eugenia bimarginata</i> | DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia brejoensis</i> | Mazine | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia candelleana</i> | DC. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Eugenia capparidifolia</i> | DC. | 3 | 3 | 2 | form4 | 0 | unprotected |
| <i>Eugenia chlorocarpa</i> | O.Berg | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eugenia chrysophyllum</i> | Poir. | 6 | 3 | 3 | common | 0.090909 | partially protected |
| <i>Eugenia citrifolia</i> | Poir. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eugenia complicata</i> | O.Berg | 10 | 3 | 5 | common | 0.151515 | partially protected |
| <i>Eugenia crassa</i> | Sobral | 2 | 1 | 2 | form5 | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|--------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Eugenia cibrata</i> | McVaugh | 6 | 3 | 2 | common | 0.047619 | partially protected |
| <i>Eugenia dichroma</i> | O.Berg | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Eugenia dictyophleba</i> | O.Berg | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Eugenia dysenterica</i> | DC. | 21 | 3 | 13 | common | 0.139773 | partially protected |
| <i>Eugenia egensis</i> | DC. | 26 | 3 | 24 | common | 0.217391 | partially protected |
| <i>Eugenia excelsa</i> | O.Berg | 12 | 3 | 7 | common | 0.333333 | partially protected |
| <i>Eugenia ferreiraiana</i> | O.Berg | 9 | 1 | 11 | form1 | 0.333333 | partially protected |
| <i>Eugenia flavescens</i> | DC. | 22 | 3 | 20 | common | 0.226667 | partially protected |
| <i>Eugenia florida</i> | DC. | 29 | 3 | 30 | common | 0.160804 | partially protected |
| <i>Eugenia gemmiflora</i> | O.Berg | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Eugenia hartmanniae</i> | Mattos | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia hiemalis</i> | Cambess. | 10 | 3 | 3 | common | 0.235294 | partially protected |
| <i>Eugenia hirta</i> | O.Berg | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eugenia incanescens</i> | Benth. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Eugenia inundata</i> | DC. | 18 | 3 | 16 | common | 0.185185 | partially protected |
| <i>Eugenia involucrata</i> | DC. | 6 | 3 | 3 | common | 0.190476 | partially protected |
| <i>Eugenia itapemirimensis</i> | Cambess. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia klotzschiana</i> | O.Berg | 10 | 3 | 4 | common | 0.171717 | partially protected |
| <i>Eugenia lambertiana</i> | DC. | 20 | 3 | 16 | common | 0.466667 | partially protected |
| <i>Eugenia laruotteana</i> | Cambess. | 4 | 1 | 3 | form1 | 0.166667 | partially protected |
| <i>Eugenia ligustrina</i> | (Sw.) Willd. | 21 | 3 | 15 | common | 0.359551 | partially protected |
| <i>Eugenia lizada</i> | O.Berg | 11 | 3 | 6 | common | 0.089109 | partially protected |
| <i>Eugenia macrantha</i> | O.Berg | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eugenia mansoi</i> | O.Berg | 5 | 3 | 2 | common | 0.142857 | partially protected |
| <i>Eugenia matogrossensis</i> | Sobral | 3 | 3 | 3 | form4 | 0.153846 | partially protected |
| <i>Eugenia minuscula</i> | McVaugh | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Eugenia monosperma</i> | Vell. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Eugenia myrcianthes</i> | Nied. | 10 | 3 | 5 | common | 0.173913 | partially protected |
| <i>Eugenia neoformosa</i> | Sobral | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Eugenia patrisii</i> | Vahl | 19 | 3 | 20 | common | 0.333333 | partially protected |
| <i>Eugenia piloensis</i> | Cambess. | 7 | 3 | 2 | common | 0.25 | partially protected |
| <i>Eugenia pluriflora</i> | DC. | 4 | 1 | 3 | form1 | 0 | unprotected |
| <i>Eugenia pohliana</i> | DC. | 2 | 1 | 3 | form5 | 0.25 | partially protected |
| <i>Eugenia protenta</i> | McVaugh | 11 | 3 | 14 | common | 0.625 | partially protected |
| <i>Eugenia punicifolia</i> | (Kunth) DC. | 35 | 3 | 39 | common | 0.173228 | partially protected |
| <i>Eugenia pyrifera</i> | Faria & Proen?Ã,Âºa | 3 | 1 | 1 | form7 | 0 | unprotected |
| <i>Eugenia sonderiana</i> | O.Berg | 9 | 3 | 5 | common | 0.22449 | partially protected |
| <i>Eugenia sparsa</i> | S.Moore | 14 | 3 | 4 | common | 0.1 | partially protected |
| <i>Eugenia stictopetala</i> | DC. | 30 | 3 | 33 | common | 0.214286 | partially protected |
| <i>Eugenia suberosa</i> | Cambess. | 19 | 3 | 8 | common | 0.161074 | partially protected |
| <i>Eugenia sulcata</i> | Spring ex Mart. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Eugenia ternatifolia</i> | Cambess. | 2 | 3 | 2 | form4 | 0.375 | partially protected |
| <i>Eugenia uniflora</i> | L. | 17 | 3 | 12 | common | 0.234043 | partially protected |
| <i>Eugenia vernicosa</i> | O.Berg | 2 | 3 | 3 | form4 | 0.076923 | partially protected |
| <i>Eugenia vetula</i> | DC. | 10 | 3 | 4 | common | 0.222222 | partially protected |
| <i>Eugenia warmingiana</i> | Kiaersk. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Euphorbia cotinifolia</i> | L. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Euphronia acuminatissima</i> | Steyermark | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Euphronia guianensis</i> | (R.H.Schomb.) Hallier f. | 7 | 3 | 8 | common | 0.555556 | partially protected |
| <i>Euplassa bahiensis</i> | (Meisn.) I.M.Johnst. | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Euplassa inaequalis</i> | (Pohl) Engl. | 23 | 3 | 19 | common | 0.147059 | partially protected |
| <i>Euplassa semicostata</i> | Plana | 2 | 1 | 3 | form5 | 0.5 | partially protected |
| <i>Exelodendron cordatum</i> | (Hook.f.) Prance | 12 | 3 | 5 | common | 0.092308 | partially protected |
| <i>Exelodendron gardneri</i> | (Hook.f.) Prance | 10 | 3 | 6 | common | 0.196429 | partially protected |
| <i>Exostyles amazonica</i> | Yakovlev | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Faramea bracteata</i> | Benth. | 14 | 3 | 7 | common | 0.166667 | partially protected |
| <i>Faramea capillipes</i> | MÃƒÂ¼ll.Arg. | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Faramea corymbosa</i> | Aubl. | 16 | 3 | 12 | common | 0.147059 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Faramea crassifolia</i> | Benth. | 7 | 3 | 8 | common | 0.6 | partially protected |
| <i>Faramea latifolia</i> | (Cham. & Schltdl.) DC. | 10 | 3 | 2 | common | 0.145455 | partially protected |
| <i>Faramea multiflora</i> | A.Rich. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Faramea nitida</i> | Benth. | 11 | 3 | 7 | common | 0.25 | partially protected |
| <i>Faramea paniculata</i> | (Aubl.) Benth. | 4 | 1 | 3 | form1 | 0 | unprotected |
| <i>Faramea stipulacea</i> | (Cham. & Schltdl.) DC. | 3 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Ferdinandusa chlorantha</i> | (Wedd.) Standl. | 14 | 3 | 12 | common | 0.416667 | partially protected |
| <i>Ferdinandusa elliptica</i> | (Pohl) Pohl | 15 | 3 | 8 | common | 0.147727 | partially protected |
| <i>Ferdinandusa nitida</i> | Ducke | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Ferdinandusa speciosa</i> | (Pohl) Pohl | 7 | 3 | 5 | common | 0.111111 | partially protected |
| <i>Ficus adhatodifolia</i> | Schott | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ficus americana</i> | Aubl. | 27 | 3 | 33 | common | 0.297101 | partially protected |
| <i>Ficus bahiensis</i> | C.C.Berg & Caraúta | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Ficus catappifolia</i> | Kunth & C.D.Bouchéf© | 4 | 3 | 4 | common | 0.4 | partially protected |
| <i>Ficus citrifolia</i> | Mill. | 28 | 3 | 25 | common | 0.200855 | partially protected |
| <i>Ficus enormis</i> | (Miq.) Miq. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ficus gomelleira</i> | Kunth & C.D.Bouchéf© | 5 | 3 | 4 | common | 0 | unprotected |
| <i>Ficus insipida</i> | Willd. | 21 | 3 | 20 | common | 0.216216 | partially protected |
| <i>Ficus laureola</i> | Warb. ex C.C.Berg & Caraúta | 2 | 1 | 2 | form5 | 0.4 | partially protected |
| <i>Ficus mathewssii</i> | (Miq.) Miq. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ficus maxima</i> | Mill. | 21 | 3 | 20 | common | 0.195652 | partially protected |
| <i>Ficus nymphaeifolia</i> | Mill. | 20 | 3 | 17 | common | 0.392857 | partially protected |
| <i>Ficus obtusifolia</i> | Kunth | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ficus pertusa</i> | L.f. | 25 | 3 | 25 | common | 0.237624 | partially protected |
| <i>Ficus trigona</i> | L.f. | 24 | 3 | 23 | common | 0.140845 | partially protected |
| <i>Fridericia bahiensis</i> | (Schauer) L.G.Lohmann | 3 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Fridericia dispar</i> | (Bureau ex K.Schum.) L.G.Lohmann | 4 | 3 | 4 | common | 0.071429 | partially protected |
| <i>Fridericia platyphylla</i> | (Cham.) L.G.Lohmann | 2 | 3 | 1 | form6 | 0.181818 | partially protected |
| <i>Galipea ciliata</i> | Taub. | 7 | 3 | 4 | common | 0.157895 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-------------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Galipea trifoliata</i> | Aubl. | 16 | 3 | 17 | common | 0.230769 | partially protected |
| <i>Garcinia brasiliensis</i> | Mart. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Gaultheria eriophylla</i> | (Pers.) Sleumer ex Burtt | 1 | 1 | 3 | form5 | 0.5 | partially protected |
| <i>Gaylussacia brasiliensis</i> | (Spreng.) Meisn. | 16 | 3 | 9 | common | 0.237668 | partially protected |
| <i>Gaylussacia montana</i> | (Pohl) Sleumer | 5 | 3 | 5 | common | 0.297297 | partially protected |
| <i>Gaylussacia pseudogaultheria</i> | Cham. & Schltdl. | 7 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Gaylussacia salicifolia</i> | Cham. & Schltdl. | 4 | 3 | 4 | common | 0.214286 | partially protected |
| <i>Genipa americana</i> | L. | 30 | 3 | 30 | common | 0.253807 | partially protected |
| <i>Geonoma deversa</i> | (Poit.) Kunth | 21 | 3 | 21 | common | 0.217391 | partially protected |
| <i>Geonoma stricta</i> | (Poit.) Kunth | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Gliricidia sepium</i> | (Jacq.) Walp. | 2 | 3 | 2 | form4 | 0.25 | partially protected |
| <i>Godmania aesculifolia</i> | (Kunth) Standl. | 13 | 3 | 8 | common | 0.217391 | partially protected |
| <i>Godmania dardanoi</i> | (J.C.Gomes) A.H.Gentry | 3 | 3 | 2 | form4 | 0 | unprotected |
| <i>Gongylolepis benthamiana</i> | R.H.Schomb. | 5 | 3 | 3 | common | 0.6 | partially protected |
| <i>Gongylolepis martiana</i> | (Baker) Steyermark & Cuatrec. | 10 | 3 | 10 | common | 0.285714 | partially protected |
| <i>Gorceixia decurrens</i> | Baker | 3 | 1 | 3 | form5 | 0.25 | partially protected |
| <i>Gordonia fruticosa</i> | (Schrad.) H.Keng | 6 | 3 | 6 | common | 0.263158 | partially protected |
| <i>Gouphia glabra</i> | Aubl. | 21 | 3 | 21 | common | 0.176471 | partially protected |
| <i>Graffenrieda weddellii</i> | Naudin | 12 | 3 | 9 | common | 0.35 | partially protected |
| <i>Guaiacum officinale</i> | L. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Guapira areolata</i> | (Heimerl) Lundell | 13 | 3 | 7 | common | 0.165468 | partially protected |
| <i>Guapira campestris</i> | (Netto) Lundell | 12 | 3 | 5 | common | 0.1 | partially protected |
| <i>Guapira graciliflora</i> | (Mart. ex J.A.Schmidt) Lundell | 21 | 3 | 11 | common | 0.151815 | partially protected |
| <i>Guapira microphylla</i> | (Heimerl) Lundell | 4 | 3 | 1 | form2 | 0 | unprotected |
| <i>Guapira noxia</i> | (Netto) Lundell | 18 | 3 | 10 | common | 0.120797 | partially protected |
| <i>Guapira obtusata</i> | (Jacq.) Little | 13 | 3 | 9 | common | 0.181818 | partially protected |
| <i>Guapira opposita</i> | (Vell.) Reitz | 29 | 3 | 28 | common | 0.22 | partially protected |
| <i>Guapira pacurero</i> | (Kunth) Little | 2 | 3 | 1 | form6 | 0.2 | partially protected |
| <i>Guapira pernambucensis</i> | (Casar.) Lundell | 8 | 3 | 4 | common | 0.352941 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|-----------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Guapira tomentosa</i> | (Casar.) Lundell | 3 | 3 | 3 | form4 | 0.608696 | partially protected |
| <i>Guarea guidonia</i> | (L.) Sleumer | 30 | 3 | 29 | common | 0.265823 | partially protected |
| <i>Guarea kunthiana</i> | A.Juss. | 27 | 3 | 28 | common | 0.244898 | partially protected |
| <i>Guatteria australis</i> | A.St.-Hil. | 6 | 3 | 3 | common | 0.25 | partially protected |
| <i>Guatteria discolor</i> | R.E.Fr. | 16 | 3 | 15 | common | 0.2 | partially protected |
| <i>Guatteria dura</i> | R.E.Fr. | 4 | 1 | 4 | form1 | 0 | unprotected |
| <i>Guatteria inundata</i> | Mart. | 11 | 3 | 8 | common | 0.125 | partially protected |
| <i>Guatteria maypurensis</i> | Kunth | 24 | 3 | 17 | common | 0.121951 | partially protected |
| <i>Guatteria rigida</i> | R.E.Fr. | 16 | 3 | 9 | common | 0.340909 | partially protected |
| <i>Guatteria rupestris</i> | Mello-Silva & Pirani | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Guatteria schomburgkiana</i> | Mart. | 24 | 3 | 28 | common | 0.166667 | partially protected |
| <i>Guatteria sellowiana</i> | Schltdl. | 11 | 3 | 9 | common | 0.272059 | partially protected |
| <i>Guazuma ulmifolia</i> | Lam. | 29 | 3 | 26 | common | 0.131222 | partially protected |
| <i>Guettarda macrantha</i> | Benth. | 19 | 3 | 10 | common | 0.1 | partially protected |
| <i>Guettarda malacophylla</i> | Standl. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Guettarda platypoda</i> | DC. | 12 | 3 | 11 | common | 0.3125 | partially protected |
| <i>Guettarda pohliana</i> | MÃƒÂ¼ll.Arg. | 8 | 3 | 4 | common | 0.307692 | partially protected |
| <i>Guettarda viburnoides</i> | Cham. & Schltdl. | 27 | 3 | 24 | common | 0.142574 | partially protected |
| <i>Guibourtia hymenaeifolia</i> | (Moric.) J.Leonard | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Gymnanthes glabrata</i> | (Mart.) Govaerts | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Hancornia speciosa</i> | Gomes | 26 | 3 | 24 | common | 0.156275 | partially protected |
| <i>Handroanthus heptaphyllus</i> | (Vell.) Mattos | 3 | 1 | 1 | form7 | 0 | unprotected |
| <i>Handroanthus impetiginosus</i> | (Mart. ex DC.) Mattos | 23 | 3 | 14 | common | 0.137615 | partially protected |
| <i>Handroanthus ochraceus</i> | (Cham.) Mattos | 33 | 3 | 26 | common | 0.149488 | partially protected |
| <i>Handroanthus pulcherrimus</i> | (Sandwith) S.O.Grose | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Handroanthus serratifolius</i> | (Vahl) S.O.Grose | 28 | 3 | 30 | common | 0.215223 | partially protected |
| <i>Haydenoxylon urbanianum</i> | (Loes.) M.P. Simmons | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Hecatostemon completus</i> | (Jacq.) Sleumer | 8 | 3 | 5 | common | 0.122449 | partially protected |
| <i>Hedyosmum brasiliense</i> | Mart. | 5 | 3 | 5 | common | 0.181818 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Heisteria citrifolia</i> | Engl. | 11 | 3 | 7 | common | 0.196262 | partially protected |
| <i>Heisteria ovata</i> | Benth. | 24 | 3 | 25 | common | 0.202899 | partially protected |
| <i>Helicteres aspera</i> | A. St.-Hil. & Naudin | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Helicteres baruensis</i> | Jacq. | 12 | 3 | 8 | common | 0.210526 | partially protected |
| <i>Helicteres brevispira</i> | A.Juss. | 21 | 3 | 16 | common | 0.139344 | partially protected |
| <i>Helicteres corylifolia</i> | Nees & Mart. | 19 | 3 | 11 | common | 0.145161 | partially protected |
| <i>Helicteres eichleri</i> | K. Schum. | 5 | 3 | 2 | common | 0.375 | partially protected |
| <i>Helicteres guazumifolia</i> | Kunth | 17 | 3 | 12 | common | 0.190476 | partially protected |
| <i>Helicteres heptandra</i> | L.B.Sm. | 9 | 3 | 8 | common | 0.195122 | partially protected |
| <i>Helicteres lhotzkyana</i> | K.Schum. | 11 | 3 | 5 | common | 0.157895 | partially protected |
| <i>Helicteres macropetala</i> | A. St.-Hil. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Helicteres muscosa</i> | Mart. | 7 | 3 | 6 | common | 0.315789 | partially protected |
| <i>Helicteres ovata</i> | Lam. | 5 | 3 | 4 | common | 0.111111 | partially protected |
| <i>Helicteres pentandra</i> | L. | 18 | 3 | 14 | common | 0.333333 | partially protected |
| <i>Helietta apiculata</i> | Benth. | 3 | 3 | 1 | form6 | 0.1 | partially protected |
| <i>Helietta glaziovii</i> | (Engl.) Pirani (Vell.) Penneys, F.A. Michelangeli, Judd & | 3 | 1 | 1 | form7 | 0.2 | partially protected |
| <i>Henriettea glabra</i> | Almeda | 3 | 1 | 1 | form7 | 0 | unprotected |
| <i>Henriettea granulata</i> | O. Berg & Triana | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Henriettea maroniensis</i> | Sagot (Cogn.) Penneys, F.A. Michelangeli, Judd & | 8 | 3 | 7 | common | 0.636364 | partially protected |
| <i>Henriettea ovata</i> | Almeda | 8 | 3 | 7 | common | 0 | unprotected |
| <i>Henriettea saldanhaei</i> | Cogn. | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Henriettea succosa</i> | (Aubl.) DC. | 6 | 3 | 8 | common | 0 | unprotected |
| <i>Heterocoma ekmaniana</i> | (Philipson) B. Loeuille, J.N. Nakaj. & Semir | 4 | 1 | 4 | form1 | 0 | unprotected |
| <i>Heterocoma lanuginosa</i> | (Glaz. ex Oliv.) B. Loeuille, J.N. Nakaj. & Semir | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Heteropterys sincorensis</i> | W.R.Anderson | 5 | 3 | 3 | common | 0.333333 | partially protected |
| <i>Heterostemon mimosoides</i> | Desf. | 12 | 3 | 14 | common | 0 | unprotected |
| <i>Hieronyma oblonga</i> | (Tul.) MÃ¶ller.Arg. | 2 | 1 | 1 | form7 | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Himatanthus articulatus</i> | (Vahl) Woodson | 24 | 3 | 26 | common | 0.273504 | partially protected |
| <i>Himatanthus bracteatus</i> | (A.DC.) Woodson | 16 | 3 | 16 | common | 0.285714 | partially protected |
| <i>Himatanthus drasticus</i> | (Mart.) Plumel | 25 | 3 | 20 | common | 0.177898 | partially protected |
| <i>Himatanthus obovatus</i> | (MÃƒÂ½ll.Arg.) Woodson | 23 | 3 | 20 | common | 0.122661 | partially protected |
| <i>Himatanthus phagedaenicus</i> | (Mart.) Woodson | 20 | 3 | 17 | common | 0.177778 | partially protected |
| <i>Himatanthus sucuuba</i> | (Spruce ex MÃƒÂ½ll.Arg.) Woodson | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Himatanthus tarapotensis</i> | (K.Schum. ex Markgr.) Plumel | 13 | 3 | 9 | common | 0.25 | partially protected |
| <i>Hirtella bicornis</i> | Mart. & Zucc. | 19 | 3 | 18 | common | 0.266667 | partially protected |
| <i>Hirtella burchellii</i> | Britton | 21 | 3 | 18 | common | 0.315789 | partially protected |
| <i>Hirtella ciliata</i> | Mart. & Zucc. | 24 | 3 | 17 | common | 0.154734 | partially protected |
| <i>Hirtella glandulosa</i> | Spreng. | 27 | 3 | 21 | common | 0.244147 | partially protected |
| <i>Hirtella gracilipes</i> | (Hook.f.) Prance | 24 | 3 | 16 | common | 0.155361 | partially protected |
| <i>Hirtella paniculata</i> | Sw. | 14 | 3 | 14 | common | 0.197368 | partially protected |
| <i>Hirtella punctillata</i> | Ducke | 11 | 3 | 11 | common | 0 | unprotected |
| <i>Hirtella racemosa</i> | Lam. | 28 | 3 | 37 | common | 0.29064 | partially protected |
| <i>Hirtella scabra</i> | Benth. | 8 | 3 | 9 | common | 0.642857 | partially protected |
| <i>Hirtella sprucei</i> | Benth. ex Hook.f. | 5 | 1 | 2 | form1 | 0 | unprotected |
| <i>Hirtella triandra</i> | Sw. | 18 | 3 | 18 | common | 0.478261 | partially protected |
| <i>Hololepis pedunculata</i> | (DC. ex Pers.) DC. | 5 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Hornschlorchia bryotrophe</i> | Nees | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Hortia oreadica</i> | Groppi, Kallunki & Pirani | 5 | 3 | 4 | common | 0 | unprotected |
| <i>Humiria balsamifera</i> | Aubl. | 28 | 3 | 36 | common | 0.293532 | partially protected |
| <i>Humiria crassifolia</i> | Mart. ex Urb. | 11 | 3 | 13 | common | 0.833333 | partially protected |
| <i>Humiriastrum mussungense</i> | Cuatrec. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Hura crepitans</i> | L. | 20 | 3 | 19 | common | 0.264706 | partially protected |
| <i>Hyeronima alchorneoides</i> | AllemÃƒÆ'Ã,Ã£o | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Hymenaea courbaril</i> | L. | 27 | 3 | 31 | common | 0.217391 | partially protected |
| <i>Hymenaea eriogyne</i> | Benth. | 10 | 3 | 4 | common | 0.217391 | partially protected |
| <i>Hymenaea longifolia</i> | (Benth.) I.M. Souza, Funch & L.P. Queiroz | 5 | 1 | 2 | form1 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|-------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Hymenaea maranhensis</i> | Lee & Lang. | 6 | 3 | 3 | common | 0 | unprotected |
| <i>Hymenaea martiana</i> | Hayne | 18 | 3 | 11 | common | 0.2 | partially protected |
| <i>Hymenaea rubriflora</i> | Ducke | 5 | 3 | 4 | common | 0 | unprotected |
| <i>Hymenaea stigonocarpa</i> | Hayne | 22 | 3 | 14 | common | 0.118674 | partially protected |
| <i>Hymenaea velutina</i> | Ducke | 6 | 3 | 3 | common | 0.4 | partially protected |
| <i>Hymenolobium heringeranum</i> | Rizzini | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Hymenolobium petraeum</i> | Ducke | 9 | 3 | 10 | common | 0.2 | partially protected |
| <i>Hypenia salzmannii</i> | (Benth.) Harley | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Hyptidendron arboreum</i> | (Benth.) Harley | 3 | 3 | 3 | form4 | 0.714286 | partially protected |
| <i>Hyptidendron canum</i> | (Pohl ex Benth.) Harley | 13 | 3 | 8 | common | 0.100649 | partially protected |
| <i>Hyptidendron claussenii</i> | (Benth.) Harley | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Hyptidendron conspersum</i> | (Benth.) Harley | 5 | 3 | 1 | form2 | 0.125 | partially protected |
| <i>Hyptidendron glutinosum</i> | (Benth.) Harley | 9 | 3 | 3 | common | 0.2 | partially protected |
| <i>Hyptis pachyphylla</i> | Epling | 4 | 1 | 1 | form3 | 0.6 | partially protected |
| <i>Ilex affinis</i> | Gardner | 18 | 3 | 13 | common | 0.22973 | partially protected |
| <i>Ilex asperula</i> | Mart. | 4 | 3 | 4 | common | 0.25 | partially protected |
| <i>Ilex brasiliensis</i> | (Spreng.) Loes. | 8 | 3 | 4 | common | 0.25641 | partially protected |
| <i>Ilex cerasifolia</i> | Reissek | 9 | 3 | 4 | common | 0.185185 | partially protected |
| <i>Ilex conocarpa</i> | Reissek | 7 | 3 | 4 | common | 0.309091 | partially protected |
| <i>Ilex dumosa</i> | Reissek | 9 | 3 | 4 | common | 0.326531 | partially protected |
| <i>Ilex floribunda</i> | Reissek ex Maxim. | 6 | 3 | 3 | common | 0.285714 | partially protected |
| <i>Ilex inundata</i> | Poepp. ex Reissek | 16 | 3 | 18 | common | 0.25 | partially protected |
| <i>Ilex jenmanii</i> | Loes. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Ilex lundii</i> | Warm. | 8 | 3 | 5 | common | 0.189189 | partially protected |
| <i>Ilex martiniana</i> | D.Don | 3 | 3 | 2 | form4 | 0.25 | partially protected |
| <i>Ilex microdonta</i> | Reissek | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ilex oligoneura</i> | Loes. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ilex paujiensis</i> | Steyermark | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ilex polita</i> | Steyermark | 1 | 1 | 1 | form7 | 1 | protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|---------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Ilex psammophila</i> | Reissek | 6 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Ilex pseudobuxus</i> | Reissek | 4 | 3 | 3 | common | 0.347826 | partially protected |
| <i>Ilex pseudovaccinium</i> | Reissek ex Maxim. | 4 | 3 | 3 | common | 0.444444 | partially protected |
| <i>Ilex retusa</i> | Klotzsch ex Reissek | 10 | 3 | 11 | common | 0.555556 | partially protected |
| <i>Ilex theezans</i> | Mart. | 7 | 3 | 6 | common | 0.285714 | partially protected |
| <i>Ilex vismiifolia</i> | Reissek | 15 | 3 | 18 | common | 0.5 | partially protected |
| <i>Indigofera blanchetiana</i> | Benth. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Inga alba</i> | (Sw.) Willd. | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Inga bourgonii</i> | (Aubl.) DC. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Inga cabelo</i> | T.D.Penn. | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Inga capitata</i> | Desv. | 24 | 3 | 29 | common | 0.38806 | partially protected |
| <i>Inga disticha</i> | Benth. | 22 | 3 | 20 | common | 0.071429 | partially protected |
| <i>Inga edulis</i> | Mart. | 24 | 3 | 24 | common | 0.384615 | partially protected |
| <i>Inga exfoliata</i> | T.D.Penn. & F.C.P. Garcia | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Inga flagelliformis</i> | (Vell.) Mart. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Inga heterophylla</i> | Willd. | 26 | 3 | 22 | common | 0.310345 | partially protected |
| <i>Inga hispida</i> | Benth. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Inga ingoides</i> | (Rich.) Willd. | 8 | 3 | 8 | common | 0.272727 | partially protected |
| <i>Inga lanceifolia</i> | Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Inga laurina</i> | (Sw.) Willd. | 12 | 3 | 7 | common | 0.210526 | partially protected |
| <i>Inga leptantha</i> | Benth. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Inga nobilis</i> | Willd. | 26 | 3 | 28 | common | 0.178571 | partially protected |
| <i>Inga pilosula</i> | (Rich.) J.F.Macbr. | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Inga ruiziana</i> | G.Don | 8 | 3 | 5 | common | 0.4 | partially protected |
| <i>Inga splendens</i> | Willd. | 20 | 3 | 20 | common | 0.380952 | partially protected |
| <i>Inga stenoptera</i> | Benth. | 16 | 3 | 16 | common | 0.173913 | partially protected |
| <i>Inga striolata</i> | T.D.Penn. | 6 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Inga subnuda</i> | Salzm. ex Benth. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Inga thibaudiana</i> | DC. | 25 | 3 | 31 | common | 0.337838 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|-------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Inga vera</i> | Willd. | 23 | 3 | 19 | common | 0.179856 | partially protected |
| <i>Insertia parviflora</i> | Vahl | 11 | 3 | 13 | common | 0.214286 | partially protected |
| <i>Ixora brevifolia</i> | Benth. | 15 | 3 | 5 | common | 0.153846 | partially protected |
| <i>Jacaranda brasiliiana</i> | (Lam.) Pers. | 21 | 3 | 16 | common | 0.209434 | partially protected |
| <i>Jacaranda carajasensis</i> | A.H.Gentry | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Jacaranda caroba</i> | (Vell.) DC. | 13 | 3 | 8 | common | 0.166667 | partially protected |
| <i>Jacaranda cuspidifolia</i> | Mart. | 16 | 3 | 10 | common | 0.126126 | partially protected |
| <i>Jacaranda irwinii</i> | A.H.Gentry | 4 | 3 | 3 | common | 0.347826 | partially protected |
| <i>Jacaranda jasminoides</i> | (Thunb.) Sandwith | 9 | 3 | 5 | common | 0.32 | partially protected |
| <i>Jacaranda micrantha</i> | Cham. | 2 | 3 | 2 | form4 | 0.333333 | partially protected |
| <i>Jacaranda obovata</i> | Cham. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Jacaranda praetermissa</i> | Sandwith | 10 | 3 | 4 | common | 0.111111 | partially protected |
| <i>Jacaranda puberula</i> | Cham. | 7 | 3 | 2 | common | 0.235294 | partially protected |
| <i>Jacaratia corumbensis</i> | Kuntze | 9 | 3 | 6 | common | 0.090909 | partially protected |
| <i>Jacaratia spinosa</i> | (Aubl.) A.DC. | 21 | 3 | 13 | common | 0.291667 | partially protected |
| <i>Jatropha curcas</i> | L. | 15 | 3 | 14 | common | 0.428571 | partially protected |
| <i>Jatropha mollissima</i> | (Pohl) Baill. | 13 | 3 | 7 | common | 0.166667 | partially protected |
| <i>Kielmeyera albopunctata</i> | Saddi | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Kielmeyera coriacea</i> | Mart. & Zucc. | 22 | 3 | 17 | common | 0.134831 | partially protected |
| <i>Kielmeyera cuspidata</i> | Saddi | 3 | 3 | 3 | form4 | 0.375 | partially protected |
| <i>Kielmeyera grandiflora</i> | (Wawra) Saddi | 15 | 3 | 8 | common | 0.155556 | partially protected |
| <i>Kielmeyera lathrophyton</i> | Saddi | 16 | 3 | 8 | common | 0.181159 | partially protected |
| <i>Kielmeyera membranacea</i> | Casar. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Kielmeyera neglecta</i> | Saddi | 4 | 3 | 5 | common | 0 | unprotected |
| <i>Kielmeyera occhioniana</i> | Saddi | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Kielmeyera paniculata</i> | Rusby | 5 | 3 | 4 | common | 0.142857 | partially protected |
| <i>Kielmeyera petiolaris</i> | Mart. & Zucc. | 11 | 3 | 6 | common | 0.12963 | partially protected |
| <i>Kielmeyera pulcherrima</i> | L.B. Sm. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Kielmeyera reticulata</i> | Saddi | 2 | 1 | 3 | form5 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Kielmeyera rosea</i> | Mart. & Zucc. | 13 | 3 | 6 | common | 0.147059 | partially protected |
| <i>Kielmeyera rubriflora</i> | Cambess. | 17 | 3 | 12 | common | 0.127143 | partially protected |
| <i>Kielmeyera speciosa</i> | A. St.-Hil. | 16 | 3 | 6 | common | 0.115854 | partially protected |
| <i>Kielmeyera tomentosa</i> | Cambess. | 5 | 3 | 4 | common | 0.210526 | partially protected |
| <i>Kielmeyera variabilis</i> | Mart. & Zucc. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Krameria tomentosa</i> | A. St.-Hil. | 4 | 3 | 4 | common | 0.235294 | partially protected |
| <i>Lacistema aggregatum</i> | (P.J.Bergius) Rusby | 31 | 3 | 24 | common | 0.270531 | partially protected |
| <i>Lacistema grandifolium</i> | Schnizl. | 13 | 3 | 9 | common | 0.428571 | partially protected |
| <i>Lacistema hasslerianum</i> | Chodat | 18 | 3 | 12 | common | 0.170404 | partially protected |
| <i>Lacistema polystachyum</i> | Schnizl. | 18 | 3 | 10 | common | 0.2 | partially protected |
| <i>Ladenbergia cujabensis</i> | Klotzsch | 12 | 3 | 4 | common | 0.209677 | partially protected |
| <i>Ladenbergia hexandra</i> | (Pohl) Klotzsch | 3 | 1 | 3 | form5 | 0.2 | partially protected |
| <i>Laetia americana</i> | L. | 12 | 3 | 8 | common | 0.555556 | partially protected |
| <i>Lafoensia glyptocarpa</i> | Koehne | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Lafoensia pacari</i> | A. St.-Hil. | 28 | 3 | 22 | common | 0.139623 | partially protected |
| <i>Lafoensia vandelliana</i> | Cham. & Schldl. | 14 | 3 | 12 | common | 0.208333 | partially protected |
| <i>Lamanonia cuneata</i> | (Cambess.) Kuntze | 3 | 1 | 2 | form5 | 0.4 | partially protected |
| <i>Lamanonia ternata</i> | Vell. | 10 | 3 | 5 | common | 0.265306 | partially protected |
| <i>Lantana camara</i> | L. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Lantana canescens</i> | Kunth | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Lantana cujabensis</i> | Schauer | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Lavoisiera nervulosa</i> | Naudin | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Leandra aurea</i> | (Cham.) Cogn. | 14 | 3 | 8 | common | 0.285714 | partially protected |
| <i>Leandra melastomoides</i> | Raddi | 4 | 3 | 2 | common | 0.384615 | partially protected |
| <i>Leandra rufescens</i> | (DC.) Cogn. | 8 | 3 | 7 | common | 0.090909 | partially protected |
| <i>Lecythis pisonis</i> | Cambess. | 16 | 3 | 18 | common | 0.333333 | partially protected |
| <i>Leocereus bahiensis</i> | Britton & Rose | 5 | 3 | 3 | common | 0.4 | partially protected |
| <i>Lepidaploa remotiflora</i> | (Rich.) H.Rob. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Leptolobium brachystachyum</i> | (Benth.) Sch.Rodr. & A.M.G.Azevedo | 4 | 3 | 3 | common | 0.47619 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Leptolobium glaziovianum</i> | (Harms) Sch.Rodr. & A.M.G.Azevedo (G.A. Aymard & V. GonzÃ¡lez) Sch. Rodr. & A.M.G. | 5 | 3 | 1 | form2 | 0.4 | partially protected |
| <i>Leptolobium stirtonii</i> | | 3 | 3 | 3 | form4 | 0.363636 | partially protected |
| <i>Leucochloron incuriale</i> | (Vell.)Barneby & J.W.Grimes | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Leucochloron minarum</i> | (Harms)Barneby & J.W.Grimes | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Libidibia ferrea</i> | (Mart. ex Tul.) L.P.Queiroz | 6 | 3 | 4 | common | 0.16 | partially protected |
| <i>Licania apetala</i> | (E.Mey.) Fritsch | 28 | 3 | 28 | common | 0.267717 | partially protected |
| <i>Licania blackii</i> | Prance | 20 | 3 | 16 | common | 0.333333 | partially protected |
| <i>Licania dealbata</i> | Hook.f. | 12 | 3 | 4 | common | 0.246914 | partially protected |
| <i>Licania egleri</i> | Prance | 18 | 3 | 17 | common | 0.4 | partially protected |
| <i>Licania gardneri</i> | (Hook.f.) Fritsch. | 15 | 3 | 8 | common | 0.207547 | partially protected |
| <i>Licania humilis</i> | Cham. & Schltdl. | 13 | 3 | 10 | common | 0.117424 | partially protected |
| <i>Licania hypoleuca</i> | Benth. | 26 | 3 | 26 | common | 0.172414 | partially protected |
| <i>Licania incana</i> | Aubl. | 12 | 3 | 16 | common | 0.461538 | partially protected |
| <i>Licania kunthiana</i> | Hook.f. | 24 | 3 | 25 | common | 0.304348 | partially protected |
| <i>Licania lasseri</i> | Maguire | 3 | 3 | 2 | form4 | 0.666667 | partially protected |
| <i>Licania licaniiflora</i> | (Sagot) S.F.Blake | 13 | 3 | 14 | common | 0.545455 | partially protected |
| <i>Licania minutiflora</i> | (Sagot) Fritsch | 11 | 3 | 12 | common | 0.222222 | partially protected |
| <i>Licania nitida</i> | Hook.f. | 4 | 3 | 3 | common | 0.307692 | partially protected |
| <i>Licania octandra</i> | (Hoffmanns. ex Schult.) Kuntze | 7 | 3 | 3 | common | 0 | unprotected |
| <i>Licania parvifolia</i> | Huber | 20 | 3 | 17 | common | 0.173913 | partially protected |
| <i>Licania rigida</i> | Benth. | 6 | 3 | 5 | common | 0.5 | partially protected |
| <i>Licania sclerophylla</i> | (Mart. ex Hook.f.) Fritsch | 7 | 3 | 3 | common | 0.4 | partially protected |
| <i>Licania steyermarkii</i> | Maguire | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Lindackeria latifolia</i> | Benth. | 13 | 3 | 14 | common | 0.5 | partially protected |
| <i>Lippia origanoides</i> | Kunth | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Lithraea brasiliensis</i> | Marchand | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Lonchocarpus punctatus</i> | Kunth | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Lonchocarpus sericeus</i> | (Poir.)DC. | 3 | 3 | 2 | form4 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Luehea candicans</i> | Mart. | 13 | 3 | 5 | common | 0.142857 | partially protected |
| <i>Luehea cymulosa</i> | Spruce ex Benth. | 4 | 3 | 1 | form2 | 0.1 | partially protected |
| <i>Luehea divaricata</i> | Mart. | 13 | 3 | 4 | common | 0.183673 | partially protected |
| <i>Luehea grandiflora</i> | Mart. | 21 | 3 | 16 | common | 0.084034 | partially protected |
| <i>Luehea paniculata</i> | Mart. | 26 | 3 | 20 | common | 0.103343 | partially protected |
| <i>Luehea speciosa</i> | Willd. | 5 | 3 | 3 | common | 0.1875 | partially protected |
| <i>Lueheopsis hoehnei</i> | Burret | 6 | 3 | 4 | common | 0 | unprotected |
| <i>Luetzelburgia auriculata</i> | (Allemao)Ducke | 11 | 3 | 7 | common | 0.083333 | partially protected |
| <i>Luetzelburgia praecox</i> | (Harms)Harms | 11 | 3 | 3 | common | 0.181818 | partially protected |
| <i>Luetzelburgia sotoi</i> | D. Cardoso, L.P. Queiroz & H.C. Lima | 1 | 1 | 3 | form5 | 0 | unprotected |
| <i>Luxemburgia corymbosa</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Luxemburgia mogolensis</i> | Feres | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Lychnophora crispa</i> | Mattf. | 5 | 3 | 5 | common | 0.571429 | partially protected |
| <i>Lychnophora diamantinana</i> | Coile & S.B.Jones | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Lychnophora ericoides</i> | Mart. | 10 | 3 | 7 | common | 0.217105 | partially protected |
| <i>Lychnophora granmogolensis</i> | (Duarte) D.J.N.Hind | 6 | 3 | 5 | common | 0.333333 | partially protected |
| <i>Lychnophora humillima</i> | Sch.Bip. | 1 | 1 | 3 | form5 | 0.666667 | partially protected |
| <i>Lychnophora markgravii</i> | G.M.Barroso | 3 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Lychnophora pohlii</i> | Sch.Bip. | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Lychnophora reticulata</i> | Gardner | 6 | 3 | 3 | common | 0.340426 | partially protected |
| <i>Lychnophora rosmarinifolia</i> | Mart. | 6 | 3 | 5 | common | 0.214286 | partially protected |
| <i>Lychnophora rupestris</i> | Semir | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Lychnophora salicifolia</i> | Mart. | 10 | 3 | 6 | common | 0.176471 | partially protected |
| <i>Lychnophora santosii</i> | H.Rob. | 2 | 1 | 3 | form5 | 0.25 | partially protected |
| <i>Lychnophora sellowii</i> | Sch.Bip. | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Lychnophora souzae</i> | H.Rob. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Lychnophora staavioides</i> | Mart. | 2 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Lychnophora tomentosa</i> | (Mart. ex DC.) Sch.Bip. | 2 | 3 | 3 | form4 | 0.25 | partially protected |
| <i>Lychnophora villosissima</i> | Mart. | 5 | 3 | 3 | common | 0.230769 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Lychnophoriopsis candelabrum</i> | (Sch.Bip.) H.Rob. | 2 | 1 | 3 | form5 | 0.2 | partially protected |
| <i>Lychnophoriopsis damazioi</i> | (Beauverd) H.Rob. | 2 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Lychnophoriopsis hatschbachii</i> | H.Rob. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Mabea angustifolia</i> | Spruce ex Benth. | 21 | 3 | 22 | common | 0.361702 | partially protected |
| <i>Mabea anomala</i> | MÃƒÂ½ll.Arg. | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Mabea fistulifera</i> | Mart. | 22 | 3 | 16 | common | 0.181185 | partially protected |
| <i>Mabea montana</i> | MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mabea nitida</i> | Spruce ex Benth. | 24 | 3 | 25 | common | 0.066667 | partially protected |
| <i>Mabea paniculata</i> | Spruce ex Benth. | 20 | 3 | 13 | common | 0.311475 | partially protected |
| <i>Mabea pohliana</i> | (Benth.) MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Mabea subsessilis</i> | Pax & K.Hoffm. | 21 | 3 | 23 | common | 0.285714 | partially protected |
| <i>Mabea taquari</i> | Aubl. | 14 | 3 | 15 | common | 0.074074 | partially protected |
| <i>Macairea pachyphylla</i> | Benth. | 20 | 3 | 16 | common | 0.541667 | partially protected |
| <i>Macairea radula</i> | (Bonpl.) DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Macairea thyrsiflora</i> | DC. | 24 | 3 | 24 | common | 0.394737 | partially protected |
| <i>Machaerium acutifolium</i> | Vogel | 27 | 3 | 22 | common | 0.137239 | partially protected |
| <i>Machaerium aristulatum</i> | (Benth.) Ducke | 6 | 3 | 2 | common | 0.117647 | partially protected |
| <i>Machaerium brasiliense</i> | Vogel | 15 | 3 | 5 | common | 0.08642 | partially protected |
| <i>Machaerium eriocarpum</i> | Benth. | 5 | 3 | 3 | common | 0.181818 | partially protected |
| <i>Machaerium ferox</i> | (Benth.) Ducke | 3 | 3 | 4 | form4 | 0.166667 | partially protected |
| <i>Machaerium inundatum</i> | (Benth.) Ducke | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Machaerium isadelphum</i> | (E.Mey.) Standl. | 13 | 3 | 9 | common | 0.121212 | partially protected |
| <i>Machaerium lunatum</i> | (L.f.) Ducke | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Machaerium mucronulatum</i> | Benth. | 3 | 3 | 1 | form6 | 0.4 | partially protected |
| <i>Machaerium opacum</i> | Vogel | 16 | 3 | 11 | common | 0.127371 | partially protected |
| <i>Machaerium punctatum</i> | Pers. | 2 | 1 | 1 | form7 | 0.25 | partially protected |
| <i>Machaerium quinata</i> | (Aubl.) Sandwith | 3 | 1 | 2 | form5 | 0.75 | partially protected |
| <i>Machaerium villosum</i> | Vogel | 14 | 3 | 9 | common | 0.150685 | partially protected |
| <i>Machaonia brasiliensis</i> | (Hoffmannss. ex Humb.) Cham. & Schldl. | 11 | 3 | 10 | common | 0.3125 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Maclura tinctoria</i> | (L.) D.Don ex Steud. | 32 | 3 | 29 | common | 0.149837 | partially protected |
| <i>Macrolobium acaciifolium</i> | (Benth.)Benth. | 2 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Macrolobium bifolium</i> | (Aubl.)Pers. | 4 | 3 | 1 | form2 | 0.1 | partially protected |
| <i>Macrolobium campestre</i> | Huber | 19 | 3 | 20 | common | 0.333333 | partially protected |
| <i>Macrolobium latifolium</i> | Vogel | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Macrolobium multijugum</i> | (DC.)Benth. | 5 | 3 | 1 | form2 | 0.071429 | partially protected |
| <i>Macrolobium urupaeense</i> | Hoehne | 4 | 1 | 4 | form1 | 0.5 | partially protected |
| <i>Macropeplus ligustrinus</i> | (Tul.) Perkins | 7 | 3 | 5 | common | 0.34375 | partially protected |
| <i>Macropeplus schwackeanus</i> | (Perkins) I.Santos & Peixoto | 1 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Magnolia ovata</i> | (A.St.-Hil.) Spreng. | 8 | 3 | 2 | common | 0.233333 | partially protected |
| <i>Magonia pubescens</i> | A. St.-Hil. | 21 | 3 | 14 | common | 0.116223 | partially protected |
| <i>Malouetia pubescens</i> | Markgr. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Malouetia tamaquarina</i> | (Aubl.) A.DC. | 20 | 3 | 21 | common | 0.3 | partially protected |
| <i>Malpighia emarginata</i> | DC. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Manihot anomala</i> | Pohl | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Manihot caeruleascens</i> | Pohl | 24 | 3 | 17 | common | 0.161491 | partially protected |
| <i>Manihot esculenta</i> | Crantz | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Manihot jacobinensis</i> | MÃƒÂ½ll.Arg. | 3 | 3 | 2 | form4 | 0.571429 | partially protected |
| <i>Manihot maracasensis</i> | Ule | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Manihot tripartita</i> | (Spreng.) MÃƒÂ½ll.Arg. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Manilkara rufula</i> | (Miq.) H.J.Lam | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Manilkara salzmannii</i> | (A.DC.) H.J.Lam | 4 | 3 | 3 | common | 0.333333 | partially protected |
| <i>Manilkara subsericea</i> | (Mart.) Dubard | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Manilkara triflora</i> | (AllemÃƒÂ£o) Monach. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Maprounea brasiliensis</i> | A.St.-Hil. | 25 | 3 | 16 | common | 0.170068 | partially protected |
| <i>Maprounea guianensis</i> | Aubl. | 28 | 3 | 26 | common | 0.233766 | partially protected |
| <i>Margaritaria nobilis</i> | L.f. | 30 | 3 | 24 | common | 0.234783 | partially protected |
| <i>Marlierea buxifolia</i> | Amshoff | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Marlierea excoriata</i> | Mart. | 4 | 3 | 3 | common | 0.444444 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Marlierea involucrata</i> | (O.Berg) Nied. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Marlierea maguirei</i> | McVaugh | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Marlierea neuwiedeana</i> | (O.Berg) Nied. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Marlierea obscura</i> | O.Berg | 2 | 3 | 2 | form4 | 0.444444 | partially protected |
| <i>Marlierea obversa</i> | D.Legrand | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Marlierea tomentosa</i> | Cambess. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Martiodendron elatum</i> | (Ducke) Gleason | 12 | 3 | 10 | common | 0.090909 | partially protected |
| <i>Martiodendron mediterraneum</i> | (Mart. ex Benth.) R.C.Koeppen | 4 | 3 | 4 | common | 0.105263 | partially protected |
| <i>Matayba arborescens</i> | (Aubl.) Radlk. | 23 | 3 | 22 | common | 0.319149 | partially protected |
| <i>Matayba discolor</i> | Radlk. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Matayba elaeagnoides</i> | Radlk. | 4 | 3 | 2 | common | 0.133333 | partially protected |
| <i>Matayba guianensis</i> | Aubl. | 34 | 3 | 41 | common | 0.185658 | partially protected |
| <i>Matayba heterophylla</i> | (Mart.) Radlk. | 10 | 3 | 5 | common | 0.15625 | partially protected |
| <i>Matayba inelegans</i> | Radlk. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Matayba intermedia</i> | Radlk. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Matayba mollis</i> | Radlk. | 5 | 3 | 3 | common | 0.151515 | partially protected |
| <i>Matayba opaca</i> | Radlk. | 17 | 3 | 21 | common | 0.421053 | partially protected |
| <i>Matayba punctata</i> | Radlk. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Matayba purgans</i> | (Poepp.) Radlk. | 13 | 3 | 10 | common | 0.25 | partially protected |
| <i>Matelea maritima</i> | (Jacq.) Woodson | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Mauria thaumatophylla</i> | Loes. | 1 | 1 | 2 | form5 | 1 | protected |
| <i>Mauritia flexuosa</i> | L.f. | 30 | 3 | 31 | common | 0.24812 | partially protected |
| <i>Mauritiella aculeata</i> | (Kunth) Burret | 18 | 3 | 21 | common | 0.176471 | partially protected |
| <i>Mauritiella armata</i> | (Mart.) Burret | 24 | 3 | 29 | common | 0.367089 | partially protected |
| <i>Maytenus boaria</i> | Molina | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Maytenus distichophylla</i> | Mart. | 8 | 3 | 7 | common | 0.266667 | partially protected |
| <i>Maytenus erythroxyla</i> | Reissek | 9 | 3 | 9 | common | 0.28125 | partially protected |
| <i>Maytenus gonoclada</i> | Mart. | 11 | 3 | 5 | common | 0.211538 | partially protected |
| <i>Maytenus guyanensis</i> | Klotzsch ex Reissek | 19 | 3 | 23 | common | 0.428571 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|-------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Maytenus ilicifolia</i> | Mart. ex Reissek | 7 | 3 | 2 | common | 0.083333 | partially protected |
| <i>Maytenus imbricata</i> | Mart. ex Reiss. | 3 | 3 | 1 | form6 | 0.2 | partially protected |
| <i>Maytenus laevis</i> | Reissek | 8 | 3 | 8 | common | 1 | protected |
| <i>Maytenus myrsinoides</i> | Reissek | 7 | 3 | 8 | common | 0.625 | partially protected |
| <i>Maytenus oblongata</i> | Reissek | 3 | 3 | 2 | form4 | 0 | unprotected |
| <i>Maytenus obtusifolia</i> | Mart. | 9 | 3 | 8 | common | 0.619048 | partially protected |
| <i>Maytenus opaca</i> | Reissek | 5 | 3 | 3 | common | 0.333333 | partially protected |
| <i>Maytenus pittieriiana</i> | Steyermark | 7 | 3 | 2 | common | 0 | unprotected |
| <i>Maytenus planifolia</i> | A.C. Sm. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Maytenus rigida</i> | Mart. | 9 | 3 | 3 | common | 0.222222 | partially protected |
| <i>Maytenus schumanniana</i> | Loes. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Melanopsisidium nigrum</i> | Colla | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Melicoccus oliviformis</i> | Kunth | 9 | 3 | 8 | common | 0 | unprotected |
| <i>Meliosma frondosa</i> | Cuatrec. & Idrobo | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Meriania urceolata</i> | Triana | 19 | 3 | 17 | common | 0.5 | partially protected |
| <i>Merianthera eburnea</i> | R.Goldenb. & Fraga | 3 | 3 | 3 | form4 | 0.307692 | partially protected |
| <i>Merianthera sopolisii</i> | (Glaz. & Cogn.) Wurdack | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Metrodorea mollis</i> | Taub. | 6 | 3 | 3 | common | 0.263158 | partially protected |
| <i>Mezilaurus crassiramea</i> | (Meisn.) Taub. ex Mez | 18 | 3 | 9 | common | 0.131579 | partially protected |
| <i>Mezilaurus itauba</i> | (Meisn.) Taub. ex Mez | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mezilaurus vanderwerffii</i> | Alves, F.M. & Baitello | 5 | 1 | 2 | form1 | 0 | unprotected |
| <i>Miconia affinis</i> | DC. | 28 | 3 | 31 | common | 0.311475 | partially protected |
| <i>Miconia alata</i> | (Aubl.) DC. | 19 | 3 | 16 | common | 0.531915 | partially protected |
| <i>Miconia albicans</i> | (Sw.) Steud. | 35 | 3 | 32 | common | 0.165468 | partially protected |
| <i>Miconia alternans</i> | Naudin | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Miconia amoena</i> | Triana | 4 | 3 | 3 | common | 0.285714 | partially protected |
| <i>Miconia ampla</i> | Triana | 12 | 3 | 8 | common | 0.375 | partially protected |
| <i>Miconia aplostachya</i> | (Bonpl.) DC. | 18 | 3 | 20 | common | 0.22 | partially protected |
| <i>Miconia argyrophylla</i> | DC. | 20 | 3 | 21 | common | 0.24 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------|------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Miconia aureoides</i> | Cogn. | 7 | 3 | 3 | common | 0 | unprotected |
| <i>Miconia borjensis</i> | Wurdack | 4 | 3 | 1 | form2 | 0.071429 | partially protected |
| <i>Miconia brevipes</i> | Benth. | 8 | 3 | 7 | common | 0.875 | partially protected |
| <i>Miconia burchellii</i> | Triana | 14 | 3 | 6 | common | 0.28125 | partially protected |
| <i>Miconia caudigera</i> | DC. | 5 | 3 | 4 | common | 0.25 | partially protected |
| <i>Miconia chamissois</i> | Naudin | 18 | 3 | 8 | common | 0.149533 | partially protected |
| <i>Miconia chrysophylla</i> | (Rich.) Urb. | 20 | 3 | 19 | common | 0.444444 | partially protected |
| <i>Miconia cinerascens</i> | Miq. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Miconia corallina</i> | Spring | 3 | 3 | 3 | form4 | 0.47619 | partially protected |
| <i>Miconia crassinervia</i> | Cogn. | 19 | 3 | 12 | common | 0.090909 | partially protected |
| <i>Miconia cuspidata</i> | Mart. ex Naudin | 24 | 3 | 18 | common | 0.304348 | partially protected |
| <i>Miconia elegans</i> | Cogn. | 17 | 3 | 6 | common | 0.257143 | partially protected |
| <i>Miconia fallax</i> | DC. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Miconia ferruginata</i> | DC. | 20 | 3 | 18 | common | 0.157986 | partially protected |
| <i>Miconia flammea</i> | Casar. | 1 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Miconia herpetica</i> | DC. | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Miconia holosericea</i> | (L.) DC. | 23 | 3 | 23 | common | 0.283019 | partially protected |
| <i>Miconia hyemalis</i> | A. St.-Hil. & Naudin | 2 | 3 | 1 | form6 | 0.666667 | partially protected |
| <i>Miconia irwinii</i> | Wurdack | 5 | 3 | 4 | common | 0.363636 | partially protected |
| <i>Miconia latecrenata</i> | (DC.) Naudin | 7 | 3 | 2 | common | 0.277778 | partially protected |
| <i>Miconia lepidota</i> | Schrink & Mart. ex DC. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Miconia leucocarpa</i> | DC. | 10 | 3 | 3 | common | 0.327586 | partially protected |
| <i>Miconia ligustroides</i> | (DC.) Naudin | 17 | 3 | 8 | common | 0.156584 | partially protected |
| <i>Miconia lourteigiana</i> | Wurdack | 9 | 3 | 6 | common | 0.625 | partially protected |
| <i>Miconia lymanii</i> | Wurdack | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Miconia macrothyrsa</i> | Benth. | 23 | 3 | 16 | common | 0.138686 | partially protected |
| <i>Miconia minutiflora</i> | (Bonpl.) DC. | 22 | 3 | 19 | common | 0.267857 | partially protected |
| <i>Miconia paucidens</i> | DC. | 4 | 3 | 2 | common | 0.193548 | partially protected |
| <i>Miconia pepericarpa</i> | Mart. ex DC. | 4 | 3 | 2 | common | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-------------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Miconia phaeophylla</i> | Triana | 16 | 3 | 22 | common | 0.466667 | partially protected |
| <i>Miconia prasina</i> | (Sw.) DC. | 26 | 3 | 26 | common | 0.239437 | partially protected |
| <i>Miconia rubiginosa</i> | (Bonpl.) DC. | 29 | 3 | 23 | common | 0.212598 | partially protected |
| <i>Miconia ruficalyx</i> | Gleason | 13 | 3 | 9 | common | 0.4 | partially protected |
| <i>Miconia sclerophylla</i> | Triana | 7 | 3 | 5 | common | 0.173913 | partially protected |
| <i>Miconia sellowiana</i> | Naudin | 11 | 3 | 5 | common | 0.364583 | partially protected |
| <i>Miconia serialis</i> | DC. | 8 | 3 | 9 | common | 0.333333 | partially protected |
| <i>Miconia setosociliata</i> | Cogn. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Miconia staminea</i> | DC. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Miconia stenostachya</i> | DC. | 34 | 3 | 37 | common | 0.177479 | partially protected |
| <i>Miconia stephananthera</i> | Ule | 12 | 3 | 9 | common | 0.285714 | partially protected |
| <i>Miconia tetraspermoides</i> | Wurdack | 10 | 3 | 8 | common | 0.6 | partially protected |
| <i>Miconia theizans</i> | (Bonpl.) Cogn. | 12 | 3 | 6 | common | 0.19802 | partially protected |
| <i>Miconia tiliifolia</i> | Naudin | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Miconia tillettii</i> | Wurdack | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Miconia tomentosa</i> | (Rich.) D. Don ex DC. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Micranthocereus albicephalus</i> | (Buining & Brederoo) F. Ritter | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Micranthocereus purpureus</i> | (GÃƒÂ¼rke) F. Ritter | 5 | 3 | 3 | common | 0.363636 | partially protected |
| <i>Micropholis gardneriana</i> | (A.DC.) Pierre | 18 | 3 | 13 | common | 0.310345 | partially protected |
| <i>Micropholis suborbicularis</i> | AubrÃƒÂ©v. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Micropholis venulosa</i> | (Mart. & Eichler ex Miq.) Pierre | 5 | 3 | 4 | common | 0.333333 | partially protected |
| <i>Mimosa acutistipula</i> | (Mart.) Benth. | 17 | 3 | 10 | common | 0.254237 | partially protected |
| <i>Mimosa adenophylla</i> | Taub. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mimosa arenosa</i> | (Willd.) Poir. | 13 | 3 | 6 | common | 0.185185 | partially protected |
| <i>Mimosa artemisihana</i> | Heringer & Paula | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Mimosa bimucronata</i> | (DC.) Kuntze | 10 | 3 | 4 | common | 0.045455 | partially protected |
| <i>Mimosa caesalpiniifolia</i> | Benth. | 3 | 3 | 2 | form4 | 0.173913 | partially protected |
| <i>Mimosa claussenii</i> | Benth. | 12 | 3 | 3 | common | 0.163934 | partially protected |
| <i>Mimosa decorticans</i> | Barneby | 4 | 3 | 1 | form2 | 0.333333 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|-----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Mimosa exalbescens</i> | Barneby | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Mimosa flocculosa</i> | Burkart | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Mimosa gemmulata</i> | Barneby | 13 | 3 | 6 | common | 0.144578 | partially protected |
| <i>Mimosa glutinosa</i> | Malme | 3 | 3 | 2 | form4 | 0.1 | partially protected |
| <i>Mimosa hexandra</i> | Micheli | 9 | 3 | 5 | common | 0.4 | partially protected |
| <i>Mimosa huanchacae</i> | Barneby | 2 | 1 | 1 | form7 | 1 | protected |
| <i>Mimosa interrupta</i> | Benth. | 6 | 3 | 2 | common | 0.108696 | partially protected |
| <i>Mimosa irrigua</i> | Barneby | 2 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Mimosa lewisii</i> | Barneby | 3 | 3 | 3 | form4 | 0.2 | partially protected |
| <i>Mimosa manidea</i> | Barneby | 4 | 3 | 2 | common | 0.555556 | partially protected |
| <i>Mimosa microptera</i> | Benth. | 2 | 3 | 2 | form4 | 0.6 | partially protected |
| <i>Mimosa myuros</i> | Barneby | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mimosa nothopteris</i> | Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Mimosa obovata</i> | Benth. | 12 | 3 | 4 | common | 0.061538 | partially protected |
| <i>Mimosa ophthalmocentra</i> | Benth. | 3 | 1 | 2 | form5 | 0.166667 | partially protected |
| <i>Mimosa paraibana</i> | Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Mimosa pithecoloboides</i> | Benth. | 8 | 3 | 5 | common | 0.112676 | partially protected |
| <i>Mimosa pteridifolia</i> | Benth. | 12 | 3 | 5 | common | 0.139241 | partially protected |
| <i>Mimosa regina</i> | Barneby | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mimosa regnellii</i> | Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Mimosa rheiptera</i> | Barneby | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Mimosa scabrella</i> | Benth. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Mimosa schomburgkii</i> | Benth. | 7 | 3 | 5 | common | 0 | unprotected |
| <i>Mimosa sericantha</i> | Benth. | 9 | 3 | 4 | common | 0.109375 | partially protected |
| <i>Mimosa tenuiflora</i> | (Willd.)Poir. | 5 | 3 | 2 | common | 0.25 | partially protected |
| <i>Mimosa verrucosa</i> | Benth. | 10 | 3 | 4 | common | 0.173077 | partially protected |
| <i>Moldenhawera acuminata</i> | Afr. Fernandes & P. Bezerra | 4 | 3 | 3 | common | 0.111111 | partially protected |
| <i>Moldenhawera emarginata</i> | (Spreng.) L.P. Queiroz & Allkin | 1 | 1 | 3 | form5 | 0.666667 | partially protected |
| <i>Moldenhawera nutans</i> | L.P. Queiroz, G.P. Lewis & Allkin | 1 | 1 | 3 | form5 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Mollia burchellii</i> | Sprague | 10 | 3 | 5 | common | 0.172414 | partially protected |
| <i>Mollia lepidota</i> | Spruce ex Benth. | 19 | 3 | 18 | common | 0.307692 | partially protected |
| <i>Mollinedia fruticulosa</i> | Perkins | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Mollinedia glabra</i> | (Spreng.) Perkins | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Moquinia racemosa</i> | (Spreng.) DC. | 8 | 3 | 5 | common | 0.285714 | partially protected |
| <i>Moquiniastrum blanchetianum</i> | (DC.) G. Sancho | 11 | 3 | 6 | common | 0.195122 | partially protected |
| <i>Moquiniastrum floribundum</i> | (Cabrera) G. Sancho | 9 | 3 | 5 | common | 0.1875 | partially protected |
| <i>Moquiniastrum hatschbachii</i> | (Cabrera) G. Sancho | 2 | 1 | 2 | form5 | 0.2 | partially protected |
| <i>Moquiniastrum oligocephalum</i> | (Gardner) G. Sancho | 7 | 3 | 6 | common | 0.4 | partially protected |
| <i>Moquiniastrum paniculatum</i> | (Less.) G. Sancho | 13 | 3 | 7 | common | 0.15493 | partially protected |
| <i>Moquiniastrum polymorphum</i> | (Less.) G. Sancho | 14 | 3 | 8 | common | 0.153846 | partially protected |
| <i>Moquiniastrum pulchrum</i> | (Cabrera) G. Sancho | 7 | 3 | 2 | common | 0.225806 | partially protected |
| <i>Moquiniastrum velutinum</i> | (Bong.) G. Sancho | 4 | 3 | 3 | common | 0.571429 | partially protected |
| <i>Morithamnus ganophyllus</i> | (Mattf. ex Pilg.) R.M.King & H.Rob. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Mouriri apiranga</i> | Spruce ex Triana | 15 | 3 | 16 | common | 0.482759 | partially protected |
| <i>Mouriri arborea</i> | Gardner | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Mouriri cearensis</i> | Huber | 4 | 3 | 4 | common | 0.75 | partially protected |
| <i>Mouriri dumetosa</i> | Cogn. | 10 | 3 | 10 | common | 0.1 | partially protected |
| <i>Mouriri elliptica</i> | Mart. | 20 | 3 | 9 | common | 0.103175 | partially protected |
| <i>Mouriri glazioviana</i> | Cogn. | 10 | 3 | 4 | common | 0.21875 | partially protected |
| <i>Mouriri grandiflora</i> | DC. | 19 | 3 | 21 | common | 0.380952 | partially protected |
| <i>Mouriri guianensis</i> | Aubl. | 30 | 3 | 27 | common | 0.192 | partially protected |
| <i>Mouriri myrtilloides</i> | (Sw.) Poir. | 8 | 3 | 5 | common | 0.2 | partially protected |
| <i>Mouriri pusa</i> | Gardner ex Gardner | 17 | 3 | 15 | common | 0.117117 | partially protected |
| <i>Mouriri vernicosa</i> | Naudin | 12 | 1 | 8 | form1 | 0.6 | partially protected |
| <i>Moutabea excoriata</i> | Mart. ex Miq. | 12 | 3 | 6 | common | 0.380952 | partially protected |
| <i>Moutabea guianensis</i> | Aubl. | 18 | 3 | 20 | common | 0.142857 | partially protected |
| <i>Muellera frutescens</i> | (Aubl.) Standl. | 3 | 1 | 3 | form5 | 0.333333 | partially protected |

