



universität
wien

MASTERARBEIT / MASTER'S THESIS

Titel der Masterarbeit / Title of the Master's Thesis

Iran's Protected area network insufficiently represents
climatic niches of endemic plants

verfasst von / submitted by

Simon Jean Ludovicy BSc

angestrebter akademischer Grad / in partial fulfilment of the requirements for the degree of
Master of Science (MSc)

Wien, 2021 / Vienna, 2021

Studienkennzahl lt. Studienblatt /
degree programme code as it appears on
the student record sheet:

UA 066 879

Studienrichtung lt. Studienblatt /
degree programme as it appears on
the student record sheet:

Masterstudium Naturschutz- und Biodiversitätsmanagement

Betreut von / Supervisor:

Univ.-Prof. Mag. Dr. Stefan Dullinger

Abstract

Protected Areas (PA) play a pivotal role in Conservation Biology. Anthropogenic climate change challenges the concept of static PAs as species may shift their distribution outside of current PA boundaries. Here, I compare the climatic niche of species with the climatic conditions that the species find within their associated PAs. Therefore, I use a newly introduced method that is particularly suitable when occurrence data of species is scarce. I focus on Iran, a centre of endemism in Southwest Asia. Based on the Iranian PA network, I assess the representation of the climatic niches of Iran's endemic plants in the national PA network under current and future climatic conditions. I find that niche representation of endemic species in PAs reaches only ca. 50% under the current climate, and that it will further decrease to 20 – 30% in the future, depending on the climate scenario used. Low and high elevation species show least climatic representation of climatic niches within PAs. In the future, endemic species with narrow ranges within Iran will lose much more climatic representation than endemics that are widespread within the country. These results suggest that the Iranian PA network is not fit for climate change. Efficient conservation planning must focus on increasing elevational gradients within PAs and on the identification of microrefugia that allow for long term species' persistence.

Keywords: climatic niche, protected areas, niche modelling, mountain ecology, climate change, endemic plants

Abstract (deutsch)

Schutzgebiete (SG) spielen eine zentrale Rolle in der Naturschutzbioologie. Der anthropogene Klimawandel fechtet das Konzept von statischen SG an, da Arten ihre Verbreitung aus den SG Grenzen hinaus verschieben. In dieser Studie vergleiche ich die klimatische Nische von Arten mit den klimatischen Bedingungen, die Arten in ihren umliegenden SG finden. Dafür benutze ich eine neu eingeführte Methode, die sich besonders eignet, wenn wenig Verbreitungsdaten von Arten vorliegen. Ich fokussiere mich auf den Iran, ein Zentrum für Endemismus in Südwest-Asien. Basierend auf dem iranischen SG-Netzwerk bewerte ich die klimatische Abdeckung der iranischen, endemischen Pflanzen im nationalen SG Netzwerk unter derzeitigen und zukünftigen klimatischen Bedingungen. Ich stelle fest, dass die Abdeckung der Nischen der Endemiten in SG nur 50% unter aktuellem Klima beträgt. In Zukunft sinkt diese Abdeckung, abhängig vom Szenario, auf 20-30%. Arten tiefer und hoher Lagen haben die geringste Repräsentierung der klimatischen Nische in SG. In Zukunft werden Arten mit eingeschränkter Verbreitung mehr klimatische Repräsentierung verlieren als weitverbreitete Arten. Diese Ergebnisse deuten darauf hin, dass das iranische SG-Netzwerk nicht fit ist für den Klimawandel. Effiziente Naturschutzplanung muss Höhengradienten innerhalb von SG erhöhen und Mikrorefugia identifizieren, die eine Persistenz der Arten über einen langen Zeitraum ermöglichen.

Schlagwörter: klimatische Nische, Schutzgebiete, Nischenmodellierung, Gebirgsökologie, Klimawandel, endemische Pflanzen

Table of Contents

Abstract.....	1
Abstract (deutsch).....	1
Einleitung	2
Manuskript.....	7
Introduction	7
Material and Methods	8
Study area	8
Species and Protected Area data	8
Climate data and scenarios.....	9
Climate niche construction	10
Climatic niche representation.....	11
Statistical analysis	11
Results.....	12
Discussion.....	15
Schlussfolgerungen	18
Acknowledgement	19
Literature	19
Appendix	27
List of tables	27
Table of figures	27

Einleitung

Der anthropogene Klimawandel sorgt für ein weltweites Ansteigen der Temperaturen auf der Erde (IPCC, 2019). Seit dem Beginn der Industriellen Revolution ist die Temperatur der Erdoberfläche kontinuierlich wärmer geworden (Shukla et al., 2019). Verantwortlich hierfür ist in erster Linie der Treibhauseffekt, der über die Menge an Treibhausgasen in der Atmosphäre gesteuert wird (Manabe, 2019). Die Treibhausgase verändern das Energiepotenzial in der Atmosphäre, indem sie die kurzwellige Strahlung der Sonne ungehindert auf die Erdoberfläche treffen lassen. Die daraufhin von Erde zurückgestrahlte langwellige Strahlung wird von den Treibhausgasen zurückgehalten und in alle Richtungen abgestrahlt. Natürlicherweise existieren mehrere Treibhausgase in der Atmosphäre, einen Großteil macht Wasserdampf H_2O aus. Andere bedeutende Treibhausegase sind Kohlendioxid (CO_2), Ozon (O_3), Lachgas (N_2O) und Methan (CH_4) (Tuckett, 2019). Durch die Verbrennung von Kohle und Gas hat der Mensch den CO_2 -Anteil in der Atmosphäre seit der Industrialisierung von 280 parts per million (ppm) auf über 400 ppm erhöht (IPCC, 2019).

Das Klima der Zukunft wird in Klimamodellen dargestellt, sogenannte GCMs (General Circulation Model) (Randall, 2000). Diese Modelle berechnen das zukünftige Klima mit Hilfe von Schlüsselkomponenten des Klimasystems, wie zum Beispiel der Atmosphäre oder den Ozeanen (z.B. Zhang et al., 2020). Wie sich die Konzentration der Treibhausgase in Zukunft verändert hängt auch von dem Handeln der Menschen heute und in Zukunft ab. Um diese Entwicklung abzuschätzen stellen Wissenschaftler in unterschiedlichen sozioökonomischen Szenarien dar, welche klimarelevante Entscheidungen in Zukunft getroffen werden könnten und welcher Treibhausgasausstoß damit verbunden ist (sogenannte Relative Concentration Pathways (RCP)(Moss et al., 2010)). Auf Basis dieser RCPs wird von verschiedenen wissenschaftlichen Institutionen mit Hilfe selbst-entwickelter GCMs das Klima der Zukunft berechnet (z.B. Séférian et al., 2019 Hajima et al., 2020). Der Weltklimarat IPCC (Intergovernmental Panel on Climate Change) fasst die Ergebnisse der verschiedenen Institutionen zusammen und bereitet sie für Entscheidungsträger auf (IPCC, 2019). Die verschiedenen Modelle stimmen darin überein, dass die Temperatur in den nächsten Dekaden ansteigen wird. Neben den steigenden Temperaturen verändern sich auch die Niederschlagsmuster (Kopala et al., 2020). Durch den höheren CO₂-Gehalt in der Atmosphäre steigt der pH-Wert der Ozeane (Doney et al., 2009). Im Eis gespeichertes Wasser an den Polen sowie in Gletschern wird ebenfalls rapide abschmelzen (Allison et al., 2009, Haeberli et al., 2007). Eine weitere wichtige Erkenntnis der Klimawandelforschung ist, dass extreme Wetterereignisse, wie Dürreperioden oder Starkniederschläge zunehmen werden (Sillmann & Roeckner, 2008).

Das Studium der Effekte des Klimawandels auf die Artenvielfalt ist ein wissenschaftliches Feld, dass in den letzten Dekaden stark an Bedeutung gewonnen hat (Bellard et al., 2012; Araujo & Rahbeck, 2006, Trew et al., 2021). Die Ergebnisse dieser Forschung zeigen, dass Klimawandel die Artenvielfalt auf unterschiedliche Art und Weise beeinflussen kann. Eine weitverbreitete und häufige Reaktion von Arten ist die Anpassung von Verbreitungsgebieten an die sich verändernde Klimageographie (Chen et al., 2011). Diese Änderungen in den Verbreitungsmustern sind gut dokumentiert (Parmesan et al., 2003). Allgemein gilt, dass Arten der Äquator-nahen Lagen dem Klimawandel folgen, indem sie sich in Richtung der Pole ausbreiten (Parmesan et al., 1999, Brommer et al., 2012). Arten der Gebirge wandern in höhere Lagen, um den zunehmend heißeren Temperaturen zu entkommen (Habel et al., 2019, Rumpf et al., 2019, Steinbauer et al., 2018). Es ändern sich jedoch nicht nur geografische Muster, sondern auch die Phänologie der Arten ist durch den Klimawandel beeinflusst (Menzel et al., 2006). Der Laubaustrieb von Pflanzen erfolgt früher im Jahr und der Laubabwurf später aufgrund der länger günstigen Bedingungen. Auch bei Tieren ist dieser Effekt sichtbar (Cohen et al., 2018). Vögel & Amphibien schieben ihren Zeitpunkt ihrer Fortpflanzung innerhalb des Jahres nach vorne (Hällfors et al., 2020, While & Uller, 2014). Für Libellen wurden frühere Flugzeiten im Zusammenhang mit steigenden Temperaturen im Frühling festgestellt (Dingemanse & Kalkman, 2008). Auch trophische Interaktionen wie etwa geografische oder phänologische Diskrepanzen zwischen Vertebraten, Insekten und Pflanzen sind eine Konsequenz des Klimawandels (Renner & Zohner, 2018).

Unter anderem um Prognosen über Veränderungen geografischer Verbreitungen von Arten unter dem Einfluss des Klimawandels zu machen, wurde Ende der 90er Jahre sogenannte Species Distribution Models (SDM) entwickelt (Thuiller et al., 2005). Sie basieren auf dem Konzept der ökologischen Nische, die in der Literatur unterschiedlich definiert wird (Ansprüche einer Art an ihre Umwelt sensu Grinnell & Hutchinson / Auswirkungen einer Art auf ihre Umwelt sensu Elton & MacArthur). Im Falle von SDMs ist die Verbreitung der Art durch biotische Interaktionen und Ressourcenlimitierung beschränkt und kommt demnach der realisierten, ökologischen Nische nach Hutchinson am nächsten (Chase & Leibold, 2009). Bis heute sind SDMs das weitestverbreitete Mittel um Aussagen über die zukünftige Verbreitung von Arten in einem geänderten Klima zu machen. SDMs sind empirische Modelle, die Umweltprädiktorvariablen mit Daten über die Verbreitung der

Zielart verbinden. Die beiden Datentypen, Umwelt und Vorkommen, werden über eine statistische Funktion verbunden. Diese Funktion kommt aus dem Bereich der frequentistischen Statistik, dem Bereich des ‚machine learning‘ oder der Bayesschen Statistik. Das Endergebnis des Models ist eine Karte, die die Vorkommenswahrscheinlichkeit der Art basierend auf den ausgewählten Umweltprädiktoren, angibt. Häufig wird die prognostizierte Karte mittels statistischem Schwellenwert in eine Präsenz-Absenz Karte umgewandelt (Allouche et al., 2006). SDM-Studien, die zukünftige Verbreitungen von Arten modellieren, zeigen ein eindeutiges Bild. Für den Großteil der Arten sinkt das potenzielle Verbreitungsgebiet (Didersky et al., 2018, Niskanen et al., 2019). Besonders gefährdet erscheinen dabei Arten, die in den Hochlagen der Gebirge vorkommen (Engler et al., 2011). Neben Prognosen zu zukünftigen Artenverbreitungen werden SDMs auch benutzt, um die Effizienz von Schutzgebieten zu ermitteln (Araújo et al., 2011).

Trotz der gewaltigen Vorteilen die SDMs der Forschung zu Biodiversität und Klimawandel gebracht haben, ist die Methode nicht frei von konzeptuellen Mängel (Schurr et al., 2012). So muss man bedenken, dass die geografische Verbreitung einer Art nicht unbedingt ihrer realisierten, ökologischen Nische nach Hutchinson entspricht. Nicht überall nämlich wo die Art vorkommt, ergibt die Zusammensetzung an Umweltvariablen eine positive Wachstumsrate. Ein Beispiel dafür sind source-sink Dynamiken (Holt, 2009). Sink-Populationen sind nicht fähig mit den herrschenden Umweltbedingungen langfristig zu überleben und können demnach SDM-Resultate verfälschen. Des Weiteren gibt es gewisse Migrationsimitierungen von Arten, die verhindern, dass Arten tatsächlich überall dort vorkommen, wo ein SDM sie hinprojizieren würde. Beispielsweise haben die Endemiten der Ostalpen ein viel größeres potenzielles Verbreitungsgebiet als sie momentan besiedeln, sie waren (bislang) jedoch nicht in der Lage sich in diese Gebiete auszubreiten (Dullinger et al., 2012b). Schließlich gibt es Arten, die in Gebieten vorkommen, in denen eine positive Wachstumsrate aufgrund einer rezent veränderten Umwelt nicht mehr möglich ist. Je nach Biologie der Art können Populationen diese Änderungen aber unterschiedliche lange tolerieren oder überdauern. Das Ergebnis ist das Entstehen einer Aussterbeschuld, unter anderem als Folge rascher Klimaveränderung (Dullinger et al., 2012b). Ein letzter kritischer Punkt in der Erstellung von SDMs ist die Anzahl an Datenpunkten, die für eine Art zur Verfügung stehen. Um zuverlässige Resultate zu produzieren, sollten gewisse Schwellenwerte nicht unterschritten werden (van Proosdij et al., 2016). Dies ist ungünstig, weil für seltene Arten, wie beispielsweise Endemiten, oft wenig Datenmaterial zur Verfügung steht.

Endemiten sind Arten, die auf ein bestimmtes geografisches Gebiet beschränkt sind (Anderson 1994; Townsend & Watson, 1998). Häufig bezieht man sich dabei auf politische Grenzen, um die Anzahl an Endemiten für ein bestimmtes Land anzugeben. Darüber hinaus spricht man von Subendemiten, wenn sich zumindest 80% der Verbreitung in einem bestimmten Land befinden. Endemismus ist besonders auf Inseln und in Gebirgen ausgeprägt und unterschiedliche Faktoren beeinflussen den Grad an Endemismus in einem Gebiet (Flantua et al., 2020). Erwähnenswert ist die historische klimatische Stabilität (Sandel et al., 2016). Sie ermöglicht ein Überleben von Arten, die in angrenzenden Gebieten mit starken klimatischen Oszillationen nicht überleben können. Die Ostalpen Österreichs liefern dafür gute Beispiele (Tribisch, 2004). Während der letzten Eiszeit waren die periglazialen Randbereiche der Alpen im Osten weitaus weniger stark vergletschert als die Zentralalpen – heute ist dafür der östlichste Teil der Alpen weitaus endemitenreicher als der zentrale Teil. Darüber hinaus ist ein weiterer Treiber von Endemismus die topographische Heterogenität (Dobrowski, 2011). Diese ermöglicht das Überdauern von ungünstigen Bedingungen in vom Regionalklima entkoppelten Mikrorefugia. Durch das Überleben in diesen Refugia wurden Populationen von ihren Schwesternpopulationen isoliert und entwickelten sich schließlich zu eigenen Arten.

Weil Endemiten eine kleinräumige Verbreitung und daher meist auch kleine Gesamtpopulationen aufweisen, spielen sie im Naturschutz eine zentrale Rolle (Manes et al., 2021). Neben dem Klimawandel bedrohen eine intensivierte Landnutzung, invasive Arten und Umweltverschmutzung zusätzlich ihr Überleben (und die Artenvielfalt im Allgemeinen (Diaz et al., 2019)). Für alpine Endemiten werden besondes starke Verbreitungsrückgänge als Folge des Klimawandels prognostiziert (Dirnböck et al., 2011). Als Gegenmaßnahme, insbesondere um die gleichzeitige Bedrohung durch andere Faktoren zu verringern, spielen Schutzgebiete (SG) eine wichtige Rolle. Die potentielle Effizienz von SG zeigt sich unter anderem in den größeren Populationen bedrohter Arten in SGs und deren Bedeutung als Zielort und Schrittstein für Arten, die ihre Areale dem Klimawandel anpassen (Thomas and Gillingham, 2015). Die Ausweitung der SGs ist daher auch ein naturschutzpolitisches Ziel, das in den letzten beiden Jahrzehnten erfreulicherweise bedeutende Fortschritte gemacht hat (Diaz et al., 2019).

Die vorliegende Arbeit beschäftigt sich mit den endemischen Pflanzen Irans und deren Repräsentation im nationalen Schutzgebietsnetzwerk. Der Iran ist ein artenreiches Land und besonders reich an Endemiten (Assadi, 2019, Noroozi et al., 2019a). Der kaukasische, sowie der irano-anatolische „Biodiversitäts-Hotspot“ befinden sich zum Teil auf iranischem Boden, wobei der Anteil am irano-anatolische Hotspot im Iran bei über 50% liegt (Mittermeier et al., 2011). Der Iran ist klimatisch sehr variabel und dies spiegelt sich in den diversen Ökosystemen wieder. Allgemein unterscheidet man drei Großklimata (Djamali et al., 2011). Diese decken sich mit den drei biogeografischen Regionen Irans. Die Nordseite der Gebirgskette Elburs ist sehr niederschlagsreich. Die hohen Temperaturen sowie ein gleichmäßiger Niederschlagsverkauf übers Jahr sorgen dafür, dass die Nordhänge des Elburs bis zu 3000m über dem Meeresspiegel mit dichtem, sommergrünem Wald bedeckt sind (Gholizadeh et al., 2020). Diese Vegetationsform, Teil des kaukasischen Biodiversitäts-Hotspot, bedeckt nur einen geringen Teil der iranischen Landfläche (Noroozi et al., 2020) und ist hinsichtlich ihrer Flora eng mit euro-sibirischen biogeografischen Region verwandt. Der Großteil Irans wird von mediterranoidem bis wüstenartigem Großklima dominiert. Niederschlagsmaxima treten im Spätherbst, Winter und Vorfrühling auf. In den Tieflagen dominieren Wüsten und Halbwüsten. Mit steigender Seehöhe findet man unterschiedliche Vegetationstypen vor (lockere Eichen-Bestände, von Sträuchern dominierte Vegetation, alpine Dornpolsterfluren)(Akhani et al., 2013; Noroozi et al., 2020). Diese Region wird dem irano-anatolischen Biodiversitäts-Hotspot zugeordnet, sowie der irano-turanischen, biogeografischen Region. Der letzte große Klimatyp ist der tropische Klimatyp. Es ist die trockenste klimatische Ausprägung im Iran und beschränkt sich auf den Süden. Das Niederschlagsregime ist ähnlich dem des mediterranoiden Klimatyps, jedoch ist die Niederschlagsmenge geringer, die Trockenperioden länger und die Temperaturen sind höher. Dieser Klimatyp entspricht der saharo-sindischen biogeografischen Region (Noroozi et al., 2020).

Die hohen Gebirgszügen Irans sind besonders reich an Endemiten (Noroozi et al., 2019). Der relative Anteil an Endemiten steigt mit der Seehöhe, wobei 62% der alpinen Pflanzen als endemisch gelten. Um Regionen mit besonders hohen Endemismuswerten zu identifizieren, hat Noroozi et al., 2018 ‚Areas of endemism‘ (AE) innerhalb Irans bestimmt. Diese AE sind Regionen, die sich in der Endemitenvielfalt signifikant von umliegenden Gebieten unterscheiden. Insgesamt wurden bei der Studie von Noroozi et al., 2018 5 AE bestimmt, wobei sich diese 5 Gebiete mit den 5 größten Gebirgsregionen Irans überdecken. Azerbijan, Alburz, Kopet-Dagh, Zagros und Yazd sind die Gebirgsketten, die jeweils ein AE beinhalten. Alle AE befinden sich in der irano-turanischen Region. Noroozi et al., 2019b hat den staatlichen Schutz dieser Gebiete thematisiert. Dabei stellte sich heraus, dass über die Hälfte dieser AE nicht durch Schutzgebiete langfristig gesichert sind. Neben dem unzureichenden rechtlichen Schutz ist der hohe Beweidungsdruck eine Gefahr für die regionale

Flora Irans, die neben einer Reduktion der Biodiversität auch zu starker Bodenerosion führt (Noroozi et al., 2008; Soofi et al., 2018).

Neben der schwachen Abdeckung durch Schutzgebiete und dem hohen Beweisungsdruck wird der Klimawandel die Endemiten Irans möglicherweise zusätzlich unter Druck setzen. Studien zu Klimawandelleffekten auf die Biodiversität Irans sind rar und beziehen sich auf wenige Arten (Safaei et al., 2021; Haidarian et al., 2021). Ein Grund dafür ist die dünne Datengrundlage, die für den Großteil der Arten zur Verfügung steht. Prognosen über zukünftige Verbreitung von Pflanzen mit Hilfe von SDMs sind daher nicht möglich. Semenchuk et al. (in review 2021) haben aber eine Methode entwickelt, die Prognosen über die zukünftige Abdeckung der klimatischen Nischen von Arten durch Schutzgebiete auch auf der Basis spärlicher Datengrundlagen ermöglicht. In meiner Master-Arbeit wende ich diese Methode für die endemischen Pflanzen im Iran an. Dabei interessiert mich, wie sich die Repräsentation der klimatischen Nischen in Schutzgebieten unter momentanen Bedingungen darstellt und wie sie sich unter verschiedenen Klimawandelszenarien ändern würde. Außerdem frage ich, welche Rolle die Größe des aktuellen Verbreitungsgebiets sowie die mittlere Seehöhe der Verbreitung einer Art für die zukünftigen Gewinne oder Verluste an klimatischer Nischenabdeckung durch Schutzgebiete spielt.