| S | author | | lat | population | ecoregion | Form | r | status |
|-------------------------------------|---|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Muellera montana</i> | (M.J. Silva & A.M.G. Azevedo) M.J. Silva & A.M.G. | | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Muellera obtusa</i> | (Benth.) M.J. Silva & A.M.G. Azevedo | | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Muntingia calabura</i> | L. | | 13 | 3 | 10 | common | 0.166667 | partially protected |
| <i>Myracrodrion urundeuva</i> | AllemÃƒÂ£o | | 19 | 3 | 12 | common | 0.088235 | partially protected |
| <i>Myrceugenia acutiflora</i> | (Kiaersk.) D.Legrand & Kausel | | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrceugenia alpigena</i> | (DC.) Landrum | | 10 | 3 | 4 | common | 0.333333 | partially protected |
| <i>Myrceugenia brevipedicellata</i> | (Burret) D.Legrand & Kausel | | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrceugenia campestris</i> | (DC.) D.Legrand & Kausel | | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrcia aegiphylloides</i> | Mattos | | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia albidotomentosa</i> | (Amshoff) McVaugh | | 3 | 1 | 3 | form5 | 1 | protected |
| <i>Myrcia albotomentosa</i> | DC. | | 19 | 3 | 5 | common | 0.116667 | partially protected |
| <i>Myrcia almasensis</i> | NicLugh. | | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Myrcia atramentifera</i> | Barb.Rodr. | | 8 | 3 | 8 | common | 0.666667 | partially protected |
| <i>Myrcia bella</i> | Cambess. | | 15 | 3 | 10 | common | 0.139752 | partially protected |
| <i>Myrcia bergiana</i> | O.Berg | | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Myrcia boliviensis</i> | (Steyer.) McVaugh | | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Myrcia bracteata</i> | (Rich.) DC. | | 21 | 3 | 22 | common | 0.5 | partially protected |
| <i>Myrcia brasiliiae</i> | Mattos & D.Legrand | | 1 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Myrcia brasiliensis</i> | Kiaersk. | | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Myrcia camapuanensis</i> | N.Silveira | | 15 | 3 | 3 | common | 0.12 | partially protected |
| <i>Myrcia chapadensis</i> | S.Moore | | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrcia citrifolia</i> | (Aubl.) Urb. | | 7 | 3 | 9 | common | 0 | unprotected |
| <i>Myrcia cujabensis</i> | O.Berg | | 5 | 3 | 1 | form2 | 0.2 | partially protected |
| <i>Myrcia cuprea</i> | (O.Berg) Kiaersk. | | 9 | 3 | 11 | common | 0.541667 | partially protected |
| <i>Myrcia decorticans</i> | DC. | | 16 | 3 | 13 | common | 0.2 | partially protected |
| <i>Myrcia densa</i> | (DC.) Sobral | | 4 | 3 | 3 | common | 0.3 | partially protected |
| <i>Myrcia eriocalyx</i> | DC. | | 5 | 3 | 3 | common | 0.285714 | partially protected |
| <i>Myrcia fenzliana</i> | O.Berg | | 17 | 3 | 8 | common | 0.179775 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------|---------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Myrcia filibracteata</i> | Mattos & D.Legrand | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia florida</i> | Lem. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrcia guianensis</i> | (Aubl.) DC. | 34 | 3 | 45 | common | 0.148814 | partially protected |
| <i>Myrcia ilheosensis</i> | Kiaersk. | 3 | 1 | 3 | form5 | 0.5 | partially protected |
| <i>Myrcia jacobinensis</i> | Mattos | 7 | 3 | 5 | common | 0.3125 | partially protected |
| <i>Myrcia lanuginosa</i> | O.Berg | 5 | 3 | 3 | common | 0.104478 | partially protected |
| <i>Myrcia laruotteana</i> | Cambess. | 11 | 3 | 6 | common | 0.307692 | partially protected |
| <i>Myrcia lutescens</i> | Cambess. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia mansoniana</i> | O.Berg | 4 | 3 | 2 | common | 0.4 | partially protected |
| <i>Myrcia mischophylla</i> | Kiaersk. | 9 | 3 | 5 | common | 0.22807 | partially protected |
| <i>Myrcia multiflora</i> | (Lam.) DC. | 33 | 3 | 33 | common | 0.237918 | partially protected |
| <i>Myrcia mutabilis</i> | (O.Berg) N.Silveira | 10 | 3 | 6 | common | 0.204545 | partially protected |
| <i>Myrcia obovata</i> | (O.Berg) Nied. | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Myrcia ochroides</i> | O.Berg | 13 | 3 | 3 | common | 0.058824 | partially protected |
| <i>Myrcia paivae</i> | O.Berg | 14 | 3 | 14 | common | 0.4 | partially protected |
| <i>Myrcia palustris</i> | DC. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia parnahibensis</i> | (O.Berg) Kiaersk. | 7 | 3 | 1 | form2 | 0.333333 | partially protected |
| <i>Myrcia platyclada</i> | DC. | 4 | 3 | 3 | common | 1 | protected |
| <i>Myrcia pubipetala</i> | Miq. | 14 | 3 | 6 | common | 0.285714 | partially protected |
| <i>Myrcia pulchra</i> | (O.Berg) Kiaersk. | 6 | 3 | 4 | common | 0.129032 | partially protected |
| <i>Myrcia racemosa</i> | (O.Berg) Kiaersk. | 2 | 3 | 1 | form6 | 0.4 | partially protected |
| <i>Myrcia racemulosa</i> | DC. | 9 | 3 | 4 | common | 0.162791 | partially protected |
| <i>Myrcia ramuliflora</i> | (O.Berg) N.Silveira | 3 | 3 | 3 | form4 | 0.25 | partially protected |
| <i>Myrcia recurvata</i> | O.Berg | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia reticulata</i> | Cambess. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Myrcia retorta</i> | Cambess. | 8 | 3 | 4 | common | 0.079545 | partially protected |
| <i>Myrcia revolutifolia</i> | McVaugh | 4 | 3 | 5 | common | 1 | protected |
| <i>Myrcia rimosaa</i> | Cambess. | 1 | 1 | 2 | form5 | 0 | unprotected |
| <i>Myrcia rosangelae</i> | NicLugh. | 4 | 3 | 5 | common | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Myrcia rotundata</i> | (Amshoff) McVaugh | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Myrcia rotundifolia</i> | (O.Berg) Kiaersk. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia rufipes</i> | DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia spectabilis</i> | DC. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Myrcia splendens</i> | (Sw.) DC. | 34 | 3 | 46 | common | 0.175732 | partially protected |
| <i>Myrcia subalpestris</i> | DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia subavenia</i> | (O.Berg) N.Silveira | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Myrcia subcordata</i> | DC. | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Myrcia subsessilis</i> | O.Berg | 14 | 3 | 8 | common | 0.8125 | partially protected |
| <i>Myrcia sylvatica</i> | (G.Mey.) DC. | 21 | 3 | 30 | common | 0.309524 | partially protected |
| <i>Myrcia tenuifolia</i> | (O.Berg) Sobral | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrcia tomentosa</i> | (Aubl.) DC. | 33 | 3 | 28 | common | 0.149744 | partially protected |
| <i>Myrcia uberavensis</i> | O.Berg | 10 | 3 | 2 | common | 0.112903 | partially protected |
| <i>Myrcia variabilis</i> | Mart. ex DC. | 16 | 3 | 8 | common | 0.084906 | partially protected |
| <i>Myrcia vauthiereana</i> | O.Berg | 1 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Myrcia venulosa</i> | DC. | 13 | 3 | 8 | common | 0.188172 | partially protected |
| <i>Myrcia vestita</i> | DC. | 13 | 3 | 6 | common | 0.122449 | partially protected |
| <i>Myrcia vittoriana</i> | Kiaersk. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Myrcianthes cavalcantei</i> | Mattos | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Myrciaria cuspidata</i> | O.Berg | 9 | 3 | 6 | common | 0.181818 | partially protected |
| <i>Myrciaria delicatula</i> | (DC.) O.Berg | 5 | 3 | 3 | common | 0.375 | partially protected |
| <i>Myrciaria floribunda</i> | (H.West ex Willd.) O.Berg | 28 | 3 | 29 | common | 0.27439 | partially protected |
| <i>Myrciaria strigipes</i> | O.Berg | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Myrciaria tenella</i> | (DC.) O.Berg | 21 | 3 | 16 | common | 0.275862 | partially protected |
| <i>Myrocarpus fastigiatus</i> | Allemand, Afo | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrospermum frutescens</i> | Jacq. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Myroxylon peruferum</i> | L.f. | 10 | 3 | 3 | common | 0.25 | partially protected |
| <i>Myrsine coriacea</i> | (Sw.) R.Br. ex Roem. & Schult. | 19 | 3 | 12 | common | 0.179054 | partially protected |
| <i>Myrsine emarginella</i> | Miq. | 7 | 3 | 5 | common | 0.263158 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------------|--------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Myrsine guianensis</i> | (Aubl.) Kuntze | 30 | 3 | 28 | common | 0.15796 | partially protected |
| <i>Myrsine lancifolia</i> | Mart. | 3 | 3 | 3 | form4 | 0.37931 | partially protected |
| <i>Myrsine leuconeura</i> | Mart. | 10 | 3 | 6 | common | 0.119048 | partially protected |
| <i>Myrsine lineata</i> | (Mez) Imkhan. | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Myrsine monticola</i> | Mart. | 12 | 3 | 5 | common | 0.155172 | partially protected |
| <i>Myrsine nitida</i> | (Mez) Pipoly | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Myrsine parvifolia</i> | A. DC. | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Myrsine parvula</i> | (Mez) Otegui | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Myrsine squarrosa</i> | (Mez) M.F.Freitas & Kin.-Gouv. | 4 | 3 | 5 | common | 0.230769 | partially protected |
| <i>Myrsine umbellata</i> | Mart. | 22 | 3 | 20 | common | 0.175141 | partially protected |
| <i>Myrsine venosa</i> | A. DC. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Myrsine villosissima</i> | Mart. | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Nectandra cissiflora</i> | Nees | 24 | 3 | 16 | common | 0.184783 | partially protected |
| <i>Nectandra cuspidata</i> | Nees & Mart. | 3 | 3 | 2 | form4 | 0.25 | partially protected |
| <i>Nectandra hihua</i> | (Ruiz & Pav.) Rohwer | 19 | 3 | 16 | common | 0.3 | partially protected |
| <i>Nectandra megapotamica</i> | (Spreng.) Mez | 6 | 3 | 2 | common | 0.111111 | partially protected |
| <i>Nectandra psammophila</i> | Nees & C. Mart. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Neea floribunda</i> | Poepp. & Endl. | 18 | 3 | 18 | common | 0.3 | partially protected |
| <i>Neea hermaphrodita</i> | S. Moore | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Neea macrophylla</i> | Poepp. & Endl. | 21 | 3 | 16 | common | 0.4 | partially protected |
| <i>Neea parviflora</i> | Poepp. & Endl. | 9 | 1 | 9 | form1 | 0.8 | partially protected |
| <i>Neea theifera</i> | Oerst. | 17 | 3 | 14 | common | 0.14321 | partially protected |
| <i>Neocabreria pennivenia</i> | (B.L.Rob.) R.M.King & H.Rob. | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Neomitranthes langsdorffii</i> | (O.Berg) Mattos | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Neomitranthes obscura</i> | (DC.) N.Silveira | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Norantea goyasensis</i> | Cambess. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Ochthocosmus attenuatus</i> | Steyermark & Luteyn | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Ochthocosmus barrae</i> | Hallier f. | 3 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Ochthocosmus longipedicellatus</i> | Steyermark & Luteyn | 3 | 3 | 2 | form4 | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------|-------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Ochthocosmus roraimae</i> | Benth. | 4 | 3 | 3 | common | 0.666667 | partially protected |
| <i>Ocotea aciphylla</i> | (Nees & Mart.) Mez | 27 | 3 | 25 | common | 0.348837 | partially protected |
| <i>Ocotea adamantina</i> | P.L.R. Moraes e van der Werff | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Ocotea arenicola</i> | L.C.S. Assis e Mello-Silva | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ocotea brachybotrya</i> | (Meisn.) Mez | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Ocotea canaliculata</i> | (Rich.) Mez | 20 | 3 | 19 | common | 0.188679 | partially protected |
| <i>Ocotea cernua</i> | (Nees) Mez | 23 | 3 | 27 | common | 0.45098 | partially protected |
| <i>Ocotea corymbosa</i> | (Meisn.) Mez | 15 | 3 | 12 | common | 0.087629 | partially protected |
| <i>Ocotea daphnifolia</i> | (Meisn.) Mez | 3 | 1 | 3 | form5 | 0.4 | partially protected |
| <i>Ocotea diospyrifolia</i> | (Meisn.) Mez | 11 | 3 | 6 | common | 0.152174 | partially protected |
| <i>Ocotea duidensis</i> | Moldenke | 3 | 3 | 2 | form4 | 0.666667 | partially protected |
| <i>Ocotea fasciculata</i> | (Nees) Mez | 13 | 3 | 13 | common | 0.117647 | partially protected |
| <i>Ocotea floribunda</i> | (Sw.) Mez | 10 | 3 | 12 | common | 0.25 | partially protected |
| <i>Ocotea glauca</i> | (Nees & Mart.) Mez | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Ocotea gracilis</i> | (Meisn.) Mez | 23 | 3 | 21 | common | 0.178571 | partially protected |
| <i>Ocotea lancifolia</i> | (Schott) Mez | 11 | 3 | 7 | common | 0.243243 | partially protected |
| <i>Ocotea langsdorffii</i> | (Meisn.) Mez | 6 | 3 | 4 | common | 0.421053 | partially protected |
| <i>Ocotea lobbii</i> | (Meisn.) Rohwer | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ocotea minarum</i> | (Nees & Mart.) Mez | 10 | 3 | 6 | common | 0.100719 | partially protected |
| <i>Ocotea notata</i> | (Nees & Mart.) Mez | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Ocotea nutans</i> | (Nees) Mez | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ocotea percoriacea</i> | Kosterm. | 9 | 3 | 5 | common | 0.25 | partially protected |
| <i>Ocotea polyantha</i> | (Nees & Mart.) Mez | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Ocotea pomaderroides</i> | (Meisn.) Mez | 10 | 3 | 6 | common | 0.208092 | partially protected |
| <i>Ocotea pulchella</i> | (Nees & Mart.) Mez | 10 | 3 | 8 | common | 0.101449 | partially protected |
| <i>Ocotea rohweri</i> | P.L.R. Moraes van der Werff | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ocotea semicompleta</i> | (Nees & Mart.) Mez | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Ocotea silvestris</i> | Vattimo | 1 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Ocotea spectabilis</i> | (Meisn.) Mez | 16 | 3 | 8 | common | 0.181818 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Ocotea spixiana</i> | (Nees) Mez | 9 | 3 | 5 | common | 0.325843 | partially protected |
| <i>Ocotea tristis</i> | (Nees & Mart.) Mez | 7 | 3 | 5 | common | 0.3 | partially protected |
| <i>Ocotea vaccinioides</i> | (Meisn.) Mez | 1 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Ocotea vegrandidis</i> | P.L.R. Moraes et van der Werff | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Ocotea velloziana</i> | (Meisn.) Mez | 9 | 3 | 7 | common | 0.307692 | partially protected |
| <i>Ocotea velutina</i> | (Nees) Mart. ex B.D.Jacks. | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Ocotea xanthocalyx</i> | (Nees) Mez | 6 | 3 | 1 | form2 | 0.2 | partially protected |
| <i>Oenocarpus bataua</i> | Mart. | 21 | 3 | 20 | common | 0.25 | partially protected |
| <i>Oenocarpus distichus</i> | Mart. | 20 | 3 | 12 | common | 0.404762 | partially protected |
| <i>Oocephalus piranii</i> | (Harley) Harley & J.F.B. Pastore | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Opuntia monacantha</i> | (Willd.) Haw. | 4 | 3 | 6 | common | 0 | unprotected |
| <i>Ormosia arborea</i> | (Vell.) Harms | 4 | 3 | 2 | common | 0.333333 | partially protected |
| <i>Ormosia costulata</i> | (Miq.) Kleinhoonte | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ormosia fastigiata</i> | Tul. | 11 | 3 | 6 | common | 0.191489 | partially protected |
| <i>Ouratea apurensis</i> | Sastre | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ouratea blanchetiana</i> | Engl. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ouratea castaneifolia</i> | (DC.) Engl. | 28 | 3 | 26 | common | 0.211832 | partially protected |
| <i>Ouratea cuspidata</i> | Tiegh. | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Ouratea glaucescens</i> | Engl. | 9 | 3 | 3 | common | 0.24 | partially protected |
| <i>Ouratea grosourdyi</i> | (Tiegh.) Steyermark | 3 | 3 | 1 | form6 | 0 | unprotected |
| <i>Ouratea hexasperma</i> | (A. St.-Hil.) Baill. | 24 | 3 | 17 | common | 0.121839 | partially protected |
| <i>Ouratea ovalis</i> | Engl. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ouratea polyantha</i> | (Triana & Planch.) Engl. | 9 | 3 | 8 | common | 0.041667 | partially protected |
| <i>Ouratea polygyna</i> | Engl. | 11 | 3 | 8 | common | 0.5625 | partially protected |
| <i>Ouratea roraimae</i> | Engl. | 11 | 3 | 10 | common | 0 | unprotected |
| <i>Ouratea salicifolia</i> | Engl. | 6 | 3 | 4 | common | 0.166667 | partially protected |
| <i>Ouratea sellowii</i> | Engl. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ouratea semiserrata</i> | (Mart. & Nees) Engl. | 4 | 3 | 3 | common | 0.333333 | partially protected |
| <i>Ouratea spectabilis</i> | (Mart. ex Engl.) Engl. | 22 | 3 | 14 | common | 0.130064 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|-----------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Ouratea subamplexicaulis</i> | Maguire & Steyermark. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Ouratea tatei</i> | Gleason | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Oxandra reticulata</i> | Maas | 13 | 3 | 6 | common | 0.2 | partially protected |
| <i>Oxandra sessiliflora</i> | R.E.Fr. | 14 | 3 | 10 | common | 0.173913 | partially protected |
| <i>Pachira flaviflora</i> | (Pulle) Fern.Alonso | 3 | 3 | 3 | form4 | 1 | protected |
| <i>Pachira minor</i> | (Sims) Hemsl. | 4 | 3 | 3 | common | 0.666667 | partially protected |
| <i>Pachira nitida</i> | Kunth | 15 | 3 | 15 | common | 0.307692 | partially protected |
| <i>Pachira paraensis</i> | (Ducke) W.S.Alverson | 13 | 3 | 12 | common | 0.444444 | partially protected |
| <i>Pagamea capitata</i> | Benth. | 11 | 3 | 11 | common | 0.625 | partially protected |
| <i>Pagamea guianensis</i> | Aubl. | 24 | 3 | 27 | common | 0.333333 | partially protected |
| <i>Pagamea thyrsiflora</i> | Spruce ex Benth. | 16 | 3 | 14 | common | 0.5625 | partially protected |
| <i>Palicourea blanchetiana</i> | Schltdl. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Palicourea crocea</i> | (Sw.) Schult. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Palicourea grandifolia</i> | (Willd. ex Schult.) Standl. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Palicourea guianensis</i> | Aubl. | 21 | 3 | 25 | common | 0.384615 | partially protected |
| <i>Palicourea longistipulata</i> | Standl. | 3 | 1 | 2 | form5 | 0.2 | partially protected |
| <i>Palicourea marcgravii</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Palicourea nitidella</i> | (MÃƒÂ¼ll.Arg.) Standl. | 3 | 3 | 4 | form4 | 0.142857 | partially protected |
| <i>Palicourea rigida</i> | Kunth | 32 | 3 | 24 | common | 0.165511 | partially protected |
| <i>Panopsis rubescens</i> | (Pohl) Pittier | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Panopsis sessilifolia</i> | (Rich.) Sandwith | 8 | 3 | 12 | common | 0.75 | partially protected |
| <i>Paralychnophora bicolor</i> | (DC.) MacLeish | 9 | 3 | 5 | common | 0.368421 | partially protected |
| <i>Paralychnophora glaziouana</i> | Loeuille | 5 | 3 | 2 | common | 0.25 | partially protected |
| <i>Paralychnophora harleyi</i> | (H. Rob.) D.J.N. Hind | 4 | 3 | 3 | common | 0.444444 | partially protected |
| <i>Parinari campestris</i> | Aubl. | 15 | 3 | 20 | common | 0.257143 | partially protected |
| <i>Parinari maguirei</i> | Prance | 3 | 1 | 3 | form5 | 1 | protected |
| <i>Parinari obtusifolia</i> | Hook.f. | 14 | 3 | 7 | common | 0.124402 | partially protected |
| <i>Parkia cachimboensis</i> | H.C.Hopkins | 6 | 1 | 4 | form1 | 0 | unprotected |
| <i>Parkia pendula</i> | (Willd.)Walp. | 1 | 1 | 1 | form7 | 1 | protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-----------------------------------|-----------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Parkia platycephala</i> | Benth. | 14 | 3 | 9 | common | 0.166667 | partially protected |
| <i>Paubrasilia echinata</i> | (Lam.) E. Gagnon, H.C. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Peltogyne campestris</i> | Ducke | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Peltogyne confertiflora</i> | (Hayne)Benth. | 20 | 3 | 9 | common | 0.103448 | partially protected |
| <i>Peltogyne crenulata</i> | Afr.Fern. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Peltogyne maranhensis</i> | Ducke | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Peltophorum dubium</i> | (Spreng.) Taub. | 9 | 3 | 3 | common | 0.085714 | partially protected |
| <i>Pera bicolor</i> | (Klotzsch) MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Pera decipiens</i> | (MÃƒÂ½ll.Arg.) MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Pera distichophylla</i> | (Mart.) Baill. | 17 | 3 | 16 | common | 0.222222 | partially protected |
| <i>Pera furfuracea</i> | MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Pera glabrata</i> | (Schott) Poepp. ex Baill. | 34 | 3 | 41 | common | 0.151079 | partially protected |
| <i>Pera heterantha</i> | (Schrank) I.M.Johnst. | 8 | 3 | 7 | common | 0.5 | partially protected |
| <i>Pereskia guamacho</i> | F.A.C.Weber | 3 | 3 | 1 | form6 | 0 | unprotected |
| <i>Persea alba</i> | Nees & Mart. | 6 | 3 | 5 | common | 0.272727 | partially protected |
| <i>Persea fusca</i> | Mez | 5 | 3 | 1 | form2 | 0.444444 | partially protected |
| <i>Persea glabra</i> | van der Werff | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Persea pedunculosa</i> | Meisn. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Persea rufotomentosa</i> | Nees & C. Mart. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Persea splendens</i> | Meisn. | 9 | 3 | 2 | common | 0 | unprotected |
| <i>Persea venosa</i> | Nees & Mart. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Phyllanthus acuminatus</i> | Vahl | 25 | 3 | 21 | common | 0.169492 | partially protected |
| <i>Phyllanthus attenuatus</i> | Miq. | 13 | 3 | 13 | common | 0.347826 | partially protected |
| <i>Phyllanthus bahiensis</i> | MÃƒÂ½ll.Arg. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Phyllanthus juglandifolius</i> | Willd. | 18 | 3 | 19 | common | 0.5 | partially protected |
| <i>Physocalymma scaberrimum</i> | Pohl | 18 | 3 | 14 | common | 0.247706 | partially protected |
| <i>Picramnia ciliata</i> | Mart. | 2 | 3 | 3 | form4 | 0.6 | partially protected |
| <i>Picramnia guianensis</i> | (Aubl.) Jans.-Jac. | 4 | 1 | 4 | form1 | 1 | protected |
| <i>Picramnia sellowii</i> | Planch. | 22 | 3 | 16 | common | 0.125 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|-------------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Pilocarpus microphyllus</i> | Stapf ex Wardleworth | 5 | 3 | 4 | common | 0.777778 | partially protected |
| <i>Pilosocereus aurisetus</i> | (Werderm.) Byles & G.D. Rowley | 5 | 3 | 3 | common | 0.307692 | partially protected |
| <i>Pilosocereus brasiliensis</i> | (Britton & Rose) Backeb. | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Pilosocereus catingicola</i> | (GÃƒÂ¼rke) Byles & G.D. Rowley | 8 | 3 | 6 | common | 0.25 | partially protected |
| <i>Pilosocereus densiareolatus</i> | F. Ritter | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Pilosocereus flavigibbosus</i> | (Buining & Brederoo) F. Ritter | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Pilosocereus fulvilanatus</i> | (Buining & Brederoo) F. Ritter | 3 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Pilosocereus glaucochrous</i> | (Werderm.) Byles & G.D. Rowley | 3 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Pilosocereus gounellei</i> | (F.A.C.Weber ex K.Schum.) Byles & G.D. Rowley | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Pilosocereus jauruensis</i> | (Buining & Brederoo) P.J.Braun | 7 | 3 | 4 | common | 0.066667 | partially protected |
| <i>Pilosocereus machristii</i> | (E.Y.Dawson) Backeb. | 12 | 3 | 3 | common | 0.130435 | partially protected |
| <i>Pilosocereus pachycladus</i> | F. Ritter | 6 | 3 | 3 | common | 0.2 | partially protected |
| <i>Pilosocereus pentaedrophorus</i> | (Labour.) Byles & G.D. Rowley | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Pilosocereus piauhyensis</i> | (GÃƒÂ¼rke) Byles & G.D. Rowley | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Pilosocereus tuberculatus</i> | (Werderm.) Byles & G.D. Rowley | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Pilosocereus vilaboensis</i> | (Diers & Esteves) P.J. Braun | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Pimenta pseudocaryophyllus</i> | (Gomes) Landrum | 7 | 3 | 6 | common | 0.4 | partially protected |
| <i>Piper aduncum</i> | L. | 27 | 3 | 26 | common | 0.177419 | partially protected |
| <i>Piper amalago</i> | L. | 19 | 3 | 13 | common | 0.206897 | partially protected |
| <i>Piper arboreum</i> | Aubl. | 29 | 3 | 26 | common | 0.288462 | partially protected |
| <i>Piper hispidum</i> | Sw. | 23 | 3 | 19 | common | 0.446809 | partially protected |
| <i>Piper mollicomum</i> | Kunth | 12 | 3 | 9 | common | 0.466667 | partially protected |
| <i>Piper tuberculatum</i> | Jacq. | 23 | 3 | 21 | common | 0.241379 | partially protected |
| <i>Piptadenia gonoacantha</i> | (Mart.) J.F.Macbr. | 5 | 3 | 3 | common | 0 | unprotected |
| <i>Piptadenia macradenia</i> | Benth. | 5 | 3 | 3 | common | 0.266667 | partially protected |
| <i>Piptadenia robusta</i> | Pittier | 5 | 3 | 3 | common | 0.166667 | partially protected |
| <i>Piptadenia stipulacea</i> | (Benth.) Ducke | 3 | 3 | 2 | form4 | 0.2 | partially protected |
| <i>Piptadenia viridiflora</i> | (Kunth) Benth. | 4 | 3 | 4 | common | 0.25 | partially protected |
| <i>Piptocarpha macropoda</i> | (DC.) Baker | 1 | 1 | 1 | form7 | 1 | protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|-----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Piptocarpha regnellii</i> | (Sch.Bip.) Cabrera | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Piptocarpha rotundifolia</i> | (Less.) Baker | 21 | 3 | 11 | common | 0.116395 | partially protected |
| <i>Piptocoma roraimensis</i> | (Steyermark.) Pruski | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Piptocoma schomburgkii</i> | (Sch.Bip.) Pruski | 5 | 3 | 5 | common | 0.666667 | partially protected |
| <i>Piptolepis monticola</i> | Loeuille | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Piranhea trifoliata</i> | Baill. | 12 | 3 | 13 | common | 0.25 | partially protected |
| <i>Pithecellobium dulce</i> | (Roxb.) Benth. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Pithecellobium roseum</i> | (Vahl) Barneby & J.W. Grimes | 5 | 3 | 3 | common | 0.1875 | partially protected |
| <i>Pityrocarpa moniliformis</i> | (Benth.) Luckow & R. W. Jobson | 16 | 3 | 9 | common | 0.15873 | partially protected |
| <i>Pityrocarpa obliqua</i> | (Pers.) Brenan | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Plathymenia foliolosa</i> | Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Plathymenia reticulata</i> | Benth. | 26 | 3 | 23 | common | 0.145125 | partially protected |
| <i>Platonia insignis</i> | Mart. | 20 | 3 | 21 | common | 0.157895 | partially protected |
| <i>Platymiscium floribundum</i> | Vogel | 16 | 3 | 7 | common | 0.243902 | partially protected |
| <i>Platymiscium pinnatum</i> | (Jacq.) Dugand | 22 | 3 | 17 | common | 0.208333 | partially protected |
| <i>Platypodium elegans</i> | Vogel | 24 | 3 | 20 | common | 0.136139 | partially protected |
| <i>Plenckia bahiensis</i> | Loes. | 1 | 1 | 2 | form5 | 1 | protected |
| <i>Plenckia populnea</i> | Reissek | 18 | 3 | 13 | common | 0.135501 | partially protected |
| <i>Pleroma candolleana</i> | (Mart. ex DC.) Triana | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Pleroma echinata</i> | Gardner | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Pleroma foveolata</i> | (Naudin) Triana | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Pleroma stenocarpa</i> | (Schrantz et Mart. ex DC.) Triana | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Podocarpus celatus</i> | de Laub. | 3 | 1 | 3 | form5 | 0.666667 | partially protected |
| <i>Podocarpus lambertii</i> | Klotzsch ex Endl. | 4 | 1 | 2 | form1 | 0.25 | partially protected |
| <i>Podocarpus sellowii</i> | Klotzsch ex Endl. | 12 | 3 | 9 | common | 0.30303 | partially protected |
| <i>Podocarpus transiens</i> | (Pilg.) de Laub. | 3 | 1 | 3 | form5 | 0.25 | partially protected |
| <i>Poecilanthe retusa</i> | Tul. | 3 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Poecilanthe falcata</i> | (Vell.) Heringer | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Poecilanthe ulei</i> | (Harms) Arroyo & Rudd | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Poeppigia procera</i> | C.Presl | 13 | 3 | 10 | common | 0.285714 | partially protected |
| <i>Pogonophora schomburgkiana</i> | Miers ex Benth. | 17 | 3 | 24 | common | 0.352941 | partially protected |
| <i>Poincianella bracteosa</i> | (Tul.) L.P.Queiroz | 10 | 3 | 6 | common | 0.125 | partially protected |
| <i>Posoqueria latifolia</i> | (Rudge) Schult. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Pouteria andarahiensis</i> | T.D.Penn. | 4 | 3 | 3 | common | 0.333333 | partially protected |
| <i>Pouteria caimito</i> | (Ruiz & Pav.) Radlk. | 20 | 3 | 26 | common | 0.45 | partially protected |
| <i>Pouteria coelomatica</i> | Rizzini | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Pouteria cuspidata</i> | (A.DC.) Baehni | 19 | 3 | 21 | common | 0.411765 | partially protected |
| <i>Pouteria furcata</i> | T.D.Penn. | 5 | 3 | 2 | common | 0.026316 | partially protected |
| <i>Pouteria gardneri</i> | (Mart. & Eichler ex Miq.) Baehni | 6 | 3 | 3 | common | 0 | unprotected |
| <i>Pouteria gardneriana</i> | (A.DC.) Radlk. | 11 | 3 | 8 | common | 0.166667 | partially protected |
| <i>Pouteria grandiflora</i> | (A.DC.) Baehni | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Pouteria guianensis</i> | Aubl. | 19 | 3 | 23 | common | 0.354839 | partially protected |
| <i>Pouteria macahensis</i> | T.D.Penn. | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Pouteria peduncularis</i> | (Mart. & Eichler ex Miq.) Baehni | 2 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Pouteria psammophila</i> | (Mart.) Radlk. | 6 | 3 | 5 | common | 0.25 | partially protected |
| <i>Pouteria ramiflora</i> | (Mart.) Radlk. | 28 | 3 | 23 | common | 0.15732 | partially protected |
| <i>Pouteria reticulata</i> | (Engl.) Eyma | 24 | 3 | 27 | common | 0.323944 | partially protected |
| <i>Pouteria rigida</i> | (Mart. & Eichler ex Miq.) Radlk. | 7 | 3 | 6 | common | 0.75 | partially protected |
| <i>Pouteria tenuisepala</i> | Pires & T.D.Penn. | 5 | 3 | 4 | common | 0.142857 | partially protected |
| <i>Pouteria torta</i> | (Mart.) Radlk. | 29 | 3 | 34 | common | 0.125592 | partially protected |
| <i>Pradosia beardii</i> | (Monach.) T.D.Penn. | 3 | 3 | 5 | form4 | 1 | protected |
| <i>Pradosia lactescens</i> | (Vell.) Radlk. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Pradosia longipedicellata</i> | Alves-Ara??jo & M.Alves | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Priogymnanthus hasslerianus</i> | (Chodat) P.S.Green | 21 | 3 | 9 | common | 0.111111 | partially protected |
| <i>Protium bahianum</i> | D.C. Daly | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Protium brasiliense</i> | Engl. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Protium heptaphyllum</i> | (Aubl.) Marchand | 34 | 3 | 44 | common | 0.175875 | partially protected |
| <i>Protium icicariba</i> | (DC.) Marchand | 2 | 3 | 1 | form6 | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|-------------------------|
| <i>Protium ovatum</i> | Engl. | 15 | 3 | 9 | common | 0.131343 | partially protected |
| <i>Protium spruceanum</i> | (Benth.) Engl. | 28 | 3 | 27 | common | 0.231405 | partially protected |
| <i>Protium unifoliolatum</i> | Engl. | 22 | 3 | 26 | common | 0.265625 | partially protected |
| <i>Prunus myrtifolia</i> | (L.) Urb. | 22 | 3 | 21 | common | 0.175758 | partially protected |
| <i>Pseudobombax campestre</i> | (Mart. & Zucc.) A. Robyns | 8 | 3 | 5 | common | 0.125 | partially protected |
| <i>Pseudobombax grandiflorum</i> | (Cav.) A.Robyns | 5 | 3 | 4 | common | 0.333333 | partially protected |
| <i>Pseudobombax longiflorum</i> | (Mart. & Zucc.) A.Robyns | 20 | 3 | 17 | common | 0.133603 | partially protected |
| <i>Pseudobombax marginatum</i> | (A.St.-Hil.) A.Robyns | 21 | 3 | 15 | common | 0.181818 | partially protected |
| <i>Pseudobombax minimum</i> | Carv.-Sobr. & L.P.Queiroz | 4 | 3 | 3 | common | 0.074074 | partially protected |
| <i>Pseudobombax parvifolium</i> | Carv.-Sobr. & L.P.Queiroz | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Pseudobombax riopretensis</i> | Ravenna | 1 | 1 | 2 | form5 | | 0.4 partially protected |
| <i>Pseudobombax tomentosum</i> | (Mart. & Zucc.) A.Robyns | 18 | 3 | 9 | common | 0.124722 | partially protected |
| <i>Pseudobrickellia angustissima</i> | (Spreng. ex Baker) R.M.King & H.Rob. | 9 | 3 | 5 | common | 0.293333 | partially protected |
| <i>Pseudolmedia laevigata</i> | TrÃƒÂfÃ©cul | 1 | 1 | 1 | form7 | | 1 protected |
| <i>Pseudopiptadenia brenanii</i> | G.P.Lewis & M.P.Lima | 4 | 3 | 2 | common | 0.333333 | partially protected |
| <i>Pseudoxandra polyphleba</i> | (Diels) R.E. Fr. | 1 | 1 | 1 | form7 | | 0 unprotected |
| <i>Psidium acutangulum</i> | Mart. ex DC. | 9 | 3 | 8 | common | 0.266667 | partially protected |
| <i>Psidium appendiculatum</i> | Kiaersk. | 6 | 3 | 5 | common | 0.375 | partially protected |
| <i>Psidium brownianum</i> | Mart. ex DC. | 2 | 3 | 1 | form6 | | 0.5 partially protected |
| <i>Psidium canum</i> | Mattos | 5 | 3 | 1 | form2 | 0.277778 | partially protected |
| <i>Psidium cattleianum</i> | Afzel. ex Sabine | 6 | 3 | 3 | common | 0.2 | partially protected |
| <i>Psidium firmum</i> | O.Berg | 1 | 1 | 1 | form7 | | 0 unprotected |
| <i>Psidium guajava</i> | L. | 22 | 3 | 20 | common | 0.296296 | partially protected |
| <i>Psidium guineense</i> | Sw. | 34 | 3 | 39 | common | 0.2 | partially protected |
| <i>Psidium myrsinoides</i> | DC. | 22 | 3 | 13 | common | 0.139368 | partially protected |
| <i>Psidium nutans</i> | O.Berg | 6 | 3 | 4 | common | 0.214286 | partially protected |
| <i>Psidium oligospermum</i> | Mart. ex DC. | 11 | 3 | 4 | common | 0.24 | partially protected |
| <i>Psidium rufum</i> | Mart. ex DC. | 12 | 3 | 4 | common | 0.155556 | partially protected |
| <i>Psidium salutare</i> | var. decussatum (DC.) Landrum | 3 | 3 | 1 | form6 | 0.25 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Psidium schenckianum</i> | Kiaersk. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Psychotria bahiensis</i> | DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Psychotria carthagensis</i> | Jacq. | 25 | 3 | 20 | common | 0.123288 | partially protected |