Manuscript

Introduction

A deadly cocktail of anthropogenic changes to the planetary ecosystem challenge the survival of many species and may result in a sixth mass extinction (Barnosky et al., 2011; Stephens et al. 2019). Effective conservation measures are key to prevent biodiversity loss (Diaz et al., 2019) and area-based conservation is an important strategy in this context. Indeed, increasing PA cover is the Aichi target of the Convention on Biological Diversity that shows the strongest progress so far (Diaz et al., 2019).

In an area of climate change, however, area-based conservation faces the challenge that the geographic ranges of species will shift (Chen et al., 2011), while PA boundaries remain static. As a consequence, the overlap between species' ranges and the area covered by PA's will change (Lawler, 2015; Hoffmann et al., 2019). Species with sufficient dispersal capacities may be least affected from this emergent mismatch because they are able to overcome the unsuitable non-PA matrix and disperse to adjacent PA's to track their climatic niche (Thomas and Gillingham, 2015). However, these relocation processes are limited by low dispersal ability, geographical barriers (Burrows et al., 2014) and/or the climatic and structural connectivity between PA's (Ward et al., 2020; Parks et al., 2021).

Climate change velocity describes the rate by which climatic conditions shift in geographical space (Laurie et al., 2009). Because of steep climatic gradients it is usually low where topographic heterogeneity is high (Sandel et al., 2016; Harrison and Noss, 2017). This allowed even low-mobility species to follow climatic changes in the past and also fostered survival in microrefugia. The implicit 'climate-change buffer' is one of the reasons why many regions with high relief energy, essentially mountains, are rich in species in general, and often also in endemics (Dobrowski, 2011). While topographic heterogeneity may also buffer regional biodiversity against future climate change impacts (Harrison and Noss, 2017), high endemicity nevertheless makes these areas potential hotspots of (global) biodiversity loss. Moreover, endemic species often are endemic because they are poor dispersers, and those, that are restricted to high elevations may lose large parts of their suitable ranges completely (Dirnböck et al., 2011, Engler et al. 2011). "Climate-smart" conservation appears thus particularly important in these endemism hotspots, for example, by designing PAs such that tracking climatic niches is facilitated without moving outside of PA boundaries (Elsen et al., 2020).

Shifts of climatically suitable ranges are often anticipated by so-called species distribution models (Guisan & Thuiller 2005), and their projections can then be used in designing 'climate-smart' PAs (Guisan et al. 2013). However, these models have data requirements that are often not met for endemic species, because their narrow ranges are often associated with a low number of occurrence records (van Proosdij et al., 2016). On the part of the PAs, the performance of these under-documented species under climate change is mostly assessed by comparing the amount of climatic conditions represented in PAs with the overall climatic conditions in a region or a country (Elsen et al., 2020). This method has the major shortcoming that it disregards the climatic needs of the target species that live in PAs and which may overlap with PA conditions to a variable extent. In a recent study, Semenchuk et al., (in review 2021) used information on species' elevational limits to approximate climatic niches of endemic species independent of occurrence data and compared these inferred niches to the climatic conditions available in PAs within the species' range boundaries. They thereby shifted the focus from the geographical to the ecological domain and asked which part of the inferred niche of species is represented in regional PAs now and in a warmer future. Because

small heterogeneous landscapes can contain more climatic variability than large but homogeneous ones, this shift of focus delivers information that can deviate from estimates of geographical range change. While the latter is clearly important (Staude et al., 2018), ecological niche loss can also have strong implications for the fate of species because it affects future intraspecific diversity and hence, among other things, the potential of evolutionary adaptation to a changing climate (e.g. Balint et al. 2011).

Iran is a country with a complex topography and covers more than 50% of the Irano-Anatolian biodiversity hotspot (Noroozi et al., 2018). Endemic species particularly contribute to the flora of the high mountain chains that intersect the country, with 62% of alpine plants considered endemic to Iran (Noroozi et al. 2015). At the same time, occurrence records of endemic vascular plants are scarce (Noroozi et al., 2019a), mainly due to the large and difficult-to-access terrain as well as limited financial resources for botanical exploration. The Iranian mountain areas are moreover poorly protected through PAs (Noroozi et al., 2019b) and possible effects of climate change on their future ranges have been hardly explored so far.

Here, I apply the method of Semenchuk et al., (in review 2021) to the endemic flora of Iran. Based on information on the geographical and elevational distribution of those endemic and subendemic vascular plants with relatively well documented occurrence, I determine the representation of each species' niche within the PA network under current climate and under future climatic conditions. Further, I analyze whether high-elevation species suffer from disproportional loss of niche representation in PAs in the future and whether there is a link between geographic range and climatic niche representation within PAs.

Material and Methods

Study area

Iran, with its 1.6 million km² total surface, is a country situated in Southwest Asia. The heterogenous geology and topography are the result of high tectonic activity (Berberian & King, 1981). Vast deserts and semi-deserts dominate central and eastern Iran. They are surrounded by high mountain ranges which restrain precipitation from reaching the interior of the country (Djamali et al., 2011). The Irano-Turanian and the Caucasian biodiversity hotspot intersect the territory of Iran highlighting its outstanding biodiversity (Mittermeier et al., 2011). Five Areas of Endemism (AE) are located within the Irano-Turanian hotspot. These AE are particularly rich in endemic species (Noroozi et al., 2018, Noroozi et al., 2019a) and each AE is associated with one of the five major mountain areas of Iran (Azerbaijan Plateau, Alborz, Kopet Dagh-Khorassan, Zagros and the Yazd-Kerman massifs). The contribution of endemics to the total flora in these areas increases with elevation and peaks in the alpine life zone (Noroozi et al., 2008). Despite their significant contribution to biodiversity, the legal protection status of the AE is insufficient (Noorozi et al., 2019b).

Species and Protected Area data

I used occurrence point data for all known 2597 endemic and subendemic (range > 80% in Iran) vascular plants in Iran (Noroozi 2019a). However, most of these species are understudied and the elevational and geographical distribution is insufficiently documented. To avoid bias resulting from these knowledge gaps, I excluded species with a recorded elevational range lower than 800 meters, leaving me with 1277 species. For this subset, I defined their elevational range by the lowest and highest elevation reported with any occurrence record, respectively. Finally, these elevational ranges were controlled and adjusted by Jalil Noroozi, an expert on Iran's endemic flora.

I buffered all reported occurrences of each species by a geographical radius of 20km and interpreted the intersection of all these buffer areas as the species' current potential extent of occurrence. I

overlaid this extent of occurrence by a 250 x 250m digital elevation model (Jarvis et al. 2008) and selected all cells that fall within the elevational range of the species. These cells I considered as the species' potential area of occupancy (cf. Fig. 1).

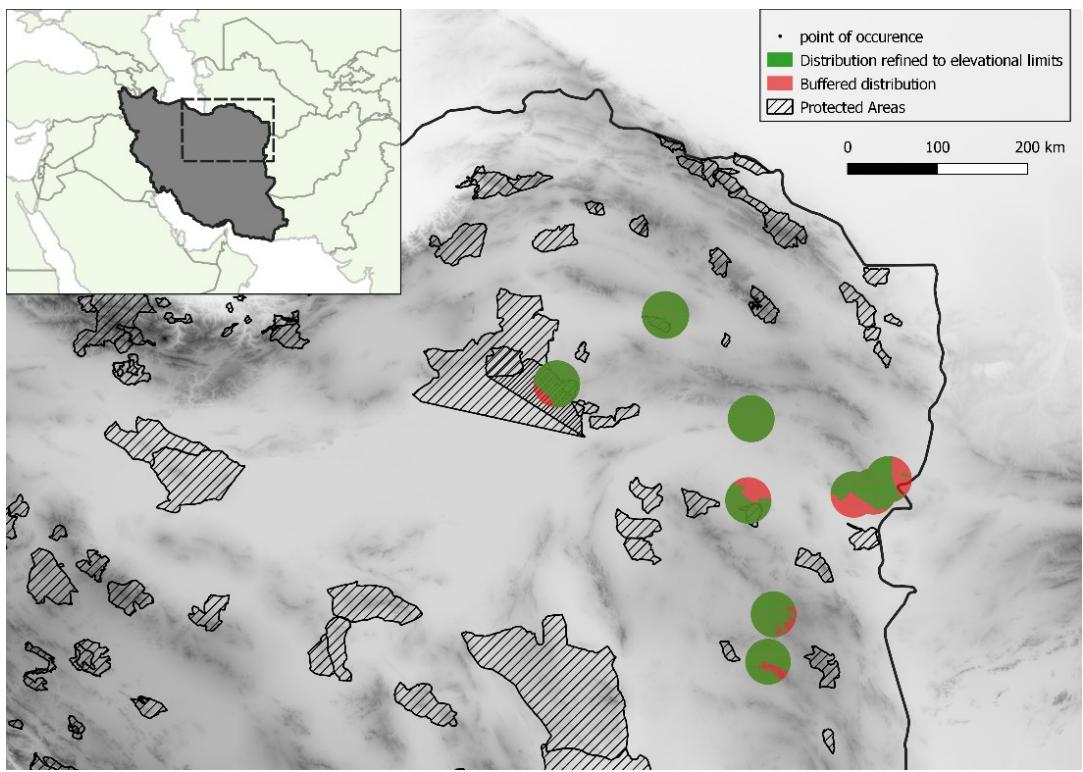
The Iranian Department of Environment provides data on the national Protected Area (PA) network (Kolahy et al., 2013). I overlaid the PA network with each species' potential area of occupancy and defined, for each species, the PAs within its current extent of occurrence as reachable and hence relevant for the respective species until the end of the century. In doing so, I implicitly assume that each species' unrecorded occurrence and/ or dispersal capacity is bound with the extent of occurrence as defined above.

Climate data and scenarios

I used mean annual temperature (Tmean) and mean annual precipitation (Prec) to create raster maps of current and future climatic conditions for Iran. Data were extracted from the WorldClim data base for current and future climate in 30" resolution (www.worldclim.org). Among Global Circulation Models (GCM) available in this database, I selected ACCES1-0 (Ackerley, D. and Dommelget, D., 2016) as it is among the best GCM's in modelling climatic key processes (McSweeney et al., 2015). To account for uncertainty in climatic predictions, I used predictions for two IPCC5 scenarios representing medium (RCP 4.5) and severe (RCP 8.5) climate change and I selected two time steps (2050 and 2070).

I overlaid the species' area of potential occupancy and the PA maps with the climatic raster maps to calculate (1) the species' climatic niches (based on current climate), (2) its future potential area of occupancy (based on its climatic niche), and (3) the climatic space within the associated PAs under both current and future climatic conditions.

Temperature and Precipitation are major drivers of species distribution (Gaston, 2003). However, constructing the climatic niches of the species based on Tmean and Prec may neglect the specific climatic requirements of individual species (Körner and Hiltbrunner, 2018). However, with the data available, species-specific climatic niche construction is impossible. Hence, I keep with Semenchuk et al., (in review 2021) and consider the selected climatic variables as best available niche descriptors , in particular as Tmean and Prec often closely correlate with other climatic variables.



*Figure 1: Derivation of potential area of occupancy from occurrence point data and information on elevational range limits illustrated for the vascular plant species *Acantholimon austro-iranicus*. The red area is the buffer (20 km) around the occurrence points, the sum of which represents the potential extent of occurrence, and the green area is the area within the buffer restricted to the elevational range limits. For more details see text. Background map is a digital elevation model. Darker shades represent higher altitudes.*

Climate niche construction

I used the function `ecospat.grid.clim.dyn` from the package `ecospat` in R (Di Cola et al. 2017) to calculate a bivariate (in Tmean and Prec) density kernel (ranging from 0 to 1) for both the species' potential area of occupancy and its associated PAs under current as well as under future climatic conditions (cf. Fig. 2). I defined the kernel of the species' potential area of occupancy under current climatic conditions as its climatic niche and the kernels of the associated PAs as the climatic niche space offered by these PAs under either current or future conditions. I used the species' potential area of extent as the 'background area' necessary for calculation of the density kernels of species and PAs and their further comparison.

The future climate will be different from the current one. Currently available (analogous) combinations of climatic factors will be complemented by new (non-analogous) combinations of climate variables. This non-analogous climate is new for the species and the individual species' response to it is unpredictable (Fitzpatrick & Hargrove, 2009). I thus restricted all analyses to analogous climates and did not consider the emergence of non-analogue climates in PAs or species' area of occupancy in my calculations. In doing so, I neglect the possibility of niche adaptation and remain conservative in my predictions.

Semenchuk et al. (in review 2021) assume that dispersal does not limit the species in colonizing its future climatic niche within their current extent of occurrence. However, the extent of occurrence remains unchanged, i.e. migration beyond its borders is not considered, based on the assumption

that many endemics are likely strongly dispersal-limited. Likewise, PA's geographic distribution are assumed to remain unchanged in the study because I want to evaluate the potential of the current PA network.

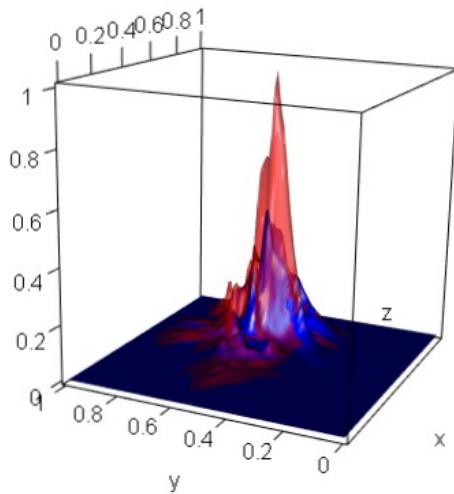


Figure 2: Bivariate density kernel (= climatic niche) of *Cousina calocephala* (red) and the climatic space in the associated PA's (blue) under current climate. The overlap between the two volumes is the suitable climatic space for the species in its associated PA's with Tmean on the x-axis, prec on the y-axis and the volume of the climatic niches of the species and the climatic space in associated PAs on the z-axis.

Climatic niche representation

The representation of the species' climatic niche in its associated PAs is the percentage of volume of the species' bivariate density grid (species' grid) covered by the PA's bivariate density grid (PA grid). To calculate this percentage, I overlaid the species' grid and the PA grid. I removed the densities in the PA grid where the species' grid was zero. Then, I divided the PA grid's remaining non-zero values by the values of the species' grid. The outcome is a value ranging from 0 (no representation) to 1 (full representation). I calculated the representation of each species for each combination of time step and scenario.

Statistical analysis

I describe the temporal variation of the climatic representation in PAs for all species and for all time steps and scenarios with a generalized linear mixed effect model using the functions glmmTMB and betareg in R (Brooks et al., 2017; Cribari-Neto et al., 2009; R Core Team, 2017). The response variable in the model is the climatic niche representation (0 to 1) and I include species as random factor. Assuming the response to follow a beta distribution I applied the beta-family in the model. Some species have no climatic representation in their associated PA's at one or several time steps. Therefore, I applied the lemon squeezer transformation of Smithson and Verkuilen (2006) to deal with the zero values in the data prior to modelling. This is a necessary step as the link function of the beta distribution is not defined for values equal to zero (or one, but climatic representation never reached one). In total, I apply two models, differing in their predictor interaction term.

For the first model, the transformed climatic niche representation is the response and the different combinations of time step and scenario serve as predictor variables. I add mean elevation as an

interaction term to the time step/scenario predictors to account for the expected effect of elevational distribution on climatic niche representation. The visual inspection of the data suggests a unimodal distribution of the response variable. Therefore, I include mean elevation as a quadratic interaction term. I tested whether the interaction between time step/ scenario and mean elevation was significant by comparing the full model to the corresponding additive model by means of a likelihood ratio test. Then, I performed Tukey-corrected pair-wise comparisons across all factor levels in the model to test whether the climatic scenarios and mean elevation differ statistically significantly in terms of niche representation, taking pair-wise contrasts as significantly different with $p < 0.05$.

In a second model, I want to evaluate whether geographical range size affects the climatic niche representation in PAs. Therefore, I group the species according to the number of sectors ($= 0.5 \times 0.5^\circ$ cells) they occur. Species present in three or less sectors are considered as range-restricted, while species present in more than three grid cells are considered as (relatively) widespread endemics (Noroozi et al., 2019a). I use this categorical variable together with time step/scenario as interactive fixed effect in my second model. The response variable is, as in the first model, the transformed climatic niche representation. Significance tests are analogous to the first model.

Results

From the 1277 species considered in my study, 85 species have no PA within their extent of occurrence or the elevational ranges of the species and its associated PAs do not match. Hence, 1192 species remain for the calculation of climatic representation of the species within their associated PAs.

In 2020, an average of 51.8% of the species' climatic niche was represented in the associated PAs. Only seven species have no representation of their climatic niches within their associated PAs whereas 27 species show over 90% climatic representation.

In 2050, mean climatic representation significantly decreases to 31.4% and 29.3% and to 26% and 22.9% in 2070 for the 4.5 and the 8.5 scenario, respectively (Fig.3). While differences among scenarios are insignificant in 2050, they become significant in 2070. In 2070, the climatic representation drops down to zero for 72 species under the moderate scenario. For the severe scenario, 69 species experience complete loss of their climatic representation in PAs in 2070. Regarding the winners in climatic representation, 169 species gain climatic representation in comparison to 2020 at any time step or scenario. In 2070, 85 species have an increased climatic representation in PAs under the moderate scenario and 59 species under the severe scenario.

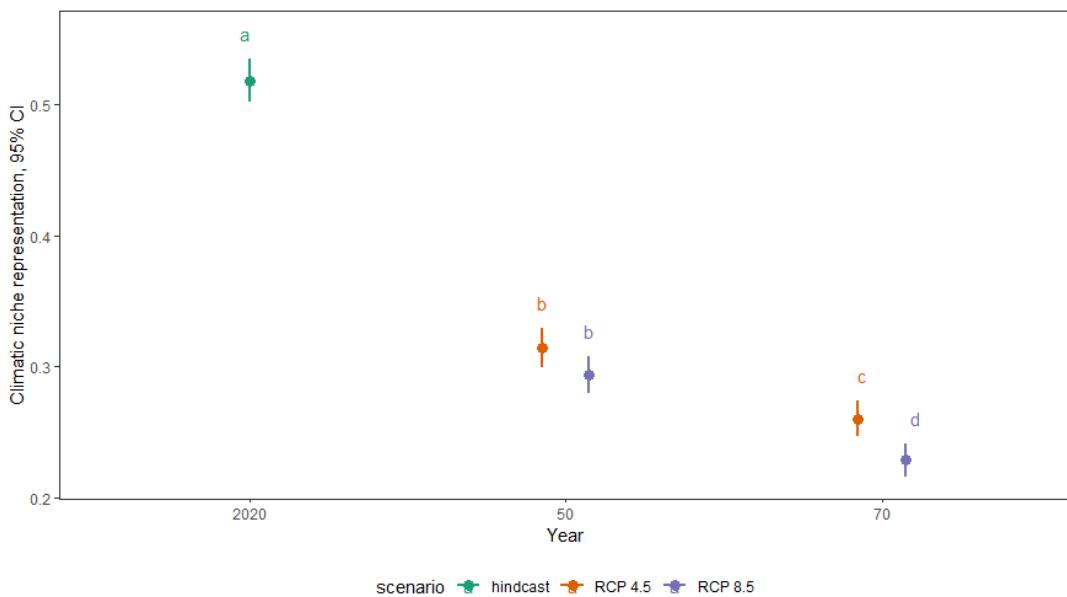


Figure 3: Climatic representation within associated PAs across all endemic species analysed. Shown are estimates and 95% confidence intervals (CI) from a generalized linear mixed effects model including a significant interaction between time step/scenario. Different colors represent different climatic scenarios, RCP 4.5 being a moderate and RCP 8.5 a severe scenario. Letters indicate significant pair-wise contrasts ($p < 0.005$).