| <i>Psychotria colorata</i> | (Willd. ex Schult.) MÃƒÂ½ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Psychotria deflexa</i> | DC. | 21 | 3 | 21 | common | 0.354839 | partially protected |
| <i>Psychotria ernestii</i> | K.Krause | 8 | 1 | 8 | form1 | 1 | protected |
| <i>Psychotria hoffmannseggiana</i> | (Willd. ex Schult.) MÃƒÂ½ll.Arg. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Psychotria mapourioides</i> | DC. | 24 | 3 | 22 | common | 0.368421 | partially protected |
| <i>Psychotria vellosiana</i> | Benth. | 10 | 3 | 6 | common | 0.152174 | partially protected |
| <i>Pterocarpus officinalis</i> | Jacq. | 12 | 3 | 14 | common | 0.285714 | partially protected |
| <i>Pterocarpus rohrii</i> | Vahl | 24 | 3 | 23 | common | 0.354839 | partially protected |
| <i>Pterocarpus santalinoides</i> | DC. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Pterocarpus villosus</i> | (Benth.)Benth. | 3 | 3 | 3 | form4 | 0.333333 | partially protected |
| <i>Pterocarpus violaceus</i> | Vogel | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Pterodon abruptus</i> | (Moric.)Benth. | 4 | 3 | 4 | common | 0.166667 | partially protected |
| <i>Pterodon emarginatus</i> | Vogel | 21 | 3 | 13 | common | 0.122563 | partially protected |
| <i>Pterogyne nitens</i> | Tul. | 17 | 3 | 12 | common | 0.078431 | partially protected |
| <i>Ptilochaeta bahiensis</i> | Turcz. | 7 | 3 | 6 | common | 0.138889 | partially protected |
| <i>Quadrella odoratissima</i> | (Jacq.) Hutch. | 3 | 3 | 2 | form4 | 0 | unprotected |
| <i>Qualea acuminata</i> | Spruce ex Warm. | 13 | 3 | 13 | common | 0 | unprotected |
| <i>Qualea cordata</i> | Spreng. | 16 | 3 | 9 | common | 0.13 | partially protected |
| <i>Qualea cryptantha</i> | (Spreng.) Warm. | 4 | 3 | 4 | common | 0.5 | partially protected |
| <i>Qualea densiflora</i> | Warm. | 8 | 3 | 2 | common | 0.272727 | partially protected |
| <i>Qualea dichotoma</i> | (Mart.) Warm. ex Wille | 15 | 3 | 8 | common | 0.163934 | partially protected |
| <i>Qualea elegans</i> | Taub. ex Benoist | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Qualea grandiflora</i> | Mart. | 26 | 3 | 23 | common | 0.137775 | partially protected |
| <i>Qualea hannekesaskiarum</i> | Marc.-Berti | 3 | 3 | 1 | form6 | 0.6 | partially protected |
| <i>Qualea lundii</i> | (Warm.) Warm. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Qualea multiflora</i> | Mart. | 20 | 3 | 17 | common | 0.132302 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------------|---------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Qualea parviflora</i> | Mart. | 23 | 3 | 16 | common | 0.133202 | partially protected |
| <i>Quiabentia zehntneri</i> | (Britton & Rose) Britton & Rose | 3 | 1 | 2 | form5 | 0 | unprotected |
| <i>Randia aculeata</i> | L. | 2 | 3 | 1 | form6 | 0 | unprotected |
| <i>Randia armata</i> | (Sw.) DC. | 29 | 3 | 29 | common | 0.258278 | partially protected |
| <i>Randia calycina</i> | Cham. | 25 | 3 | 19 | common | 0.313131 | partially protected |
| <i>Randia hebecarpa</i> | Benth. | 2 | 3 | 1 | form6 | 0 | unprotected |
| <i>Randia obcordata</i> | S.Watson | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Raputia brevipedunculata</i> | Kallunki | 2 | 1 | 3 | form5 | 1 | protected |
| <i>Rauvolfia mattfeldiana</i> | Markgr. | 4 | 1 | 2 | form1 | 0.5 | partially protected |
| <i>Remijia amazonica</i> | K.Schum. | 12 | 3 | 9 | common | 0.25 | partially protected |
| <i>Remijia densiflora</i> | Benth. | 4 | 3 | 4 | common | 0.666667 | partially protected |
| <i>Remijia ferruginea</i> | (A.St.-Hil.) DC. | 5 | 3 | 3 | common | 0.214286 | partially protected |
| <i>Remijia firmula</i> | (Mart.) Wedd. | 19 | 3 | 17 | common | 0.384615 | partially protected |
| <i>Remijia roraimae</i> | (Benth.) K.Schum. | 5 | 3 | 3 | common | 0.75 | partially protected |
| <i>Retiniphyllum kuhlmannii</i> | Standl. | 8 | 3 | 5 | common | 0.102564 | partially protected |
| <i>Retiniphyllum laxiflorum</i> | (Benth.) N.E.Br. | 5 | 3 | 5 | common | 0.666667 | partially protected |
| <i>Retiniphyllum schomburgkii</i> | (Benth.) MÃƒÂ¼ll.Arg. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Rhamnidium elaeocarpum</i> | Reissek | 19 | 3 | 13 | common | 0.105386 | partially protected |
| <i>Rhynchanthera cordata</i> | DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Rhynchanthera grandiflora</i> | (Aubl.) DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Richeria grandis</i> | Vahl | 22 | 3 | 23 | common | 0.333333 | partially protected |
| <i>Rinorea guianensis</i> | Aubl. | 19 | 3 | 21 | common | 0.2 | partially protected |
| <i>Rinorea pubiflora</i> | (Benth.) Sprague & Sandwith | 17 | 3 | 17 | common | 0.166667 | partially protected |
| <i>Rinorea riana</i> | Kuntze | 12 | 3 | 9 | common | 0.615385 | partially protected |
| <i>Rinoreocarpus ulei</i> | (Melch.) Ducke | 14 | 3 | 12 | common | 0.636364 | partially protected |
| <i>Rochefortia spinosa</i> | (Jacq.) Urb. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Rogersonanthus arboreus</i> | (Britton) Maguire & B.M.Boom | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Rosenbergiodendron densiflorum</i> | (K.Schum.) Fagerl. | 8 | 3 | 6 | common | 0.157895 | partially protected |
| <i>Rosenbergiodendron formosum</i> | (Jacq.) Fagerl. | 8 | 3 | 3 | common | 0.086957 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Roucheria schomburgkii</i> | Planch. | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Roupala dielsii</i> | J.F. Macbr. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Roupala meisneri</i> | Sleumer | 2 | 1 | 2 | form5 | 0.4 | partially protected |
| <i>Roupala montana</i> | Aubl. | 35 | 3 | 32 | common | 0.163007 | partially protected |
| <i>Rourea discolor</i> | Baker | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Rourea doniana</i> | Baker | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Rourea gardneriana</i> | Planch. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Rourea induta</i> | Planch. | 21 | 3 | 13 | common | 0.1258 | partially protected |
| <i>Rourea martiana</i> | Baker | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Rudgea burchelliana</i> | MÃƒÂ¼ll.Arg. | 4 | 3 | 1 | form2 | 0.2 | partially protected |
| <i>Rudgea cornifolia</i> | (Kunth) Standl. | 23 | 3 | 21 | common | 0.290323 | partially protected |
| <i>Rudgea crassiloba</i> | (Benth.) B.L.Rob. | 16 | 3 | 17 | common | 0.083333 | partially protected |
| <i>Rudgea eriloba</i> | Benth. | 12 | 3 | 8 | common | 0.105263 | partially protected |
| <i>Rudgea hostmanniana</i> | Benth. | 3 | 1 | 3 | form5 | 1 | protected |
| <i>Rudgea palicoureoides</i> | (Mart.) MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Rudgea viburnoides</i> | (Cham.) Benth. | 24 | 3 | 18 | common | 0.103203 | partially protected |
| <i>Ruizterania retusa</i> | (Spruce ex Warm.) Marc.-Berti | 18 | 3 | 15 | common | 0.166667 | partially protected |
| <i>Ruprechtia brachysepala</i> | Meisn. | 16 | 3 | 12 | common | 0.133333 | partially protected |
| <i>Ruprechtia exploratricis</i> | Sandwith | 5 | 3 | 5 | common | 0.333333 | partially protected |
| <i>Ryania canescens</i> | Eichler | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Ryania dentata</i> | (Kunth) Miq. | 5 | 3 | 5 | common | 0.047619 | partially protected |
| <i>Ryania riedeliana</i> | Eichler | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Ryania speciosa</i> | Vahl | 23 | 3 | 20 | common | 0.176471 | partially protected |
| <i>Sacoglottis guianensis</i> | Benth. | 26 | 3 | 29 | common | 0.34375 | partially protected |
| <i>Sacoglottis mattogrossensis</i> | Malme | 21 | 3 | 21 | common | 0.391304 | partially protected |
| <i>Salacia crassifolia</i> | (Mart. ex Schult.) G. Don | 20 | 3 | 13 | common | 0.140523 | partially protected |
| <i>Salacia elliptica</i> | (Mart.) G.Don | 16 | 3 | 7 | common | 0.2 | partially protected |
| <i>Salacia grandifolia</i> | (Mart.) G. Don | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Salvertia convallariodora</i> | A. St.-Hil. | 25 | 3 | 20 | common | 0.133047 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Salzmannia nitida</i> | DC. | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Samanea inopinata</i> | (Harms)Barneby & J.W.Grimes | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Samanea tubulosa</i> | (Benth.)Barneby & J.W.Grimes | 9 | 3 | 7 | common | 0.181818 | partially protected |
| <i>Sambucus australis</i> | Cham. & Schldl. | 7 | 3 | 8 | common | 0.555556 | partially protected |
| <i>Sapindus saponaria</i> | L. | 24 | 3 | 17 | common | 0.166667 | partially protected |
| <i>Sapium argutum</i> | (MÃƒÂ¼ll.Arg.) Huber | 5 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Sapium glandulosum</i> | (L.) Morong | 32 | 3 | 29 | common | 0.25 | partially protected |
| <i>Sapium haematospermum</i> | MÃƒÂ¼ll.Arg. | 10 | 3 | 5 | common | 0.045455 | partially protected |
| <i>Sapium obovatum</i> | Klotzsch ex MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Sapium pallidum</i> | (MÃƒÂ¼ll.Arg.) Huber | 4 | 3 | 3 | common | 0 | unprotected |
| <i>Sapium sellowianum</i> | (MÃƒÂ¼ll.Arg.) Klotzsch ex Baill. | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Sarcaulus brasiliensis</i> | (A.DC.) Eyma | 19 | 3 | 17 | common | 0.222222 | partially protected |
| <i>Savia sessiliflora</i> | (Sw.) Willd. | 3 | 3 | 4 | form4 | 0.4 | partially protected |
| <i>Schefflera burchellii</i> | (Seem.) Frodin & Fiaschi | 14 | 3 | 5 | common | 0.117117 | partially protected |
| <i>Schefflera coriacea</i> | (Marchal ex Thurn) Harms | 2 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Schefflera distractiflora</i> | (Harms) Frodin | 16 | 3 | 9 | common | 0.229167 | partially protected |
| <i>Schefflera macrocarpa</i> | (Cham. & Schldl.) Frodin | 13 | 3 | 11 | common | 0.123994 | partially protected |
| <i>Schefflera malmei</i> | (Harms) Frodin | 13 | 3 | 5 | common | 0.086957 | partially protected |
| <i>Schefflera morototoni</i> | (Aubl.) Maguire, Steyermark & Frodin | 13 | 3 | 5 | common | 0 | unprotected |
| <i>Schefflera selloi</i> | (Marchal) Frodin & Fiaschi | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Schefflera sessiliflora</i> | Frodin | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Schefflera varisiana</i> | Frodin | 7 | 3 | 6 | common | 0.347826 | partially protected |
| <i>Schefflera villosissima</i> | Fiaschi & Pirani | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Schefflera vinosa</i> | (Cham. & Schldl.) Frodin & Fiaschi | 20 | 3 | 13 | common | 0.136674 | partially protected |
| <i>Schinopsis brasiliensis</i> | Engl. | 9 | 3 | 10 | common | 0.212766 | partially protected |
| <i>Schinus terebinthifolius</i> | Raddi | 19 | 3 | 12 | common | 0.121622 | partially protected |
| <i>Schizocalyx cuspidatus</i> | (A.St.-Hil.) Kainul. & B.Bremer | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Schoepfia brasiliensis</i> | A.DC. | 14 | 3 | 9 | common | 0.448276 | partially protected |
| <i>Schoepfia lucida</i> | Pulle | 6 | 3 | 6 | common | 0.3 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Sebastiania brasiliensis</i> | Spreng. | 16 | 3 | 10 | common | 0.169811 | partially protected |
| <i>Sebastiania brevifolia</i> | (MÃƒÂ¢ll.Arg.) MÃƒÂ¢ll.Arg. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Sebastiania commersoniana</i> | (Baill.) L.B.Sm. & Downs | 6 | 3 | 2 | common | 0.214286 | partially protected |
| <i>Seguieria americana</i> | L. | 11 | 3 | 10 | common | 0.285714 | partially protected |
| <i>Seguieria macrophylla</i> | Benth. | 12 | 3 | 11 | common | 0.357143 | partially protected |
| <i>Seguieria paraguayensis</i> | Morong | 5 | 3 | 3 | common | 0 | unprotected |
| <i>Senaea coerulea</i> | Taub. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Senegalia bahiensis</i> | (Benth.) Seigler & Ebinger | 8 | 3 | 6 | common | 0 | unprotected |
| <i>Senegalia bonariensis</i> | (Gillies ex Hook. & Arn.) Seigler & Ebinger | 3 | 1 | 2 | form5 | 1 | protected |
| <i>Senegalia globosa</i> | (Bocage & Miotto) L.P. Queiroz | 3 | 1 | 3 | form5 | 0.166667 | partially protected |
| <i>Senegalia langsdorffii</i> | (Benth.) Seigler & Ebinger | 11 | 3 | 4 | common | 0.285714 | partially protected |
| <i>Senegalia lewisi</i> | (Bocage & Miotto) L.P.Queiroz | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Senegalia loretensis</i> | (J.F. Macbr.) Seigler & Ebinger | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Senegalia piauhensis</i> | (Benth.) Seigler & Ebinger | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Senegalia polyphylla</i> | (DC.) Britton | 15 | 3 | 8 | common | 0.076923 | partially protected |
| <i>Senegalia riparia</i> | (Kunth) Britton | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Senegalia tamarindifolia</i> | (L.) Britton & Rose | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Senegalia tenuifolia</i> | (L.) Britton & Rose | 26 | 3 | 24 | common | 0.241758 | partially protected |
| <i>Senna acuruensis</i> | (Benth.) H.S.Irwin & Barneby | 3 | 3 | 2 | form4 | 0.125 | partially protected |
| <i>Senna affinis</i> | (Benth.) H.S.Irwin & Barneby | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Senna alata</i> | (L.) Roxb. | 26 | 3 | 19 | common | 0.246154 | partially protected |
| <i>Senna appendiculata</i> | (Vogel) Wiersema | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Senna atomaria</i> | (L.) H.S.Irwin & Barneby | 4 | 3 | 1 | form2 | 0.166667 | partially protected |
| <i>Senna bacillaris</i> | (L.f.) H.S.Irwin & Barneby | 14 | 3 | 12 | common | 0.235294 | partially protected |
| <i>Senna biglandularis</i> | A.O. Araujo & V.C. Souza | 7 | 3 | 2 | common | 0.157895 | partially protected |
| <i>Senna cearensis</i> | Afr.Fern. | 2 | 3 | 2 | form4 | 0.333333 | partially protected |
| <i>Senna corifolia</i> | (Benth.) H.S.Irwin & Barneby | 4 | 3 | 4 | common | 0.315789 | partially protected |
| <i>Senna gardneri</i> | (Benth.) H.S.Irwin & Barneby | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Senna georgica</i> | H.S.Irwin & Barneby | 15 | 3 | 13 | common | 0.307692 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Senna lechriosperma</i> | H.S.Irwin & Barneby | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Senna macranthera</i> | (Collad.)H.S.Irwin & Barneby | 13 | 3 | 9 | common | 0.153846 | partially protected |
| <i>Senna macrophylla</i> | (Kunth)H.S.Irwin & Barneby | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Senna multijuga</i> | (Rich.) H.S.Irwin & Barneby | 29 | 3 | 24 | common | 0.298077 | partially protected |
| <i>Senna oblongifolia</i> | (Vogel)H.S.Irwin & Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Senna quinquangulata</i> | (Rich.)H.S.Irwin & Barneby | 6 | 3 | 8 | common | 0.3 | partially protected |
| <i>Senna reniformis</i> | (G.Don)H.S.Irwin & Barneby | 7 | 3 | 4 | common | 0.259259 | partially protected |
| <i>Senna reticulata</i> | (Willd.)H.S.Irwin & Barneby | 7 | 3 | 8 | common | 0.166667 | partially protected |
| <i>Senna rizzinii</i> | H.S.Irwin & Barneby | 4 | 3 | 3 | common | 0.294118 | partially protected |
| <i>Senna robiniifolia</i> | (Benth.)H.S.Irwin & Barneby | 2 | 3 | 2 | form4 | 0.25 | partially protected |
| <i>Senna rostrata</i> | (Mart.)H.S.Irwin & Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Senna rugosa</i> | (G.Don)H.S.Irwin & Barneby | 24 | 3 | 16 | common | 0.125402 | partially protected |
| <i>Senna silvestris</i> | (Vell.) H.S.Irwin & Barneby | 29 | 3 | 32 | common | 0.236152 | partially protected |
| <i>Senna skinneri</i> | (Benth.)H.S.Irwin & Barneby | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Senna spectabilis</i> | (DC.)H.S.Irwin & Barneby | 11 | 3 | 9 | common | 0.057143 | partially protected |
| <i>Senna trachypus</i> | (Benth.)H.S.Irwin & Barneby | 4 | 3 | 3 | common | 0.444444 | partially protected |
| <i>Senna velutina</i> | (Vogel)H.S.Irwin & Barneby | 19 | 3 | 9 | common | 0.164706 | partially protected |
| <i>Senna viciifolia</i> | (Benth.)H.S.Irwin & Barneby | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Senna wurdackii</i> | H.S.Irwin & Barneby | 3 | 3 | 1 | form6 | 0.2 | partially protected |
| <i>Sida ulei</i> | Ulbr. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Sida viarum</i> | A.St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Sideroxylon obtusifolium</i> | (Roem. & Schult.) T.D.Penn. | 11 | 3 | 8 | common | 0.125 | partially protected |
| <i>Sigmatanthus trifoliatus</i> | Huber ex Emmerich | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Simaba ferruginea</i> | A. St.-Hil. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Simaba floribunda</i> | A. St.-Hil. | 4 | 3 | 3 | common | 0.25 | partially protected |
| <i>Simaba insignis</i> | A. St.-Hil. & Tul. | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Simaba maiana</i> | Casar. | 13 | 3 | 5 | common | 0.136364 | partially protected |
| <i>Simaba orinocensis</i> | Kunth | 17 | 3 | 16 | common | 0.26087 | partially protected |
| <i>Simaba polyphylla</i> | (Cavalcante) W.W. Thomas | 1 | 1 | 1 | form7 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|----------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Simaba subcymosa</i> | A. St.-Hil. & Tul. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Simarouba amara</i> | Aubl. | 25 | 3 | 27 | common | 0.343511 | partially protected |
| <i>Simarouba versicolor</i> | A. St.-Hil. | 24 | 3 | 21 | common | 0.158898 | partially protected |
| <i>Simira corumbensis</i> | (Standl.) Steyerm. | 14 | 3 | 7 | common | 0.102941 | partially protected |
| <i>Simira hexandra</i> | (S.Moore) Steyerm. | 5 | 3 | 3 | common | 0.111111 | partially protected |
| <i>Simira rubescens</i> | (Benth.) Bremek. ex Steyerm. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Siparuna brasiliensis</i> | (Spreng.) A. DC. | 13 | 3 | 4 | common | 0.125 | partially protected |
| <i>Siparuna guianensis</i> | Aubl. | 31 | 3 | 38 | common | 0.185381 | partially protected |
| <i>Siphoneugena crassifolia</i> | (DC.) ProenÃƒÂ§a & Sobral | 1 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Siphoneugena densiflora</i> | O.Berg | 9 | 3 | 6 | common | 0.162011 | partially protected |
| <i>Siphoneugena dussii</i> | (Krug & Urb.) ProenÃƒÂ§a | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Sloanea eichleri</i> | K.Schum. | 21 | 3 | 21 | common | 0.266667 | partially protected |
| <i>Sloanea retusa</i> | Uittien | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Sloanea terniflora</i> | (Moc. & SessÃƒÂ© ex DC.) Standl. | 3 | 1 | 4 | form5 | 0.6 | partially protected |
| <i>Socratea exorrhiza</i> | (Mart.) H.Wendl. | 1 | 3 | 1 | form6 | 0.381818 | partially protected |
| <i>Solanum asperum</i> | Rich. | 22 | 3 | 12 | common | 0.159091 | partially protected |
| <i>Solanum bicolor</i> | Willd. ex Roem. & Schult. | 5 | 3 | 2 | common | 0.055556 | partially protected |
| <i>Solanum caavurana</i> | Vell. | 4 | 3 | 3 | common | 0.444444 | partially protected |
| <i>Solanum campaniforme</i> | Roem. & Schult. | 11 | 3 | 12 | common | 0.875 | partially protected |
| <i>Solanum conglobatum</i> | Dunal | 4 | 3 | 3 | common | 0.428571 | partially protected |
| <i>Solanum crinitum</i> | Lam. | 25 | 3 | 20 | common | 0.247706 | partially protected |
| <i>Solanum itatiaiae</i> | Glaz. ex Edmonds | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Solanum lacerdae</i> | DusÃƒÂ©n | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Solanum lycocarpum</i> | A. St.-Hil. | 19 | 3 | 12 | common | 0.091803 | partially protected |
| <i>Solanum mauritianum</i> | Scop. | 4 | 3 | 1 | form2 | 0.25 | partially protected |
| <i>Solanum proteanthum</i> | Bohs | 17 | 3 | 12 | common | 0.238095 | partially protected |
| <i>Solanum pseudoquina</i> | A. St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Solanum rhytidioandrum</i> | Sendtn. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Solanum rufescens</i> | Sendtn. | 1 | 1 | 1 | form7 | 0.5 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|--------------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Solanum scuticum</i> | M. Nee | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Solanum stipulaceum</i> | Roem. & Schult. | 4 | 3 | 2 | common | 0.375 | partially protected |
| <i>Solanum swartzianum</i> | Roem. & Schult. | 3 | 3 | 5 | form4 | 0.571429 | partially protected |
| <i>Solanum velleum</i> | Roem. & Schult. | 3 | 3 | 2 | form4 | 0.444444 | partially protected |
| <i>Sorocea bonplandii</i> | (Baill.) W.C.Burger, Lanj. & de Boer | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Sorocea duckei</i> | W.C. Burger | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Sorocea guilleminiana</i> | Gaudich. | 24 | 3 | 21 | common | 0.24 | partially protected |
| <i>Sorocea hilarii</i> | Gaudich. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Sorocea sprucei</i> | (Baill.) J.F.Macbr. | 14 | 3 | 11 | common | 0.116279 | partially protected |
| <i>Sphinctanthes microphyllus</i> | K.Schum. | 5 | 1 | 3 | form1 | 0.5 | partially protected |
| <i>Sphinga platyloba</i> | (DC.)Barneby & J.W.Grimes | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Spiranthera odoratissima</i> | A. St.-Hil. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Spiranthera parviflora</i> | Sandwith | 5 | 1 | 8 | form1 | 1 | protected |
| <i>Spondias mombin</i> | L. | 30 | 3 | 24 | common | 0.284672 | partially protected |
| <i>Spondias tuberosa</i> | Arruda | 5 | 3 | 4 | common | 0.4 | partially protected |
| <i>Staminodianthus racemosus</i> | (Hoehne) D.B.O.S.Cardoso & H.C.Lima | 18 | 3 | 15 | common | 0.222222 | partially protected |
| <i>Stenopadus talaumifolius</i> | S.F.Blake | 4 | 1 | 2 | form1 | 1 | protected |
| <i>Sterculia striata</i> | A. St.-Hil. & Naudin | 19 | 3 | 11 | common | 0.107843 | partially protected |
| <i>Stigmaphyllon paralias</i> | A.Juss. | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Stillingia argutedentata</i> | Jabl. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Stillingia bodenbenderi</i> | (Kuntze) D.J.Rogers | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Stillingia oppositifolia</i> | Baill. ex MÃƒÂ¼ll.Arg. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Stillingia saxatilis</i> | MÃƒÂ¼ll.Arg. | 8 | 3 | 5 | common | 0.35 | partially protected |
| <i>Stillingia trapezoidea</i> | Ule | 3 | 1 | 2 | form5 | 0.2 | partially protected |
| <i>Stillingia uleana</i> | Pax & K.Hoffm. | 8 | 3 | 5 | common | 0.333333 | partially protected |
| <i>Strychnos pseudoquina</i> | A. St.-Hil. | 18 | 3 | 10 | common | 0.106195 | partially protected |
| <i>Strychnos rubiginosa</i> | A.DC. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Stryphnodendron adstringens</i> | (Mart.)Coville | 21 | 3 | 13 | common | 0.13089 | partially protected |
| <i>Stryphnodendron coriaceum</i> | Benth. | 14 | 3 | 6 | common | 0.083333 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|-----------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Stryphnodendron fissuratum</i> | Martins | 2 | 1 | 3 | form5 | 0.333333 | partially protected |
| <i>Stryphnodendron guianense</i> | (Aubl.) Benth. | 18 | 3 | 23 | common | 0.172414 | partially protected |
| <i>Stryphnodendron polyphyllum</i> | Mart. | 15 | 3 | 5 | common | 0.072581 | partially protected |
| <i>Stryphnodendron pulcherrimum</i> | (Willd.) Hochr. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Stryphnodendron rotundifolium</i> | Mart. | 22 | 3 | 18 | common | 0.136087 | partially protected |
| <i>Stylogyne orinocensis</i> | (Kunth) Mez | 14 | 3 | 10 | common | 0.2 | partially protected |
| <i>Stylogyne serpentina</i> | Mez | 6 | 1 | 5 | form1 | 0.666667 | partially protected |
| <i>Stylosanthes capitata</i> | Vogel | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Stylosanthes guianensis</i> | (Aubl.) Sw. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Styrax aureus</i> | Mart. | 4 | 3 | 3 | common | 0.388889 | partially protected |
| <i>Styrax camporum</i> | Pohl | 19 | 3 | 10 | common | 0.146293 | partially protected |
| <i>Styrax ferrugineus</i> | Nees & Mart. | 20 | 3 | 13 | common | 0.143048 | partially protected |
| <i>Styrax glaber</i> | Sw. | 3 | 1 | 3 | form5 | 0 | unprotected |
| <i>Styrax griseus</i> | P.W. Fritsch | 11 | 3 | 4 | common | 0.193548 | partially protected |
| <i>Styrax maninul</i> | B.Walln. | 4 | 3 | 3 | common | 0.344828 | partially protected |
| <i>Styrax martii</i> | Seub. | 5 | 3 | 3 | common | 0.242424 | partially protected |
| <i>Styrax pallidus</i> | A.DC. | 4 | 3 | 3 | common | 0.666667 | partially protected |
| <i>Styrax pauciflorus</i> | A. DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Styrax pedicellatus</i> | (Perkins) B.Walln. | 7 | 3 | 5 | common | 0.333333 | partially protected |
| <i>Styrax rotundatus</i> | (Perkins) P.W.Fritsch | 8 | 3 | 5 | common | 0.357143 | partially protected |
| <i>Swartzia acuminata</i> | Willd. ex Vogel | 8 | 3 | 14 | common | 0.5 | partially protected |
| <i>Swartzia acutifolia</i> | Vogel | 4 | 3 | 2 | common | 0.5 | partially protected |
| <i>Swartzia apetala</i> | Raddi | 14 | 3 | 11 | common | 0.162162 | partially protected |
| <i>Swartzia aptera</i> | DC. | 4 | 1 | 4 | form1 | 0 | unprotected |
| <i>Swartzia aymardii</i> | Barneby | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Swartzia bahiensis</i> | R.S.Cowan | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Swartzia brachyrachis</i> | Harms | 15 | 3 | 18 | common | 0.888889 | partially protected |
| <i>Swartzia capixabensis</i> | Mansano | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Swartzia dipetala</i> | Vogel | 3 | 3 | 1 | form6 | 0.142857 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|-----------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Swartzia flaemingii</i> | Raddi | 19 | 3 | 14 | common | 0.227273 | partially protected |
| <i>Swartzia grandifolia</i> | Benth. | 2 | 3 | 2 | form4 | 0.571429 | partially protected |
| <i>Swartzia jorori</i> | Harms | 11 | 3 | 8 | common | 0.173913 | partially protected |
| <i>Swartzia latifolia</i> | Benth. | 11 | 3 | 8 | common | 0.35 | partially protected |
| <i>Swartzia laurifolia</i> | Benth. | 18 | 3 | 20 | common | 0.419355 | partially protected |
| <i>Swartzia laxiflora</i> | Benth. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Swartzia macrostachya</i> | Benth. | 9 | 3 | 8 | common | 0.111111 | partially protected |
| <i>Swartzia parvifolia</i> | Schery | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Swartzia rondoniensis</i> | Torke & Mansano | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Swartzia xanthopetala</i> | Sandwith | 2 | 3 | 3 | form4 | 0.2 | partially protected |
| <i>Sweetia fruticosa</i> | Spreng. | 17 | 3 | 11 | common | 0.057471 | partially protected |
| <i>Syagrus cocoides</i> | Mart. | 13 | 3 | 12 | common | 0.222222 | partially protected |
| <i>Syagrus comosa</i> | (Mart.) Mart. | 19 | 3 | 12 | common | 0.190955 | partially protected |
| <i>Syagrus deflexa</i> | Noblick & Lorenzi | 2 | 1 | 1 | form7 | 0.6 | partially protected |
| <i>Syagrus flexuosa</i> | (Mart.) Becc. | 21 | 3 | 12 | common | 0.142367 | partially protected |
| <i>Syagrus glaucescens</i> | Glaz. ex Becc. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Syagrus oleracea</i> | (Mart.) Becc. | 15 | 3 | 8 | common | 0.063492 | partially protected |
| <i>Syagrus romanzoffiana</i> | (Cham.) Glassman | 6 | 3 | 3 | common | 0.25 | partially protected |
| <i>Syagrus sancona</i> | (Kunth) H.Karst. | 5 | 3 | 5 | common | 0.25 | partially protected |
| <i>Syagrus schizophylla</i> | (Mart.) Glassman | 3 | 3 | 4 | form4 | 0.333333 | partially protected |
| <i>Symmeria paniculata</i> | Benth. | 22 | 3 | 21 | common | 0.263158 | partially protected |
| <i>Symphonia globulifera</i> | L.f. | 24 | 3 | 25 | common | 0.352941 | partially protected |
| <i>Sympphyopappus itatiayensis</i> | (Hieron.) R.M.King & H.Rob. | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Sympphyopappus lymansmithii</i> | B.L.Rob. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Symplocos angulata</i> | Brand | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Symplocos arbutifolia</i> | Casar. | 3 | 3 | 3 | form4 | 0.3 | partially protected |
| <i>Symplocos celastrinea</i> | Mart. ex Miq. | 11 | 3 | 4 | common | 0.258065 | partially protected |
| <i>Symplocos glaberrima</i> | Gontsch. | 3 | 1 | 2 | form5 | 0.166667 | partially protected |
| <i>Symplocos guianensis</i> | (Aubl.) GÃƒÂ¶rké | 7 | 3 | 6 | common | 0.555556 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|--------------------------------------|---|-----|------------|-----------|--------|----------|---------------------|
| <i>Symplocos microstyla</i> | Aranha, P.W.Fritsch & Almeda | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Symplocos nitens</i> | (Pohl) Benth. | 17 | 3 | 12 | common | 0.114286 | partially protected |
| <i>Symplocos oblongifolia</i> | Casar. | 12 | 3 | 7 | common | 0.205128 | partially protected |
| <i>Symplocos platyphylla</i> | (Pohl) Benth. | 6 | 3 | 1 | form2 | 0.125 | partially protected |
| <i>Symplocos pubescens</i> | Klotzsch ex Benth. | 12 | 3 | 7 | common | 0.258065 | partially protected |
| <i>Symplocos pycnobotrya</i> | Mart. ex Miq. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Symplocos pycnophylla</i> | Sleumer | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Symplocos rhamnifolia</i> | A. DC. | 4 | 3 | 2 | common | 0.333333 | partially protected |
| <i>Symplocos tenuifolia</i> | Brand | 4 | 3 | 3 | common | 0.285714 | partially protected |
| <i>Symplocos tetrandra</i> | Mart. | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Syzygium jambos</i> | (L.) Alston | 5 | 3 | 1 | form2 | 0.3 | partially protected |
| <i>Tabebuia aurea</i> | (Silva Manso) Benth. & Hook.f. ex S.Moore | 29 | 3 | 27 | common | 0.149688 | partially protected |
| <i>Tabebuia cassinoides</i> | (Lam.) DC. | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Tabebuia elliptica</i> | (DC.) Sandwith | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Tabebuia insignis</i> | (Miq.) Sandwith | 28 | 3 | 21 | common | 0.180952 | partially protected |
| <i>Tabebuia nodosa</i> | (Griseb.) Griseb. | 6 | 3 | 4 | common | 0.125 | partially protected |
| <i>Tabebuia pilosa</i> | A.H.Gentry | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Tabebuia roseoalba</i> | (Ridl.) Sandwith | 20 | 3 | 15 | common | 0.116564 | partially protected |
| <i>Tabebuia stenocalyx</i> | Sprague & Stapf | 2 | 3 | 1 | form6 | 0.333333 | partially protected |
| <i>Tabernaemontana catharinensis</i> | A.DC. | 8 | 3 | 7 | common | 0.421053 | partially protected |
| <i>Tabernaemontana flavicans</i> | Willd. ex Roem. & Schult. | 18 | 3 | 19 | common | 0.388889 | partially protected |
| <i>Tabernaemontana hystrix</i> | Steud. | 3 | 3 | 3 | form4 | 0.125 | partially protected |
| <i>Tabernaemontana laeta</i> | Mart. | 3 | 3 | 2 | form4 | 0.5 | partially protected |
| <i>Tabernaemontana siphilitica</i> | (L.f.) Leeuwenb. | 27 | 3 | 22 | common | 0.166667 | partially protected |
| <i>Tachigali aurea</i> | Tul. | 21 | 3 | 13 | common | 0.134935 | partially protected |
| <i>Tachigali pilgeriana</i> | (Harms) Oliveira-Filho | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Tachigali rubiginosa</i> | (Mart. ex Tul.) Oliveira-Filho | 16 | 3 | 10 | common | 0.264368 | partially protected |
| <i>Tachigali subvelutina</i> | (Benth.) Oliveira-Filho | 18 | 3 | 13 | common | 0.140065 | partially protected |
| <i>Tachigali vulgaris</i> | L.F. Gomes da Silva & H.C. Lima | 22 | 3 | 15 | common | 0.159091 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Talisia esculenta</i> | (A. St.-Hil.) Radlk. | 21 | 3 | 22 | common | 0.106796 | partially protected |
| <i>Talisia hexaphylla</i> | Vahl | 11 | 1 | 7 | form1 | 0.25 | partially protected |
| <i>Talisia subalbens</i> | (Mart.) Radlk. | 2 | 3 | 1 | form6 | 0.5 | partially protected |
| <i>Tapirira guianensis</i> | Aubl. | 32 | 3 | 42 | common | 0.249042 | partially protected |
| <i>Tapirira obtusa</i> | (Benth.) J.D.Mitch. | 23 | 3 | 20 | common | 0.233766 | partially protected |
| <i>Tapura amazonica</i> | Poepp. | 10 | 3 | 4 | common | 0.291667 | partially protected |
| <i>Taralea cordata</i> | Ducke | 14 | 3 | 17 | common | 0.454545 | partially protected |
| <i>Taralea crassifolia</i> | (Benth.) Ducke | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Taralea reticulata</i> | (Benth.) Ducke | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Terminalia actinophylla</i> | Mart. | 14 | 3 | 5 | common | 0.235294 | partially protected |
| <i>Terminalia argentea</i> | Mart. | 20 | 3 | 13 | common | 0.115385 | partially protected |
| <i>Terminalia eichleriana</i> | Alwan & Stace | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Terminalia fagifolia</i> | Mart. | 21 | 3 | 13 | common | 0.144486 | partially protected |
| <i>Terminalia glabrescens</i> | Mart. | 21 | 3 | 11 | common | 0.12628 | partially protected |
| <i>Terminalia lucida</i> | Hoffmanns. ex Mart. | 15 | 3 | 12 | common | 0.4 | partially protected |
| <i>Terminalia triflora</i> | (Griseb.) Lillo | 4 | 3 | 2 | common | 0.333333 | partially protected |
| <i>Ternstroemia brasiliensis</i> | Cambess. | 6 | 3 | 4 | common | 0.5 | partially protected |
| <i>Ternstroemia candolleana</i> | Wawra | 14 | 3 | 10 | common | 0.44 | partially protected |
| <i>Ternstroemia carnosa</i> | Cambess. | 6 | 3 | 5 | common | 0.285714 | partially protected |
| <i>Ternstroemia dentata</i> | (Aubl.) Sw. | 20 | 3 | 22 | common | 0.230769 | partially protected |
| <i>Ternstroemia punctata</i> | (Aubl.) Sw. | 16 | 3 | 14 | common | 0.7 | partially protected |
| <i>Tessaria integrifolia</i> | Ruiz & Pav. | 11 | 3 | 11 | common | 0.083333 | partially protected |
| <i>Tetragastris altissima</i> | (Aubl.) Swart | 22 | 3 | 21 | common | 0.253521 | partially protected |
| <i>Thespesia populnea</i> | (L.) Sol. ex CorrÃƒÂ³a | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Thiloa glaucocarpa</i> | (Mart.) Eichler | 14 | 3 | 10 | common | 0.171429 | partially protected |
| <i>Thyrsodium spruceanum</i> | Benth. | 18 | 3 | 20 | common | 0.428571 | partially protected |
| <i>Tibouchina macrochiton</i> | Cogn. | 2 | 3 | 1 | form6 | 0.4 | partially protected |
| <i>Tibouchina papyrus</i> | (Pohl) Toledo | 5 | 3 | 2 | common | 0.206897 | partially protected |
| <i>Tibouchina rigidula</i> | (Naudin) Wurdack | 2 | 1 | 2 | form5 | 0 | unprotected |