Under current climatic conditions, climatic representation is not equally distributed across elevation but peaks at a mean elevation of about 2500 m.a.s.l. (Fig. 4). Under future climate this peak decreases in elevation to about 2000 m.a.s.l. and the decrease in climatic representation becomes particularly pronounced at highest elevations. As a consequence, a likelihood ratio test between a model including the interaction of elevation with time step and scenario and an additive model was highly significant ($p < 0.0001$)

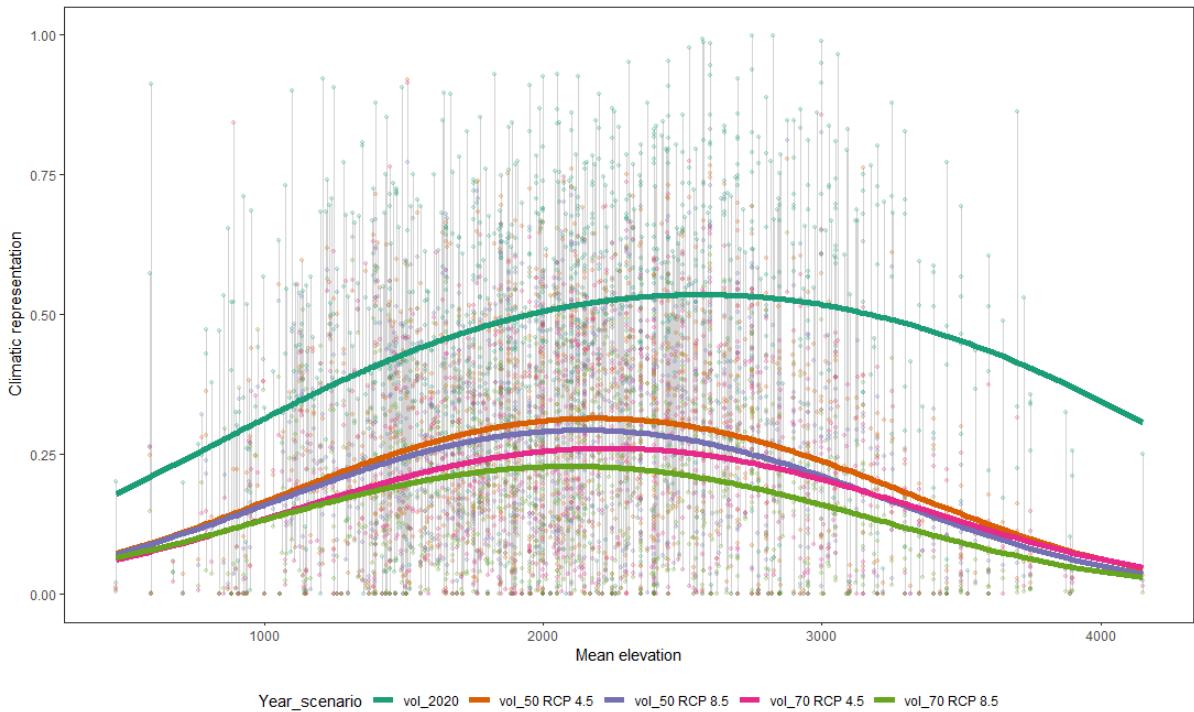


Figure 4: Representation of species' climatic niches within PAs in relation to mean elevation. Colored lines represent different combinations of time steps (2050 & 2070) and scenarios (RCP 4.5 & RCP 8.5). Grey vertical lines connect data points of one species.

Current climatic representation did not differ among restricted or widespread species. Under future scenarios and time steps, however, range-restricted species lost significantly more climatic representation than widespread endemics (Tukey-corrected pair-wise comparisons $p<0.05$). Hence a model including an interaction between the predictor variables time-step/ scenario and the categorial variable "range-restricted" was explaining the data significantly better than an additive model ($p < 0.0001$). For range-restricted species, the current climatic representation of 47.9% drops to 13.5% in 2050 and further to 8.9% in 2070 for RCP 8.5. The climatic representation of widespread species in PAs is at 48.5% today, and shrinks down to 26.9% in 2050 and 19.5% in 2070 for RCP 8.5. Under RCP 4.5, decreases in climatic representation are less pronounced for widespread species, but almost equally pronounced as in RCP 8.5 for range-restricted species. As a consequence, for widespread species, the difference between the scenarios was significant at both time steps (Tukey-corrected pair-wise comparisons $p<0.05$), while it was insignificant for range-restricted species at both time steps.

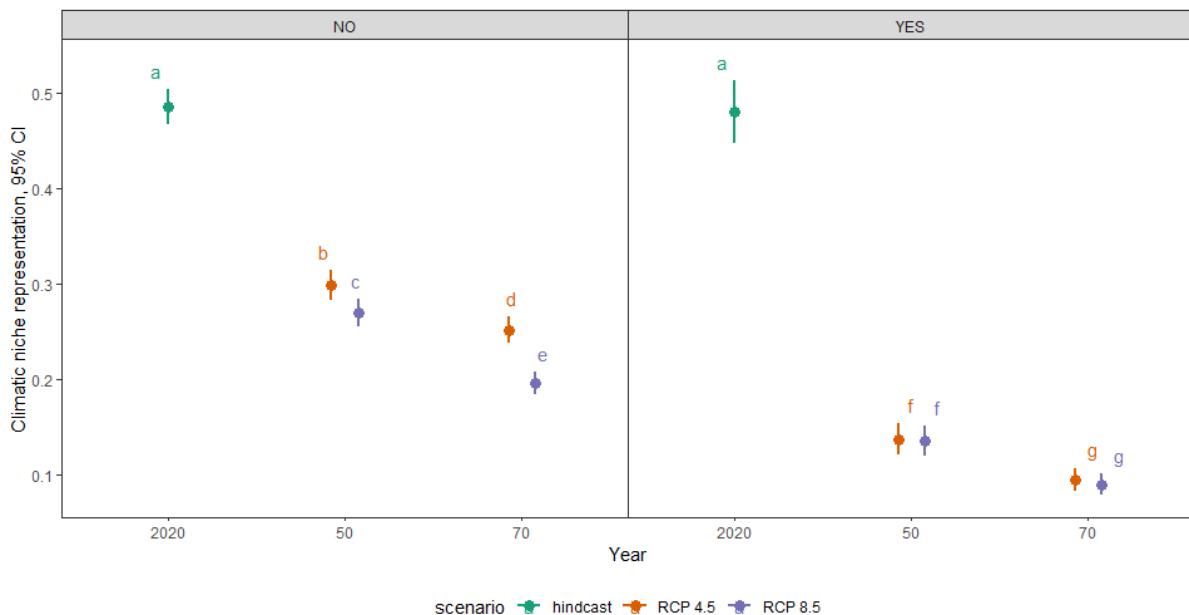


Figure 5: Climatic representation within associated PAs for widespread and range-restricted species separately. Shown are estimates and 95% confidence intervals (CI) from a generalized linear mixed effects model including a significant interaction between time step/scenario and extent of geographic distribution grouping. Different colors represent different climatic scenarios, RCP 4.5 being a moderate and RCP 8.5 a severe scenario. Letters indicate significant pair-wise contrasts ($p < 0.005$).

Discussion

Under current climatic conditions, the Iranian PA-network covers the climatic niches of Iran's endemic plants only to about 50%, on average. With climate change, the fraction of endemics' niches protected further decreases to values below 30%. These findings suggest that the Iranian PA network is not prepared for the challenges imposed by anthropogenic climate change, at least not with respect to protecting the endemic flora of the country.

My results suggest that the climatic niches of species thriving at lowest and highest elevations have particularly poor representation within PAs. Under current conditions, this suggests that PAs cover the largest fraction of the total Iranian mountains at intermediate elevations. As the climate warms, the climatic niches of the species become increasingly rare in geographical space because colder temperatures are not realized anymore in the landscape, i.e. mountains are not high enough. As a consequence, the fraction of the niche covered by PAs also shrinks. These findings corroborate earlier evidence that endemics of high elevations are particularly vulnerable to climate change (Dirnböck et al., 2011). By contrast, the endemics of lower elevations mostly thrive in lower parts of mountain chains, but have a great proportion of their (spatially) associated PAs located in extensive, desert PAs bordering these mountains. As many of these endemics are not adapted to the dry desert climate, but need somewhat moister conditions offered at least locally in mountainous terrain, their climatic niches are hardly covered by desert PAs under current conditions. In the future, when conditions in desert PAs becomes even warmer and drier, this mismatch even increases and is obviously only partly buffered by an increased overlap with climatic conditions offered by higher lying PAs, because PA-cover has its peak at elevations at least 1000 m above the foothills of the mountains (Malakoutikhah et al., 2018). The species the climatic ranges of which are currently represented best by the Iranian PA occur predominantly around 2500 m. In the future, this peak

shifts to a lower elevation (~2000 m) because the species that currently have their peak at this elevation will then have their niches represented in the greater area of PAs higher up.

Range-restricted and widespread endemics significantly differ in their climatic representation within PAs in the future. While under current climatic conditions climatic representation is similar, representation losses in the future are much more pronounced for range-restricted endemics. Whatever the reason of this difference, it suggests that the magnitude of ecological response to climate change may indeed be related to range size (Williams et al., 2021) and that climatic niche size may be correlated to future climatic disequilibrium (Perret et al., 2018). Hence, weak niche representation of range-restricted species in PAs might increase their extinction risk in comparison to species with a wider distribution. (Staude et al., 2018).

As distributions of endemic plants in Iran are associated with topographic heterogeneity, many species will likely find microrefugia within their range (Noroozi et al., 2019a). Climatic conditions of such microrefugia are decoupled from regional climate and vary on a small spatial scale (Dobrowski, 2011). The coarse resolution of the climatic data used to create species' climatic niches misses such small-scale variation in the climatic conditions. Hence, finer-scaled climate data is needed to integrate the critical role of microrefugia for species' survival (Potter et al., 2013). This also applies to survival of species in PAs in general, and to representation of species' niches in a particular area. However, a shift of the macroclimate away from the climatic requirements of species will most likely reduce the area of appropriate climate available to the species (Scherrer & Körner 2012). So even if a macroclimatic study like mine cannot quantify area or niche loss accurately, the main trends of decrease or increase are most likely captured correctly.

A critical assumption of the study is the limited dispersal ability of the species. Species are assumed to be unable to disperse outside of their current range extent and this constrains the colonization of new PAs. For endemic plants, the lack of dispersal ability often is an important reason of their narrow ranges (Essl et al., 2011). However, genetic shifts in the dispersal ability might occur rapidly under high environmental pressure (Bradshaw et al., 2001). The field of dispersal genetics is poorly investigated and therefore predictions about the genetic response to climate change are contentious (Saastamoinen et al., 2018). In this context, the longevity of endemic plants and the predicted pace of climate change both counteract anticipated gene shifts (Cotto et al., 2017).

Another key assumption of the study is the species' (potential) distribution which is defined as the entire space within the elevational range limits. This is unlikely for most of the species as several factors further restrict the distribution of a species. Besides temperature and precipitation, other abiotic factors such as geology control species distributions (Gaston et al., 2003). Biotic interactions further interfere with abiotic species distribution patterns. Indeed, climate change-induced tree line shifts are common and put alpine species under competitive pressure (Valavi et al., 2019, Gatti et al., 2019). Facilitation is another key component shaping biodiversity in alpine environments (Cavieres et al., 2014). Cushion plants in Iran's alpine habitats ameliorate local microclimatic conditions and allow the establishment of species that are adapted to less severe environments (Butterfield et al., 2013). Hence, the future distribution of such cushion plants (e.g. many species of *Acantholimon*) will affect future diversity patterns. Finally, the study does not account for changes in climatic representation that are based on species-specific ecological niche differences among spatially separated subpopulations (Wasof et al., 2013).