| S | author | lat | population | ecoregion | Form | r | status |
|------------------------------------|------------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Tibouchina robusta</i> | Cogn. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Tibouchina sellowiana</i> | Cogn. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Tibouchina stenocarpa</i> | (DC.) Cogn. | 21 | 3 | 14 | common | 0.180685 | partially protected |
| <i>Tibouchina trichopoda</i> | Baill. | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Tococa guianensis</i> | Aubl. | 27 | 3 | 23 | common | 0.292308 | partially protected |
| <i>Tocoyena brasiliensis</i> | Mart. | 21 | 3 | 15 | common | 0.295775 | partially protected |
| <i>Tocoyena bullata</i> | (Vell.) Mart. | 10 | 3 | 8 | common | 0.259259 | partially protected |
| <i>Tocoyena formosa</i> | (Cham. & Schleidl.) K.Schum. | 30 | 3 | 28 | common | 0.157788 | partially protected |
| <i>Tocoyena hispidula</i> | Standl. | 6 | 3 | 7 | common | 0.296296 | partially protected |
| <i>Tocoyena longiflora</i> | Aubl. | 4 | 3 | 1 | form2 | 0.5 | partially protected |
| <i>Tocoyena neglecta</i> | N.E.Br. | 2 | 3 | 3 | form4 | 0.428571 | partially protected |
| <i>Tocoyena sellowiana</i> | (Cham. & Schleidl.) K.Schum. | 3 | 1 | 2 | form5 | 0.125 | partially protected |
| <i>Toulia crassifolia</i> | Radlk. | 7 | 3 | 3 | common | 0.114286 | partially protected |
| <i>Tournefortia ruficunda</i> | Salzm. ex DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Trema micrantha</i> | (L.) Blume | 28 | 3 | 34 | common | 0.190184 | partially protected |
| <i>Trembleya laniflora</i> | (D. Don) Cogn. | 3 | 3 | 3 | form4 | 0.466667 | partially protected |
| <i>Trembleya parviflora</i> | (D. Don) Cogn. | 14 | 3 | 8 | common | 0.193548 | partially protected |
| <i>Trichilia casaretti</i> | C. DC. | 2 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Trichilia clausenii</i> | C. DC. | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Trichilia elegans</i> | A.Juss. | 26 | 3 | 16 | common | 0.161616 | partially protected |
| <i>Trichilia pallida</i> | Sw. | 26 | 3 | 21 | common | 0.191667 | partially protected |
| <i>Trichilia stellatotomentosa</i> | Kuntze | 11 | 3 | 5 | common | 0.190476 | partially protected |
| <i>Triplaris americana</i> | L. | 22 | 3 | 21 | common | 0.188679 | partially protected |
| <i>Triplaris caracasana</i> | Cham. | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Triplaris gardneriana</i> | Wedd. | 2 | 1 | 1 | form7 | 0 | unprotected |
| <i>Trischidium molle</i> | (Benth.) H.E. Ireland | 2 | 3 | 2 | form4 | 0 | unprotected |
| <i>Unonopsis aurantiaca</i> | Maas & Westra | 1 | 1 | 1 | form7 | 0.5 | partially protected |
| <i>Unonopsis bahiensis</i> | Maas & Orava | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Unonopsis floribunda</i> | Diels | 16 | 3 | 14 | common | 0.294118 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|---------------------------------|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Unonopsis guatterioides</i> | (A.DC.) R.E.Fr. | 27 | 3 | 26 | common | 0.226563 | partially protected |
| <i>Vachellia farnesiana</i> | (L.) Wight & Arn. | 18 | 3 | 14 | common | 0.173333 | partially protected |
| <i>Vachellia macracantha</i> | (Humb. & Bonpl. ex Willd.) Seigler & Ebinger | 4 | 3 | 2 | common | 0.130435 | partially protected |
| <i>Vantanea bahiaeensis</i> | Cuatrec. | 2 | 3 | 1 | form6 | 0.4 | partially protected |
| <i>Vantanea compacta</i> | (Schnizl.) Cuatrec. | 5 | 3 | 4 | common | 0.26087 | partially protected |
| <i>Vantanea minor</i> | Benth. | 3 | 3 | 2 | form4 | 0.666667 | partially protected |
| <i>Vantanea morii</i> | Cuatrec. | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Vantanea obovata</i> | (Nees & Mart.) Benth. | 9 | 3 | 4 | common | 0.36 | partially protected |
| <i>Vasconcellea glandulosa</i> | A. DC. | 1 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Vasconcellea microcarpa</i> | (Jacq.) A.DC. | 14 | 3 | 11 | common | 0.111111 | partially protected |
| <i>Vatairea fusca</i> | (Ducke) Ducke | 1 | 3 | 1 | form6 | 0 | unprotected |
| <i>Vatairea macrocarpa</i> | (Benth.) Ducke | 26 | 3 | 20 | common | 0.142857 | partially protected |
| <i>Vellozia gigantea</i> | N.L.Menezes & Mello-Silva | 2 | 1 | 2 | form5 | 0.666667 | partially protected |
| <i>Vellozia squamata</i> | Pohl | 14 | 3 | 3 | common | 0.284211 | partially protected |
| <i>Vellozia tubiflora</i> | (A.Rich.) Kunth | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Verbesina glabrata</i> | Hook. & Arn. | 3 | 3 | 4 | form4 | 0.444444 | partially protected |
| <i>Verbesina luetzelburgii</i> | Mattf. | 3 | 1 | 2 | form5 | 0.25 | partially protected |
| <i>Vernonanthura discolor</i> | H.Rob. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Vernonanthura divaricata</i> | (Spreng.) H.Rob. | 4 | 3 | 1 | form2 | 0.363636 | partially protected |
| <i>Vernonanthura fagifolia</i> | (Gardner) H.Rob. | 3 | 1 | 3 | form5 | 0.285714 | partially protected |
| <i>Vernonia brasiliiana</i> | (L.) Druce | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Virola sebifera</i> | Aubl. | 33 | 3 | 35 | common | 0.171895 | partially protected |
| <i>Virola subsessilis</i> | (Benth.) Warb. | 12 | 3 | 3 | common | 0.086957 | partially protected |
| <i>Vismia cayennensis</i> | (Jacq.) Pers. | 8 | 3 | 7 | common | 0.153846 | partially protected |
| <i>Vismia gracilis</i> | Hieron. | 22 | 3 | 15 | common | 0.321101 | partially protected |
| <i>Vismia guianensis</i> | (Aubl.) Pers. | 30 | 3 | 32 | common | 0.261538 | partially protected |
| <i>Vismia japurensis</i> | Rchb.f. | 4 | 3 | 2 | common | 0 | unprotected |
| <i>Vismia martiana</i> | Reichardt | 5 | 3 | 4 | common | 0.75 | partially protected |
| <i>Vismia micrantha</i> | Mart. ex A. St.-Hil. | 3 | 3 | 3 | form4 | 0.466667 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------|---------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Vismia minutiflora</i> | Ewan | 13 | 3 | 10 | common | 0.153846 | partially protected |
| <i>Vismia pentagyna</i> | (Spreng.) Ewan | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Vismia schultesii</i> | N.Robson | 12 | 3 | 11 | common | 0.375 | partially protected |
| <i>Vismia tenuinervia</i> | (M.E.Berg) N.Robson | 7 | 3 | 6 | common | 0.4 | partially protected |
| <i>Vitex cymosa</i> | Bertero ex Spreng. | 10 | 3 | 6 | common | 0.166667 | partially protected |
| <i>Vitex flavens</i> | Kunth | 7 | 3 | 7 | common | 0.538462 | partially protected |
| <i>Vitex gardneriana</i> | Schauer | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Vitex megapotamica</i> | (Spreng.) Moldenke | 6 | 3 | 2 | common | 0.142857 | partially protected |
| <i>Vitex orinocensis</i> | Kunth | 15 | 3 | 17 | common | 0.1875 | partially protected |
| <i>Vitex pashiniana</i> | Moldenke | 11 | 3 | 9 | common | 0.291667 | partially protected |
| <i>Vitex polygama</i> | Cham. | 24 | 3 | 19 | common | 0.185096 | partially protected |
| <i>Vitex pseudolea</i> | Rusby | 7 | 3 | 3 | common | 0.375 | partially protected |
| <i>Vitex rufescens</i> | A.Juss. | 6 | 3 | 6 | common | 0.4375 | partially protected |
| <i>Vitex schaueriana</i> | Moldenke | 5 | 3 | 3 | common | 0.428571 | partially protected |
| <i>Vitex sellowiana</i> | Cham. | 1 | 1 | 3 | form5 | 0.5 | partially protected |
| <i>Vitex triflora</i> | Vahl | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Vochysia cinnamomea</i> | Pohl | 16 | 3 | 8 | common | 0.057554 | partially protected |
| <i>Vochysia discolor</i> | Warm. | 2 | 1 | 3 | form5 | 0.285714 | partially protected |
| <i>Vochysia divergens</i> | Pohl | 11 | 3 | 7 | common | 0.235294 | partially protected |
| <i>Vochysia elliptica</i> | Mart. | 14 | 3 | 6 | common | 0.142857 | partially protected |
| <i>Vochysia emarginata</i> | Vahl | 9 | 3 | 5 | common | 0.310345 | partially protected |
| <i>Vochysia ferruginea</i> | Mart. | 24 | 3 | 26 | common | 0.21875 | partially protected |
| <i>Vochysia gardneri</i> | Warm. | 16 | 3 | 7 | common | 0.176849 | partially protected |
| <i>Vochysia glaberrima</i> | Warm. | 2 | 1 | 2 | form5 | 1 | protected |
| <i>Vochysia haenkeana</i> | Mart. | 16 | 3 | 12 | common | 0.280255 | partially protected |
| <i>Vochysia petraea</i> | Warm. | 1 | 1 | 1 | form7 | 0.4 | partially protected |
| <i>Vochysia pruinosa</i> | Pohl | 10 | 3 | 3 | common | 0.170732 | partially protected |
| <i>Vochysia rufa</i> | Mart. | 17 | 3 | 14 | common | 0.139808 | partially protected |
| <i>Vochysia thyrsoides</i> | Pohl | 16 | 3 | 11 | common | 0.190594 | partially protected |