To conclude, my results suggest that the endemic flora in Iran will increasingly lack adequate area-based protection in the future. Identification of microrefugia could help to close conservation gaps and make the Iranian PA network fit for climate change. In this context, the tight link between water

availability and microrefugia emphasizes research on the spatial distribution of ‘hydrologic climate buffers’ (Gentilli et al., 2020, McLaughlin et al., 2017). From a species perspective, protection of range-restricted species distributed at low and high altitudes should be prioritized as compared to those predominating at intermediate elevations. The methods applied here are certainly too coarse to allow for accurate quantifications of niche representation loss, but they give a first impression on the extent of threat of endemic biodiversity in this understudied biodiversity hotspot.

Schlussfolgerungen

Meine Master-Arbeit bewertet zum ersten Mal Klimawandaleffekte für einen großen Teil der endemischen Flora Irans. Dabei fokussiere ich mich auf die klimatische Nische der Art. In der Tat ist das Klima auf einer makroökologischen Skala ein Hauptfaktor für die Artenverteilung in der Landschaft (Gaston et al., 2003). Ich beziehe mich ausschließlich auf die Abdeckung der klimatischen Nischen in SG, weil die intensivierte Landnutzung auch ohne Klimawandel bereits massive Bestandesrückgänge der Artenvielfalt außerhalb von SG fordert. SG bilden die letzte Bastionen, in denen Arten basierend auf einem rechtlichen Fundament von Bedrohung durch Landnutzung zumindest theoretisch geschützt sind. Der Klimawandel unterscheidet jedoch nicht zwischen geschützten und ungeschützten Flächen. Dies unterstreicht das theoretische Konzept, das dieser Arbeit zugrunde liegt.

Bereits unter momentanen Klimabedingungen muss man nach meinen Ergebnissen dem iranischen SG-Netzwerk eine ungenügende Note eintragen. Mit durchschnittlich 50% Abdeckung der klimatischen Nische in SG ist die endemische Flora Irans bereits jetzt unzureichend geschützt. Das starke Ungleichgewicht zwischen dem Beitrag dieser Arten zur nationalen und globalen Artenvielfalt und der Aufmerksamkeit, die ihnen in naturschutzpolitischen Entscheidungsprozessen gewidmet wird, ist auffällig. Meine Ergebnisse unterstreichen den unzureichenden Schutz von Irans floristisch wertvollsten Gebieten, auf den bereits Noroozi et al., 2019b aufmerksam macht. Tatsächlich korrelieren meine Ergebnisse mit den seinen, wenn auch der methodische Zugang ein anderer ist. Um die momentane Situation zu verbessern ist es erforderlich, neue SG auszuweisen. Schließlich muss verhindert werden, dass Beweidung die potenziellen Erfolge dieser SG zunichtemacht. Irans ländlicher Raum ist von der Landwirtschaft dominiert. Durch Aufklärung der lokalen Bevölkerung über die Bedeutung von SG kann eine Symbiose zwischen Naturschutz und Bevölkerung entstehen. Gleichzeitig wird dieses Unterfangen nur dann funktionieren, wenn man die finanzielle Situation der Menschen verbessert. In dem von wirtschaftlichen Sanktionen erschöpften Iran können SG auch dabei helfen, die Ökonomie der ländlichen Region zu verbessern. Ökotourismus ist ein wachsender Wirtschaftszweig und der Iran hat dafür die perfekten Voraussetzungen (Fennell, 2020).

Damit solche Ideen auch in die Realität umgesetzt werden können, braucht es zuallererst die Planung neuer SG und ein effizientes Schutzgebietsmanagement. Meine Ergebnisse zeigen nämlich eine dramatische Reduzierung der Abdeckung der klimatischen Nischen der Endemiten in der Zukunft. Je nach Szenario und Zeitpunkt sinken die Werte auf durchschnittlich 20-30%. Diese Verluste an klimatischer Abdeckung sind nicht homogen entlang des Höhengradienten verteilt. Das schlechte Abschneiden von Tiefland- und Hochgebirgsarten ist Segen und Fluch zugleich. Neue SG einrichten ist möglich, mit einem immer stärker werdenden Klimawandel werden die Arten der Geschwindigkeit der Erwärmung allerdings nicht folgen können. Schließlich setzen die Berggipfel den einer möglichen Migration von Arten ein Ende. Daraus schließe ich, dass der Schutz der Tieflandarten mit vergleichsweise mäßigem Aufwand verbessert werden kann. Das Schicksal der Hochgebirgsarten dagegen wird vor allem von dem Ausmaß an klimatischer Erwärmung bestimmt werden.

Ich habe die geografische Verbreitung der Arten in 2 Klassen (weitverbreitet und eingeschränkt verbreitet) eingeteilt. Dabei erkennt man eine bessere Abdeckung der klimatischen Nischen von weitverbreiteten Arten. In der Praxis bedeutet dies, dass bei der Planung von SG Arten mit beschränkten Verbreitungen höher gewichtet werden sollten als weitverbreitete Arten.

Die Planung und Implementierung neuer SG ist unumgänglich, wenn der Iran seine Artenvielfalt erhalten will. Konflikte mit traditioneller Landnutzung durch die ländlichen Bevölkerung sollten nach

Möglichkeit verhindert werden. Eine positive Grundhaltung der lokalen Bevölkerung, durch Aufklärung und Einbindung in die SG-Gestaltung ist für den Naturschutz-Erfolg wesentlich. Um die Planung von SGs effektiver zu gestalten muss außerdem die Datengrundlage verbessert werden, insbesondere auch im Hinblick auf die Verbreitung schützenswerter Arten. Dafür ist es notwendig, genügend finanzielle Ressourcen bereit zu stellen. Gebiete mit einer hohen Anzahl an Endemiten, insbesondere solcher mit kleinem Verbreitungsgebiet, sollten bei der Einrichtung von SG als prioritär betrachtet behandelt werden. Eine klimawandelangepasste Planung potenzieller SG sollte Gebiete mit möglichst ausgeprägten klimatischen Gradienten bevorzugen. In ariden Gebieten wie dem Iran spielt dabei neben den Tempaturgradienten die Wasserverfügbarkeit eine entscheidende Rolle (Rodriguez-Iturbe et al., 2007). Gebiete, deren Wasserhaushalt von der umgebenden Landschaft entkoppelt ist, können als Refugien für speziell angepasste Arten dienen (McLaughlin et al., 2017). Als Grundlage für eine gezielte Planung könnten in dieser Hinsicht Modelle dienen, die die Wechselwirkungen von Vegetation und Bodenfeuchtigkeit simulieren (Tietjen et al., 2009). Schließlich spielt für die langfristige Erhaltung der Artenvielfalt aber eine konsequente, weltweite Klimawandelpolitik die wichtigste Rolle.

Acknowledgement

I want to thank Stefan Dullinger for being an exemplary supervisor under challenging circumstances during COVID-19 pandemic. I also want to thank Jalil Noroozi for providing the data on the Iranian flora and for sharing his knowledge with me. Thanks to Philipp Semenchuk who helped me a lot with the methodological part of the Thesis. Thanks to Johannes Wessely and Dietmar Moser for intellectual input. Finally, I want to thank my family and friends for never ending support and encouragement.

Literature

- Ackerley, D., & Dommenget, D. (2016). Atmosphere-only GCM (ACCESS1.0) simulations with prescribed land surface temperatures. *Geoscientific Model Development*, 9(6), 2077-2098.
- Akhani, H., Mahdavi, P., Noroozi, J., & Zarrinpour, V. (2013). Vegetation patterns of the Irano-Turanian steppe along a 3,000 m altitudinal gradient in the Alborz Mountains of Northern Iran. *Folia Geobotanica*, 48(2), 229-255.
- Allison, I., Alley, R. B., Fricker, H. A., Thomas, R. H., & Warner, R. C. (2009). Ice sheet mass balance and sea level. *Antarctic Science*, 21(5), 413-426.
- Allouche, O., Tsoar, A., & Kadmon, R. (2006). Assessing the accuracy of species distribution models: prevalence, kappa and the true skill statistic (TSS). *Journal of applied ecology*, 43(6), 1223-1232.
- Araújo, M. B., & Rahbek, C. (2006). How does climate change affect biodiversity? *Science*, 313(5792), 1396-1397.
- Assadi, M. (2019). Flora of Iran. *Iran Nature*, 4(2), 29-41.
- Bálint, M., Domisch, S., Engelhardt, C. H. M., Haase, P., Lehrian, S., Sauer, J., ... & Nowak, C. (2011). Cryptic biodiversity loss linked to global climate change. *Nature climate change*, 1(6), 313-318.

- Barnosky, A. D., Matzke, N., Tomaia, S., Wogan, G. O., Swartz, B., Quental, T. B., ... & Ferrer, E. A. (2011). Has the Earth's sixth mass extinction already arrived?. *Nature*, 471(7336), 51-57.
- Bebber, D. P., & Butt, N. (2017). Tropical protected areas reduced deforestation carbon emissions by one third from 2000–2012. *Scientific reports*, 7(1), 1-7.
- Berberian, M., & King, G. C. P. (1981). Towards a paleogeography and tectonic evolution of Iran. *Canadian journal of earth sciences*, 18(2), 210-265.
- Bradshaw, W. E., & Holzapfel, C. M. (2001). Genetic shift in photoperiodic response correlated with global warming. *Proceedings of the National Academy of Sciences*, 98(25), 14509-14511.
- Broennimann, O., Fitzpatrick, M. C., Pearman, P. B., Petitpierre, B., Pellissier, L., Yoccoz, N. G., ... & Guisan, A. (2012). Measuring ecological niche overlap from occurrence and spatial environmental data. *Global ecology and biogeography*, 21(4), 481-497.
- Brommer, J. E., Lehikoinen, A., & Valkama, J. (2012). The breeding ranges of Central European and Arctic bird species move poleward.
- Brooks, M. E., Kristensen, K., Van Benthem, K. J., Magnusson, A., Berg, C. W., Nielsen, A., ... & Bolker, B. M. (2017). glmmTMB balances speed and flexibility among packages for zero-inflated generalized linear mixed modeling. *The R journal*, 9(2) 378-400.
- Burrows, M. T., Schoeman, D. S., Richardson, A. J., Molinos, J. G., Hoffmann, A., Buckley, Loczanska, E. S. (2014). Geographical limits to species-range shifts are suggested by climate velocity. *Nature*, 507(7493), 492-495.
- Butterfield, B. J., Cavieres, L. A., Callaway, R. M., Cook, B. J., Kikvidze, Z., Lortie, C. J., ... & Brooker, R. W. (2013). Alpine cushion plants inhibit the loss of phylogenetic diversity in severe environments. *Ecology Letters*, 16(4), 478-486.
- Cavieres, L. A., Brooker, R. W., Butterfield, B. J., Cook, B. J., Kikvidze, Z., Lortie, C. J., ... & Callaway, R. M. (2014). Facilitative plant interactions and climate simultaneously drive alpine plant diversity. *Ecology letters*, 17(2), 193-202.
- Chase, J. M., & Leibold, M. A. (2009). *Ecological niches*. University of Chicago Press.
- Chen, I. C., Hill, J. K., Ohlemüller, R., Roy, D. B., & Thomas, C. D. (2011). Rapid range shifts of species associated with high levels of climate warming. *Science*, 333(6045), 1024-1026.
- Cohen, J. M., Lajeunesse, M. J., & Rohr, J. R. (2018). A global synthesis of animal phenological responses to climate change. *Nature Climate Change*, 8(3), 224-228.
- Cotto, O., Wessely, J., Georges, D., Klonner, G., Schmid, M., Dullinger, S., ... & Guillaume, F. (2017). A dynamic eco-evolutionary model predicts slow response of alpine plants to climate warming. *Nature Communications*, 8(1), 1-9.
- Cribari-Neto, F., & Zeileis, A. (2009). Beta regression in R.
- Di Cola, V., Broennimann, O., Petitpierre, B., Breiner, F. T., d'Amen, M., Randin, C., ... & Guisan, A. (2017). ecospat: an R package to support spatial analyses and modeling of species niches and distributions. *Ecography*, 40(6), 774-787.
- Díaz, S., Settele, J., Brondízio, E. S., Ngo, H. T., Agard, J., Arneth, A., ... & Zayas, C. N. (2019). Pervasive human-driven decline of life on Earth points to the need for transformative change. *Science*, 366(6471).

- Dingemanse, N. J., & Kalkman, V. J. (2008). Changing temperature regimes have advanced the phenology of Odonata in the Netherlands. *Ecological Entomology*, 33(3), 394-402.
- Dirnböck, T., Essl, F., & Rabitsch, W. (2011). Disproportional risk for habitat loss of high-altitude endemic species under climate change. *Global Change Biology*, 17(2), 990-996.
- Djamali, M., Akhani, H., Khoshravesh, R., Andrieu-Ponel, V., Ponel, P., & Brewer, S. (2011). Application of the global bioclimatic classification to Iran: implications for understanding the modern vegetation and biogeography. *Ecologia mediterranea*, 37(1), 91-114.
- Dobrowski, S. Z. (2011). A climatic basis for microrefugia: the influence of terrain on climate. *Global change biology*, 17(2), 1022-1035.
- Dullinger, S., Gatringer, A., Thuiller, W., Moser, D., Zimmermann, N. E., Guisan, A., ... & Hülber, K. (2012a). Extinction debt of high-mountain plants under twenty-first-century climate change. *Nature climate change*, 2(8), 619-622.
- Dullinger, S., Willner, W., Plutzar, C., Englisch, T., Schrott-Ehrendorfer, L., Moser, D., ... & Niklfeld, H. (2012b). Post-glacial migration lag restricts range filling of plants in the European Alps. *Global Ecology and Biogeography*, 21(8), 829-840.
- Dyderski, M. K., Paź, S., Frelich, L. E., & Jagodziński, A. M. (2018). How much does climate change threaten European forest tree species distributions?. *Global change biology*, 24(3), 1150-1163.
- Elsen, P. R., & Tingley, M. W. (2015). Global mountain topography and the fate of montane species under climate change. *Nature Climate Change*, 5(8), 772-776.
- Elsen, P. R., Monahan, W. B., & Merenlender, A. M. (2018). Global patterns of protection of elevational gradients in mountain ranges. *Proceedings of the National Academy of Sciences*, 115(23), 6004-6009.
- Engler, R., Randin, C. F., Thuiller, W., Dullinger, S., Zimmermann, N. E., Araújo, M. B., ... & Guisan, A. (2011). 21st century climate change threatens mountain flora unequally across Europe. *Global Change Biology*, 17(7), 2330-2341.
- Essl, F., Dullinger, S., Plutzar, C., Willner, W., & Rabitsch, W. (2011). Imprints of glacial history and current environment on correlations between endemic plant and invertebrate species richness. *Journal of Biogeography*, 38(3), 604-614.
- Fitzpatrick, M. C., & Hargrove, W. W. (2009). The projection of species distribution models and the problem of non-analog climate. *Biodiversity and Conservation*, 18(8), 2255-2261.
- Flantua, S. G., Payne, D., Borregaard, M. K., Beierkuhnlein, C., Steinbauer, M. J., Dullinger, S., ... & Field, R. (2020). Snapshot isolation and isolation history challenge the analogy between mountains and islands used to understand endemism. *Global Ecology and Biogeography*, 29(10), 1651-1673.
- Gaston, K. J. (2003). *The structure and dynamics of geographic ranges*. Oxford University Press on Demand.
- Gatti, R. C., Callaghan, T., Velichevskaya, A., Dudko, A., Fabbio, L., Battipaglia, G., & Liang, J. (2019). Accelerating upward treeline shift in the Altai Mountains under last-century climate change. *Scientific reports*, 9(1), 1-13.

- Geldmann, J., Barnes, M., Coad, L., Craigie, I. D., Hockings, M., & Burgess, N. D. (2013). Effectiveness of terrestrial protected areas in reducing habitat loss and population declines. *Biological Conservation*, 161, 230-238.
- Gentili, R., Baroni, C., Panigada, C., Rossini, M., Tagliabue, G., Armiraglio, S., ... & Salvatore, M. C. (2020). Glacier shrinkage and slope processes create habitat at high elevation and microrefugia across treeline for alpine plants during warm stages. *Catena*, 193, 104626.
- Gholizadeh, H., Naqinezhad, A., & Chytrý, M. (2020). Classification of the Hyrcanian forest vegetation, Northern Iran. *Applied Vegetation Science*, 23(1), 107-126.
- Guisan, A., & Thuiller, W. (2005). Predicting species distribution: offering more than simple habitat models. *Ecology letters*, 8(9), 993-1009.
- Guisan, A., Tingley, R., Baumgartner, J. B., Naujokaitis-Lewis, I., Sutcliffe, P. R., Tulloch, A. I., ... & Buckley, Y. M. (2013). Predicting species distributions for conservation decisions. *Ecology letters*, 16(12), 1424-1435.
- Habel, J. C., Rödder, D., Schmitt, T., Gros, P., & Ulrich, W. (2021). Climate change drives mountain butterflies towards the summits. *Scientific Reports*.
- Haeblerli, W., Hoelzle, M., Paul, F., & Zemp, M. (2007). Integrated monitoring of mountain glaciers as key indicators of global climate change: the European Alps. *Annals of glaciology*, 46, 150-160.
- Haidarian, M., Tamartash, R., Jafarian-Jeloudar, Z., Tarkesh, M., & Tataian, M. R. (2021). The Effects of Climate Change on the Future Distribution of Astragalus adscendens in Central Zagros, Iran. *Journal of Rangeland Science*, 11(2), 152-170.
- Hajima, T., Watanabe, M., Yamamoto, A., Tatebe, H., Noguchi, M. A., Abe, M., ... & Kawamiya, M. (2020). Development of the MIROC-ES2L Earth system model and the evaluation of biogeochemical processes and feedbacks. *Geoscientific Model Development*, 13(5), 2197-2244.
- Hälfors, M. H., Antão, L. H., Itter, M., Lehikoinen, A., Lindholm, T., Roslin, T., & Saastamoinen, M. (2020). Shifts in timing and duration of breeding for 73 boreal bird species over four decades. *Proceedings of the National Academy of Sciences*, 117(31), 18557-18565.
- Harrison, S., & Noss, R. (2017). Endemism hotspots are linked to stable climatic refugia. *Annals of Botany*, 119(2), 207-214.
- Hochkirch, A., Samways, M. J., Gerlach, J., Böhm, M., Williams, P., Cardoso, P., ... & Dijkstra, K. D. B. (2021). A strategy for the next decade to address data deficiency in neglected biodiversity. *Conservation Biology*, 35(2), 502-509.
- Hoffmann, S., Irl, S. D., & Beierkuhnlein, C. (2019). Predicted climate shifts within terrestrial protected areas worldwide. *Nature communications*, 10(1), 1-10.
- IPCC, 2019: Climate Change and Land: an IPCC special report on climate change, desertification, land degradation, sustainable land management, food security, and greenhouse gas fluxes in terrestrial ecosystems
- Jarvis, A. (2008). Hole-field seamless SRTM data, International Centre for Tropical Agriculture (CIAT). <http://srtm.csi.cgiar.org>.