| S | author | lat | population | ecoregion | Form | r | status |
|----------------------------------|-----------------------------|-----|------------|-----------|--------|----------|---------------------|
| <i>Vochysia tucanorum</i> | Mart. | 22 | 3 | 17 | common | 0.158333 | partially protected |
| <i>Vochysia venezuelana</i> | Stafleu | 5 | 3 | 4 | common | 0.129032 | partially protected |
| <i>Weinmannia discolor</i> | Gardner | 2 | 1 | 2 | form5 | 0 | unprotected |
| <i>Weinmannia guyanensis</i> | Klotzsch ex Engl. | 1 | 3 | 1 | form6 | 1 | protected |
| <i>Weinmannia humilis</i> | Engl. | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Weinmannia paulliniifolia</i> | Pohl | 2 | 3 | 3 | form4 | 0.2 | partially protected |
| <i>Wunderlichia azulensis</i> | Maguire & G.M.Barroso | 1 | 1 | 1 | form7 | 1 | protected |
| <i>Wunderlichia bahiensis</i> | Maguire & G.M.Barroso | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Wunderlichia cruelesiana</i> | Taub. | 9 | 3 | 6 | common | 0.24 | partially protected |
| <i>Wunderlichia mirabilis</i> | Riedel ex Baker | 10 | 3 | 6 | common | 0.257732 | partially protected |
| <i>Ximenia americana</i> | L. | 21 | 3 | 19 | common | 0.183099 | partially protected |
| <i>Ximenia coriacea</i> | Engl. | 3 | 1 | 2 | form5 | 0.166667 | partially protected |
| <i>Ximenia intermedia</i> | (Chodat & Hassl.) DeFilipps | 16 | 3 | 8 | common | 0.083333 | partially protected |
| <i>Xylopia aromatica</i> | (Lam.) Mart. | 33 | 3 | 39 | common | 0.146377 | partially protected |
| <i>Xylopia frutescens</i> | Aubl. | 19 | 3 | 17 | common | 0.37037 | partially protected |
| <i>Xylopia laevigata</i> | (Mart.) R.E. Fr. | 2 | 1 | 1 | form7 | 0.666667 | partially protected |
| <i>Xylopia sericea</i> | A.St.-Hil. | 21 | 3 | 19 | common | 0.209738 | partially protected |
| <i>Xylosma benthamii</i> | (Tul.) Triana & Planch. | 26 | 3 | 26 | common | 0.111111 | partially protected |
| <i>Xylosma venosa</i> | N.E.Br. | 3 | 1 | 1 | form7 | 0 | unprotected |
| <i>Zanthoxylum acuminatum</i> | (Sw.) Sw. | 11 | 3 | 8 | common | 0.25 | partially protected |
| <i>Zanthoxylum caribaeum</i> | Lam. | 23 | 3 | 19 | common | 0.271186 | partially protected |
| <i>Zanthoxylum fagara</i> | (L.) Sarg. | 15 | 3 | 7 | common | 0.125 | partially protected |
| <i>Zanthoxylum hamadryadicum</i> | Pirani | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Zanthoxylum rhoifolium</i> | Lam. | 29 | 3 | 31 | common | 0.137815 | partially protected |
| <i>Zanthoxylum riedelianum</i> | Engl. | 22 | 3 | 19 | common | 0.110778 | partially protected |
| <i>Zanthoxylum rigidum</i> | Humb. & Bonpl. ex Willd. | 9 | 3 | 5 | common | 0.08 | partially protected |
| <i>Zanthoxylum stelligerum</i> | Turcz. | 6 | 3 | 3 | common | 0.222222 | partially protected |
| <i>Zanthoxylum syncarpum</i> | Tul. | 4 | 3 | 4 | common | 0.294118 | partially protected |
| <i>Zeyheria montana</i> | Mart. | 20 | 3 | 15 | common | 0.127451 | partially protected |

| S | author | | lat | population | ecoregion | Form | r | status |
|-----------------------------|--|--|-----|------------|-----------|--------|----------|---------------------|
| <i>Zeyheria tuberculosa</i> | (Vell.) Bureau ex Verl. | | 7 | 3 | 4 | common | 0 | unprotected |
| <i>Ziziphus cotinifolia</i> | Reissek | | 1 | 1 | 1 | form7 | 0 | unprotected |
| <i>Ziziphus joazeiro</i> | Mart. | | 14 | 3 | 10 | common | 0.121212 | partially protected |
| <i>Zollernia glabra</i> | (Spreng.)Yakovlev | | 2 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Zollernia ilicifolia</i> | (Brongn.)Vogel | | 2 | 1 | 2 | form5 | 0.333333 | partially protected |
| <i>Zygia cataractae</i> | (Kunth)L.Rico | | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Zygia claviflora</i> | (Spruce ex Benth.) Barneby & J.W. Grimes | | 3 | 3 | 1 | form6 | 0.125 | partially protected |
| <i>Zygia coccinea</i> | (G.Don)L.Rico | | 2 | 1 | 2 | form5 | 0.5 | partially protected |
| <i>Zygia inaequalis</i> | (Willd.)Pittier | | 1 | 1 | 1 | form7 | 0.333333 | partially protected |
| <i>Zygia latifolia</i> | (L.)Fawc. & Rendle | | 5 | 3 | 5 | common | 0.333333 | partially protected |

Table 4. Proportion of protected species in each form of rarity.

| Status of protected | common | form1 | form2 | form3 | form4 | form5 | form6 | form7 | total/status of protected |
|-------------------------------------|---------------|--------------|--------------|--------------|--------------|--------------|--------------|--------------|----------------------------------|
| unprotected | 45 | 17 | 2 | | 27 | 53 | 39 | 165 | 348 |
| partially protected | 1,254 | 20 | 24 | 1 | 95 | 123 | 59 | 74 | 1,650 |
| protected | 6 | 8 | | | 6 | 21 | 29 | 135 | 205 |
| Total species/form of rarity | 1,305 | 45 | 26 | 1 | 128 | 197 | 127 | 374 | 2,203 |

Table 5. The ratio (r) between the occurrence into PA and total occurrences for each form and common species. The form of rarity (Forms), the number of species per form (S), the number of all records per form (n), the r minimum (rMin) maximum (rMax) median (rMD) and semi-amplitude of the 95% confidence interval (SCI).

| Forms | S | n | rMin | rMax | rMD | SCI |
|--------------|----------|----------|-------------|-------------|------------|------------|
| Common | 1,254 | 177,858 | 0.02 | 0.88 | 0.14 | 0.0003 |
| Form1 | 20 | 256 | 0.12 | 0.80 | 0.13 | 0.023 |
| Form2 | 24 | 336 | 0.07 | 0.6 | 0.20 | 0.013 |
| Form3 | 1 | 10 | 0.6 | 0.6 | 0.6 | 0 |
| Form4 | 95 | 1,027 | 0.07 | 0.72 | 0.37 | 0.009 |
| From5 | 123 | 684 | 0.12 | 0.80 | 0.38 | 0.012 |
| Form6 | 59 | 482 | 0.10 | 0.66 | 0.38 | 0.01 |
| Form7 | 74 | 282 | 0.20 | 0.66 | 0.50 | 0.01 |

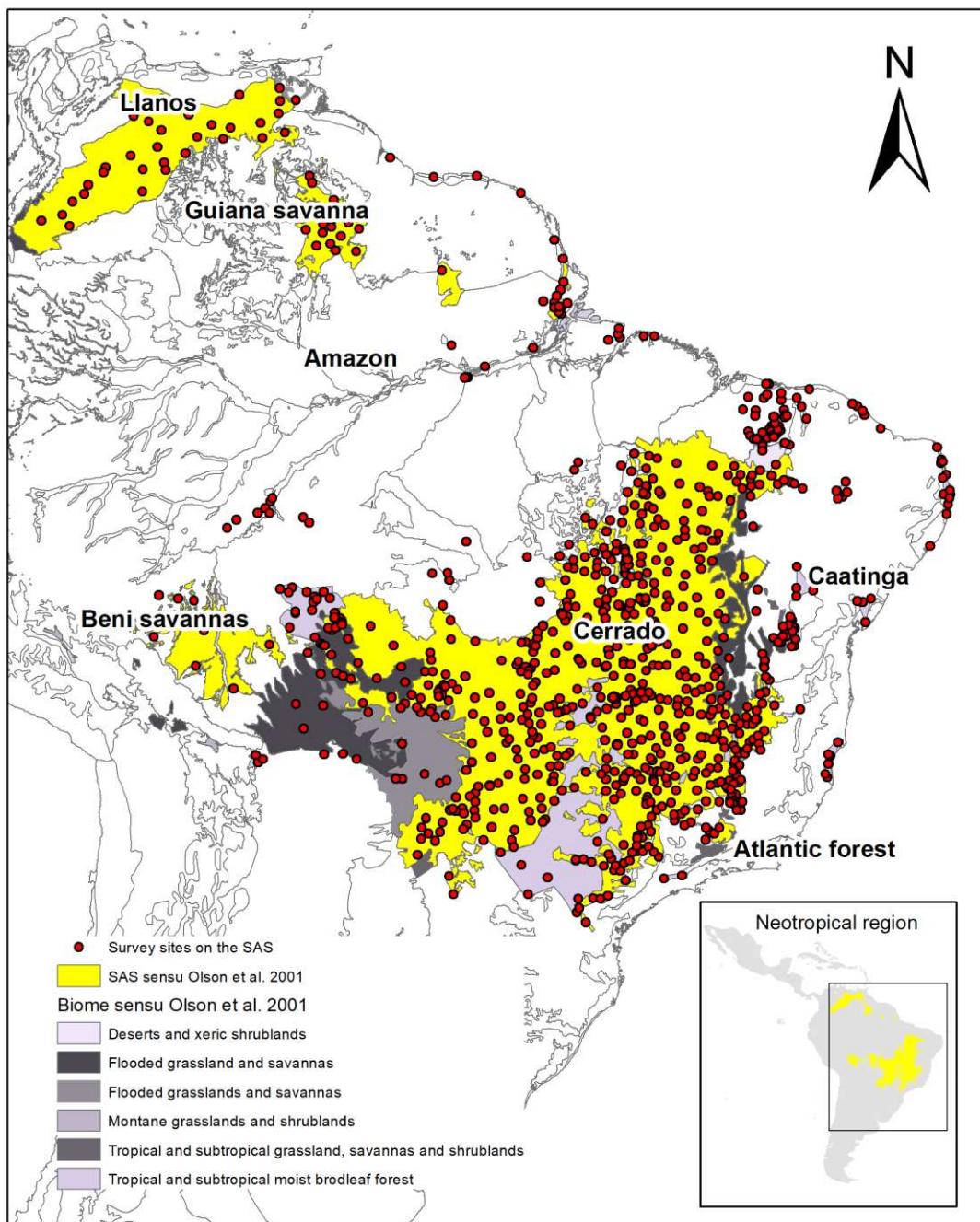


Fig. 1. Main ecoregions of the South American savannas (SAS). In yellow, the restrict concept of Olson et al. (2001). The red points indicate the localisation of the surveys (n=949).

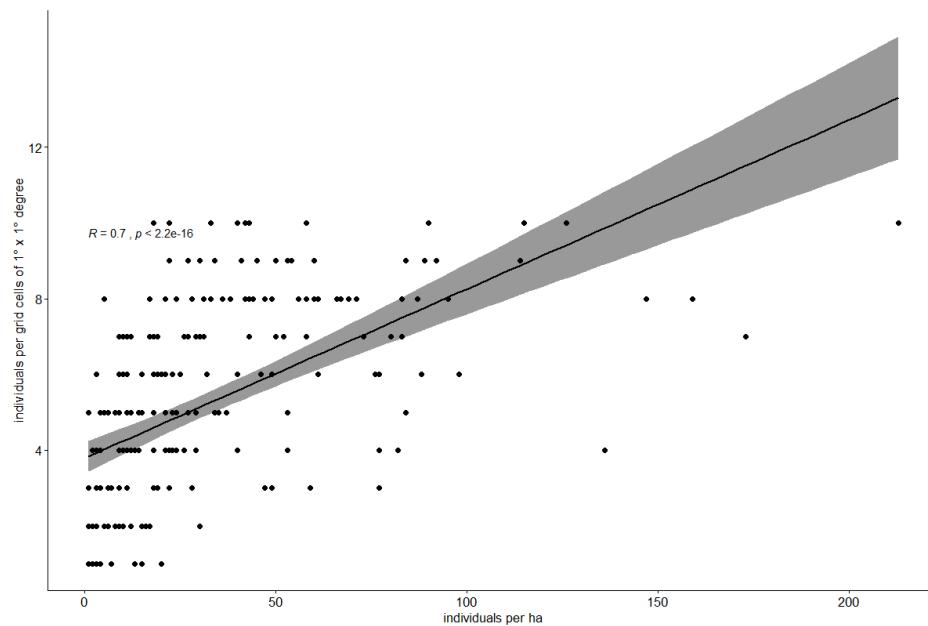


Fig. 2. Scatter plot showing the correlation between the number of individuals per site and grid cells. On the x and y axis are both number of individuals per species. Because for South America savannas the phytosociological surveys are scarce, we used Brazilian Cerrado as a base area. A site corresponds to an area with a minimum size of one hectare with DBH $\geq 4.7\text{cm}$ as a minimum sample size made in Cerrado. The grid area comprises grid cells by 1×1 degree for all Cerrado as in the Maciel and Martins (2019). The line shows the fit of the spearman model and the gray band the 95% confidence interval. The spearman coefficient (R) and the p-value for 0.05 (p) are displayed.

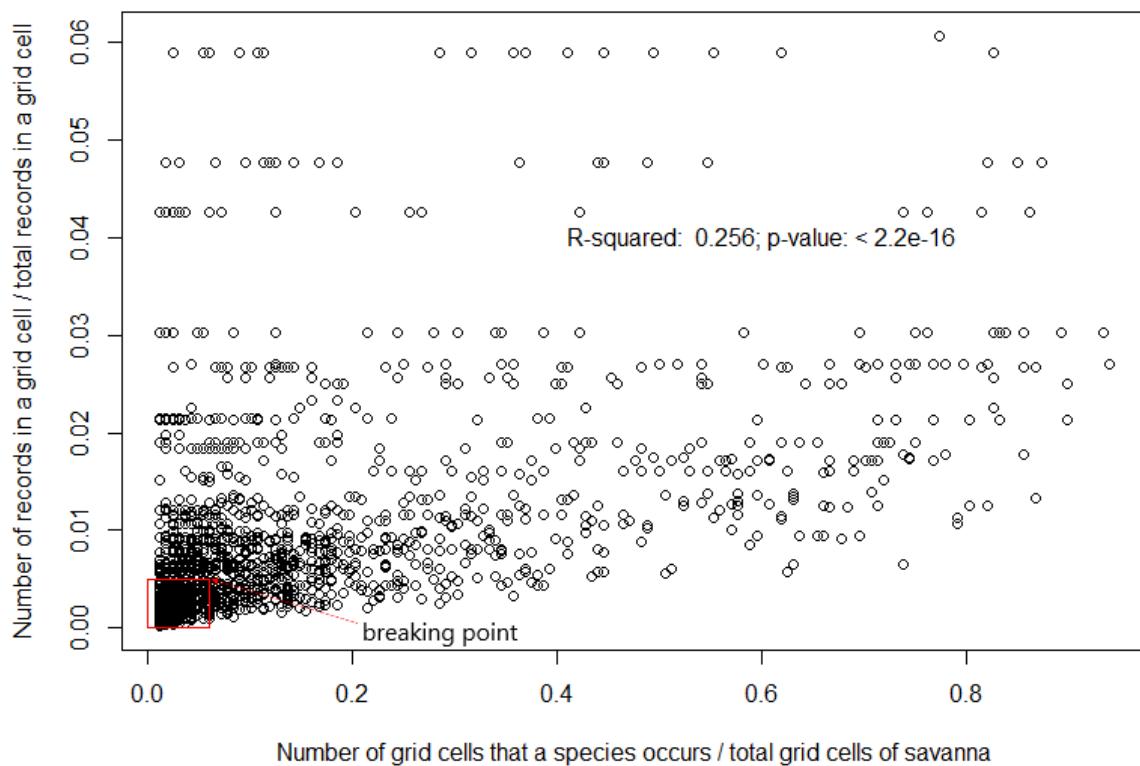


Fig. 3. Scatter plot with the breaking point (indicated by red arrow). Each circle indicates a species. The species within the red square were considered scarce population.

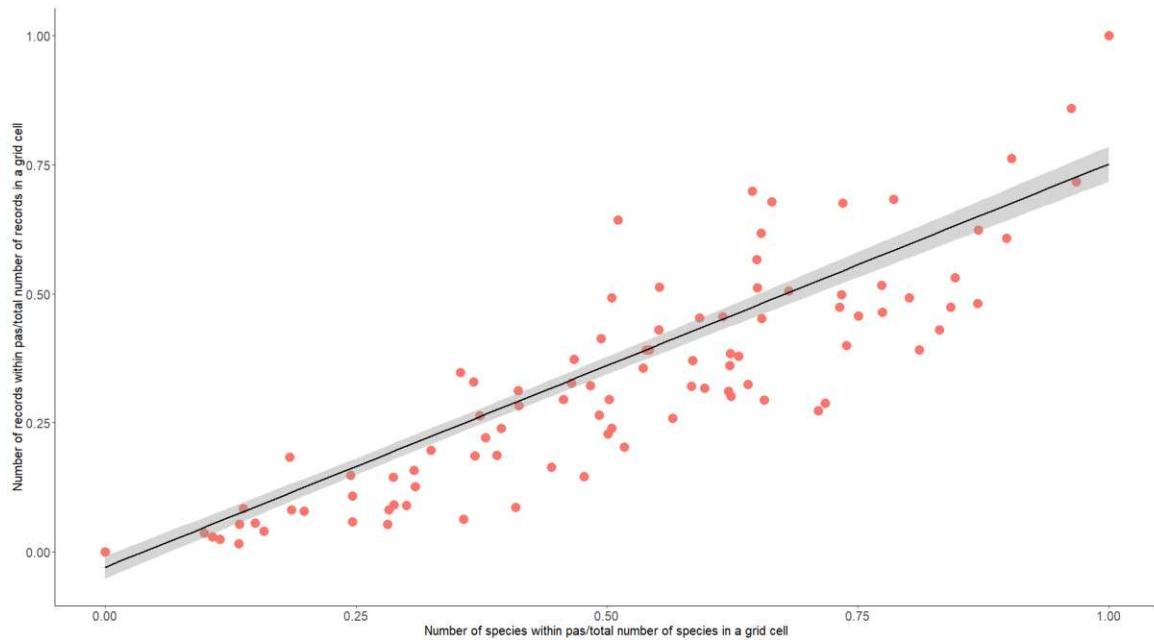


Fig. 4. Scatterplot with a straight line of the best fit generalized linear model. The x-axis is the ratio between the number of records within the protected areas and all records in each grid cell of 1° of latitude by 1° of longitude. The y-axis is the ratio between the number of species within PAs and the total species per grid cells (A).

Supplementary material chapter 4

Supplementary information

Model the dynamics of the vegetation in FORMIND (details could be found in Fischer et al. 2016)

FORMIND calculates biomass using the following general equation:

$$B=4.D2.H.f.$$

where the calculation represents the volume of the tree stem (according to its geometry) multiplied by three factors, which describe the biomass content more concisely. First, f [-] denotes a type-specific form factor, which accounts for deviations of the stem from a cylindrical shape. Second, the parameter ρ [tODM/m³] represents wood density, which describes how much organic dry matter per unit of volume the stem contains. Third, division by the parameter σ [tODM/tODM], which represents the fraction of the total aboveground biomass attributed to the stem, results in the total aboveground biomass (B).

Table 1. Reproduction capacity (RP) and seeders (SE), phonologic leaf with deciduously (DC) or evergreen (EV), animal (ZO), and no animal dispersal (NZ). More details for plant functional types parametrization can be also find in Dantas et al. 2018.

| Parameter | Description | Unit | PFT1 | PFT2 | PFT3 | PFT4 | PFT5 | PFT6 | PFT7 | PFT8 | Reference |
|-------------------|---|---|------|------|------|------|------|------|------|-----------------------|-----------------------------------|
| Reproduction | | SE | SE | SE | SE | RP | RP | RP | RP | RP | |
| Phenology | | DC | DC | EV | EV | DC | DC | EV | EV | EV | |
| Dispersal spectra | | NZ | ZO | NZ | ZO | NZ | ZO | NZ | ZO | (Hughes et al., 1994) | |
| TimeEnd | Total simulation time | Years | 1000 | | | | | | | | |
| K | Light extinction coefficient | $\text{m}^2 \text{ ground}$ $\text{m}^{-2} \text{ leaf}$ | 0.6 | | | | | | | | |
| Io | average irradiance above canopy | Mmol (photons) $\text{m}^{-2} \text{ s}^{-1}$ | 700 | | | | | | | | |
| I_{seed} | Minimum light intensity (relative to full light above canopy) required for establishment of seeds | % of I_0 | 0.7 | | | | | | | | |
| Hmax | Maximum size (height) of 10 a tree | m | 7 | 7 | 8 | 6.5 | 8 | 7 | 6.5 | 7.5 | derived from inventory data |

| Parameter | Description | Unit | PFT1 | PFT2 | PFT3 | PFT4 | PFT5 | PFT6 | PFT7 | PFT8 | Reference |
|--|----------------------------------|--|--------|------|------|------|------|------|------|------|--|
| | function of a single tree | | | | | | | | | | |
| A | Slope of light response curve | $\mu\text{mol } (\text{CO}_2) \cdot \text{mol}$ (photons) $^{-1}$ | 0.6 | 0.6 | 0.6 | 0.6 | 0.6 | 0.6 | 0.6 | 0.6 | (Franco et al., 2005). |
| A | predator efficiency | - | 0.1 | | | | | | | | (Dantas et al., 2018) |
| Th | predator handling time | Years | 0.0001 | | | | | | | | |
| Fire tolerance = probability for death (1: no tolerance 4: tolerant against fire) (group-specific) | | | 2 | 2 | 2 | 2 | 3 | 3 | 3 | 3 | (da Silva Rios and Sousa-Silva, 2017) (Ojeda et al., |

| Parameter | Description | Unit | PFT1 | PFT2 | PFT3 | PFT4 | PFT5 | PFT6 | PFT7 | PFT8 | Reference |
|--------------------|--|------|------|------|------|------|------|------|------|------|-----------------------------|
| Specific leaf mass | (SLM, g m ⁻²) | | 77 | 77 | 176 | 176 | 77 | 77 | 176 | 176 | Prado and De Moraes, (1997) |
| Leaf mass bases | [P_{Nmax} μmol (CO ₂)kg ⁻¹ s ⁻¹] | | 117 | 117 | 52 | 52 | 117 | 117 | 52 | 52 | Prado and De Moraes, (1997) |

Table 2. Inventories data for three areas. Reproduction capacity (RP) and seeders (SE), phonologic leaf with deciduously (DC) or evergreen (EV), animal (ZO), and no animal dispersal (NZ).

| Site | PFT | Reproduction | Leaf phenology | Dispersal mode | No of species | Stems (ha ⁻¹) | Max. height | Basal area (m ² ha ⁻¹) | Max. increment of stem (mm years ⁻¹) |
|------|-----|--------------|----------------|----------------|---------------|---------------------------|-------------|---|--|
| WS1 | 1 | SE | DC | NZ | 15 | 196 | 7 | 1.554 | 2.757 |
| | 2 | SE | DC | ZO | 9 | 163 | 7 | 1.31 | 2.547 |
| | 3 | SE | EV | NZ | 6 | 16 | 8 | 0.122 | 0.740 |
| | 4 | SE | EV | NZ | 15 | 240 | 6.5 | 1.724 | 2.070 |
| | 5 | RP | DC | ZO | 9 | 447 | 8 | 3.038 | 2.604 |
| | 6 | RP | DC | NZ | 17 | 372 | 7 | 2.428 | 1.877 |
| | 7 | RP | EV | ZO | 2 | 177 | 5 | 0.833 | 1.700 |
| | 8 | RP | EV | NZ | 9 | 164 | 7.5 | 1.182 | 1.220 |
| WS2 | 1 | SE | DC | ZO | 16 | 240 | 9 | 2.164 | 3.175 |
| | 2 | SE | DC | NZ | 20 | 274 | 9 | 4.815 | 3.632 |
| | 3 | SE | EV | ZO | 1 | 10 | 5 | 0.127 | 0.537 |
| | 4 | SE | EV | NZ | 9 | 69 | 8.5 | 0.709 | 3.527 |
| | 5 | RP | DC | NZ | 6 | 105 | 7.5 | 1.335 | 2.373 |
| | 6 | RP | DC | ZO | 11 | 71 | 6.5 | 0.711 | 1.248 |
| | 7 | RP | EV | NZ | 2 | 16 | 6 | 0.135 | 0.957 |
| | 8 | RP | EV | ZO | 5 | 68 | 8 | 0.575 | 0.650 |
| WS3 | 1 | SE | DC | NZ | 16 | 115 | 12 | 1.801 | 1.336 |
| | 2 | SE | DC | ZO | 18 | 354 | 12 | 6.981 | 4.269 |
| | 3 | SE | EV | NZ | 3 | 35 | 10 | 1.099 | 3.825 |
| | 4 | SE | EV | ZO | 10 | 92 | 11 | 1.196 | 4.442 |
| | 5 | RP | DC | NZ | 7 | 244 | 14 | 2.744 | 1.601 |
| | 6 | RP | DC | ZO | 19 | 339 | 9 | 3.067 | 2.534 |
| | 7 | RP | EV | NZ | 3 | 82 | 8.5 | 0.712 | 1.112 |
| | 8 | RP | EV | ZO | 8 | 130 | 8 | 1.287 | 1.671 |

Table 3. Aboveground biomass (ABV) for three phytophysiognomy of Cerrado, open savanna (OS), wood savanna (WS) and Savanna Woodlands (SW). Control scenario (C), climate change scenario (CC), climate change + defaunation (CC .DE), climate change + fire change scenario (CC.CF), and synergism (CC.CF.DE).

| Phytophysiognomy | ABV | Reference |
|-------------------------|------------|---------------------------------|
| OS | 8.03 | (Barbosa and Fearnside, 2005) |
| | 8.62 | (Kauffman et al., 1994) |
| WS | 23.66 | |
| | 28.2 | (Fearnside et al., 2009) |
| | 25.2 | (Felfili et al., 2004) |
| | 12.4 | (Felfili et al., 2004) |
| | 15.3 | (Felfili et al., 2004) |
| | 16.1 | (Felfili et al., 2004) |
| | 8.9 | (Felfili et al., 2004) |
| | 17.2 | (Felfili et al., 2002) |
| | 55.6 | (Torello-Raventos et al., 2013) |
| | 34.1 | (Torello-Raventos et al., 2013) |
| SW | 51.21 | (Fearnside et al., 2009) |
| | 76.47 | (Teodoro, 2014) |

Table 4. Parameters used in each model. Control scenario (C), climate change scenario (CC), climate change + defaunation (CC .DE), climate change + change in the fire (CC.CF), and synergism (CC.DS.CF).

| Scenario | Climate variable | Fire variable | Defaunation variable | | |
|------------------|-------------------------|---------------|----------------------------|---------------------------------|--|
| Change parameter | Annual temperature (°C) | Dry season | Fire frequency in 10 years | Seed predator attack efficiency | Survival rate of seeds – post disperse |
| C | Mean 25 | 6 | 2-3 | 10% | 10% |
| CC | +2 °C | 7 | 2-3 | 10% | 10% |
| CC.DS | +2 °C | 7 | 2-3 | 50% | 5% |
| CC.FC | +2 °C | 7 | 5 | 10% | 10% |
| CC.DS.FC | +2 °C | 7 | 5 | 50% | 5% |

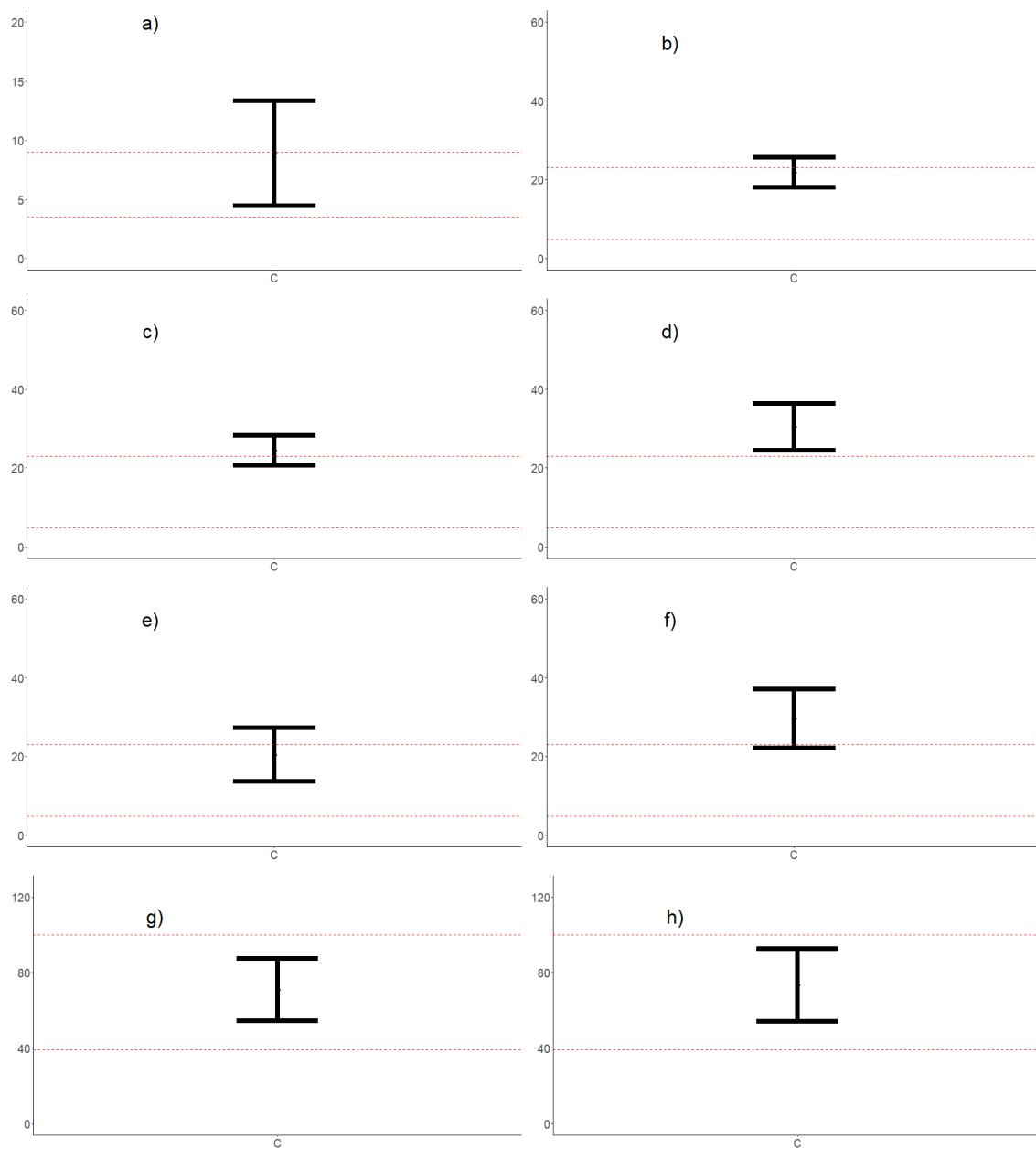


Fig. 1. Aboveground biomass (mean \pm SD) in eight Neotropical savanna communities in baseline scenario for open savanna (a), woody savanna (b, c, d, e, and f, respectively) and savanna woodland (g and h). Different areas are indicated by letters Red lines indicate the mean and minimum biomass recorded for each community in previous studies (references in Table S4).

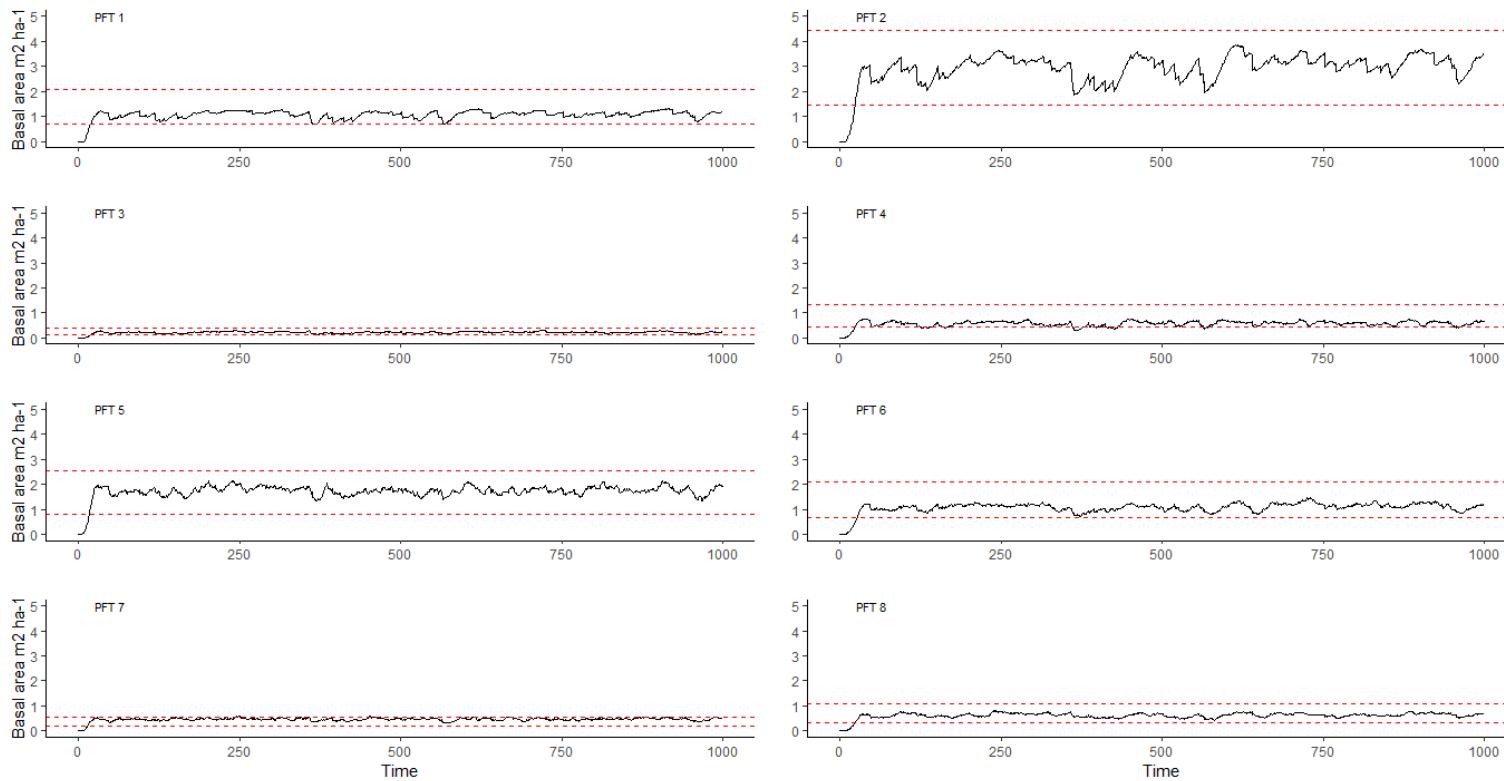


Fig. 2. The models validation for each plant functional type (PFT). The black line is a base area modeled in FORMIND and red line is a base area derived from data inventory. More details about model validation can be find in Dantas et al. 2018.

Anexo 1 Declaração de Bioética



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DECLARAÇÃO

Em observância ao §5º do Artigo 1º da Informação CCPG-UNICAMP/001/15, referente a Bioética e Biossegurança, declaro que o conteúdo de minha Tese de Doutorado, intitulada *LINKING PATTERNS AND PROCESSES IN TREE PLANT COMMUNITIES TO CONSERVATION AND MANAGEMENT OF THE SOUTH AMERICAN SAVANNAS*, desenvolvida no Programa de Pós-Graduação em Biociências e Tecnologia de Produtos Bioativos do Instituto de Biologia da Unicamp, não versa sobre pesquisa envolvendo seres humanos, animais ou temas afetos a Biossegurança.

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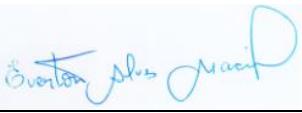
Nome do(a) orientador(a): Fernando Roberto Martins

Data: 18 de setembro de 2020

Anexo 2 Declaração dos Direitos Autorais**Declaração**

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