- Keppel, G., Mokany, K., Wardell-Johnson, G. W., Phillips, B. L., Welbergen, J. A., & Reside, A. E. (2015). The capacity of refugia for conservation planning under climate change. *Frontiers in Ecology and the Environment*, 13(2), 106-112.
- Kolahî, M., Sakai, T., Moriya, K., Makhdoum, M. F., & Koyama, L. (2013). Assessment of the effectiveness of protected areas management in Iran: Case study in Khojir National Park. *Environmental management*, 52(2), 514-530.
- Konapala, G., Mishra, A. K., Wada, Y., & Mann, M. E. (2020). Climate change will affect global water availability through compounding changes in seasonal precipitation and evaporation. *Nature Communications*, 11(1), 1-10.
- Körner, C., & Hiltbrunner, E. (2018). The 90 ways to describe plant temperature. *Perspectives in Plant Ecology, Evolution and Systematics*, 30, 16-21.
- Lawler, J. J. et al. (2015) 'The theory behind, and the challenges of, conserving nature's stage in a time of rapid change', *Conservation Biology*, 29(3), pp. 618–629. doi: 10.1111/cobi.12505.
- Malakoutikhah, S., Fakheran, S., Hemami, M. R., Tarkesh, M., & Senn, J. (2018). Altitudinal heterogeneity and vulnerability assessment of protected area network for climate change adaptation planning in central Iran. *Applied geography*, 92, 94-103.
- Manabe, S. (2019). Role of greenhouse gas in climate change. *Tellus A: Dynamic Meteorology and Oceanography*, 71(1), 1620078.
- Manes, S., Costello, M. J., Beckett, H., Debnath, A., Devenish-Nelson, E., Grey, K. A., ... & Vale, M. M. (2021). Endemism increases species' climate change risk in areas of global biodiversity importance. *Biological Conservation*, 109070.
- McLaughlin, B. C., Ackerly, D. D., Klos, P. Z., Natali, J., Dawson, T. E., & Thompson, S. E. (2017). Hydrologic refugia, plants, and climate change. *Global change biology*, 23(8), 2941-2961.
- McSweeney, C. F., Jones, R. G., Lee, R. W., & Rowell, D. P. (2015). Selecting CMIP5 GCMs for downscaling over multiple regions. *Climate Dynamics*, 44(11), 3237-3260.
- Menzel, A., Sparks, T. H., Estrella, N., Koch, E., Aasa, A., Ahas, R., ... & Zust, A. N. A. (2006). European phenological response to climate change matches the warming pattern. *Global change biology*, 12(10), 1969-1976.
- Mittermeier, R. A., Turner, W. R., Larsen, F. W., Brooks, T. M., & Gascon, C. (2011). Global biodiversity conservation: the critical role of hotspots. In *Biodiversity hotspots* (pp. 3-22). Springer, Berlin, Heidelberg.
- Moss, R. H., Edmonds, J. A., Hibbard, K. A., Manning, M. R., Rose, S. K., Van Vuuren, D. P., ... & Wilbanks, T. J. (2010). The next generation of scenarios for climate change research and assessment. *Nature*, 463(7282), 747-756.
- Noroozi, J., Akhani, H., & Breckle, S. W. (2008). Biodiversity and phytogeography of the alpine flora of Iran. *Biodiversity and Conservation*, 17(3), 493-521.
- Noroozi, J., Talebi, A., Doostmohammadi, M., Rumpf, S. B., Linder, H. P., & Schneeweiss, G. M. (2018). Hotspots within a global biodiversity hotspot-areas of endemism are associated with high mountain ranges. *Scientific reports*, 8(1), 1-10.

- Norozi, J., Talebi, A., Doostmohammadi, M., Manafzadeh, S., Asgarpour, Z., & Schneeweiss, G. M. (2019a). Endemic diversity and distribution of the Iranian vascular flora across phytogeographical regions, biodiversity hotspots and areas of endemism. *Scientific reports*, 9(1), 1-12.
- Norozi, J., Naqinezhad, A., Talebi, A., Doostmohammadi, M., Plutzar, C., Rumpf, S. B., ... & Schneeweiss, G. M. (2019b). Hotspots of vascular plant endemism in a global biodiversity hotspot in Southwest Asia suffer from significant conservation gaps. *Biological Conservation*, 237, 299-307.
- Norozi, J. (Ed.). (2020). *Plant biogeography and vegetation of high mountains of Central and South-West Asia*. Springer International Publishing.
- Parks, S. A., Carroll, C., Dobrowski, S. Z., & Allred, B. W. (2020). Human land uses reduce climate connectivity across North America. *Global change biology*, 26(5), 2944-2955.
- Parmesan, C., Ryrholm, N., Stefanescu, C., Hill, J. K., Thomas, C. D., Descimon, H., ... & Warren, M. (1999). Poleward shifts in geographical ranges of butterfly species associated with regional warming. *Nature*, 399(6736), 579-583.
- Parmesan, C., & Yohe, G. (2003). A globally coherent fingerprint of climate change impacts across natural systems. *Nature*, 421(6918), 37-42.
- Pascual, U., Adams, W. M., Díaz, S., Lele, S., Mace, G. M., & Turnhout, E. (2021). Biodiversity and the challenge of pluralism. *Nature Sustainability*, 1-6.
- Perret, D. L., Leslie, A. B., & Sax, D. F. (2019). Naturalized distributions show that climatic disequilibrium is structured by niche size in pines (*Pinus L.*). *Global Ecology and Biogeography*, 28(4), 429-441.
- Potter, K. A., Arthur Woods, H., & Pincebourde, S. (2013). Microclimatic challenges in global change biology. *Global change biology*, 19(10), 2932-2939.
- van Proosdij, A. S., Sosef, M. S., Wieringa, J. J., & Raes, N. (2016). Minimum required number of specimen records to develop accurate species distribution models. *Ecography*, 39(6), 542-552.
- R Core Team (2017) *R: A Language and Environment for Statistical Computing*. Vienna, Austria: R 551 Foundation for Statistical Computing.
- Rada, S., Schweiger, O., Harpke, A., Kühn, E., Kuras, T., Settele, J., & Musche, M. (2019). Protected areas do not mitigate biodiversity declines: A case study on butterflies. *Diversity and Distributions*, 25(2), 217-224.
- Renner, S. S., & Zohner, C. M. (2018). Climate change and phenological mismatch in trophic interactions among plants, insects, and vertebrates. *Annual review of ecology, evolution, and systematics*, 49, 165-182.
- Rodríguez-Iturbe, I., & Porporato, A. (2007). *Ecohydrology of water-controlled ecosystems: soil moisture and plant dynamics*. Cambridge University Press.
- Rumpf, S. B., Hülber, K., Zimmermann, N. E., & Dullinger, S. (2019). Elevational rear edges shifted at least as much as leading edges over the last century. *Global Ecology and Biogeography*, 28(4), 533-543.
- Saastamoinen, M., Bocedi, G., Cote, J., Legrand, D., Guillaume, F., Wheat, C. W., ... & del Mar Delgado, M. (2018). Genetics of dispersal. *Biological Reviews*, 93(1), 574-599.

- Safaei, M., Rezayan, H., Firouzabadi, P. Z., & Sadidi, J. (2021). Optimization of species distribution models using a genetic algorithm for simulating climate change effects on Zagros forests in Iran. *Ecological Informatics*, 63, 101288.
- Sandel, B., Monnet, A. C., Govaerts, R., & Vorontsova, M. (2017). Late Quaternary climate stability and the origins and future of global grass endemism. *Annals of Botany*, 119(2), 279-288.
- Séférian, R., Nabat, P., Michou, M., Saint-Martin, D., Voldoire, A., Colin, J., ... & Madec, G. (2019). Evaluation of CNRM earth system model, CNRM-ESM2-1: Role of earth system processes in present-day and future climate. *Journal of Advances in Modeling Earth Systems*, 11(12), 4182-4227.
- Semenchuk P., Moser D., Essl F., Schindler S., Wessely J., Gattringer A., Dullinger S. (2021) Manuscript submitted for publication
- Shukla, P. R., Skea, J., Calvo Buendia, E., Masson-Delmotte, V., Pörtner, H. O., Roberts, D. C., ... & Malley, J. (2019). IPCC, 2019: Climate Change and Land: an IPCC special report on climate change, desertification, land degradation, sustainable land management, food security, and greenhouse gas fluxes in terrestrial ecosystems.
- Sillmann, J., & Roeckner, E. (2008). Indices for extreme events in projections of anthropogenic climate change. *Climatic Change*, 86(1), 83-104.
- Smithson, M., & Verkuilen, J. (2006). A better lemon squeezer? Maximum-likelihood regression with beta-distributed dependent variables. *Psychological methods*, 11(1), 54.
- Smolik, M., Dullinger, S., Essl, F., Kleinbauer, I., Leitner, M., Peterseil, J., ... & Vogl, G. (2010). Integrating species distribution models and interacting particle systems to predict the spread of an invasive alien plant. *Journal of Biogeography*, 37(3), 411-422.
- Soofi, M., Ghoddousi, A., Zeppenfeld, T., Shokri, S., Soufi, M., Jafari, A., ... & Waltert, M. (2018). Livestock grazing in protected areas and its effects on large mammals in the Hyrcanian forest, Iran. *Biological Conservation*, 217, 377-382.
- Steinbauer, M. J., Grytnes, J. A., Jurasiński, G., Kulonen, A., Lenoir, J., Pauli, H., ... & Wipf, S. (2018). Accelerated increase in plant species richness on mountain summits is linked to warming. *Nature*, 556(7700), 231-234.
- Stephens, L., Fuller, D., Boivin, N., Rick, T., Gauthier, N., Kay, A., ... & Ellis, E. (2019). Archaeological assessment reveals Earth's early transformation through land use. *Science*, 365(6456), 897-902.
- Thomas, C. D. and Gillingham, P. K. (2015) 'The performance of protected areas for biodiversity under climate change', *Biological Journal of the Linnean Society*, 115(3), pp. 718–730. doi: 10.1111/bij.12510.
- Townsend Peterson, A., & Watson, D. M. (1998). Problems with areal definitions of endemism: the effects of spatial scaling. *Diversity and Distributions*, 4(4), 189-194.
- Trew, B. T., & Maclean, I. M. (2021). Vulnerability of global biodiversity hotspots to climate change. *Global Ecology and Biogeography*, 30(4), 768-783.
- Tribsch, A. (2004). Areas of endemism of vascular plants in the Eastern Alps in relation to Pleistocene glaciation. *Journal of Biogeography*, 31(5), 747-760.
- Tuckett, R. (2019). Greenhouse gases. In *Encyclopedia of Analytical Science* (pp. 362-372). Elsevier.

Ward, M., Saura, S., Williams, B., Ramírez-Delgado, J. P., Arafah-Dalmau, N., Allan, J. R., ... & Watson, J. E. (2020). Just ten percent of the global terrestrial protected area network is structurally connected via intact land. *Nature communications*, 11(1), 1-10.

Wasof, S., Lenoir, J., Gallet-Moron, E., Jamoneau, A., Brunet, J., Cousins, S. A., ... & Decocq, G. (2013). Ecological niche shifts of understorey plants along a latitudinal gradient of temperate forests in north-western Europe. *Global Ecology and Biogeography*, 22(10), 1130-1140.

Wessely, J., Hülber, K., Gattringer, A., Kuttner, M., Moser, D., Rabitsch, W., ... & Essl, F. (2017). Habitat-based conservation strategies cannot compensate for climate-change-induced range loss. *Nature Climate Change*, 7(11), 823-827.

While, G. M., & Uller, T. (2014). Quo vadis amphibia? Global warming and breeding phenology in frogs, toads and salamanders. *Ecography*, 37(10), 921-929.

Williams, J. W., Ordonez, A., & Svenning, J. C. (2021). A unifying framework for studying and managing climate-driven rates of ecological change. *Nature Ecology & Evolution*, 5(1), 17-26.

Valavi, R., Shafizadeh-Moghadam, H., Matkan, A., Shakiba, A., Mirbagheri, B., & Kia, S. H. (2019). Modelling climate change effects on Zagros forests in Iran using individual and ensemble forecasting approaches. *Theoretical and Applied Climatology*, 137(1), 1015-1025.

Zhang, R. H., Yu, Y., Song, Z., Ren, H. L., Tang, Y., Qiao, F., ... & Wang, L. (2020). A review of progress in coupled ocean-atmosphere model developments for ENSO studies in China. *Journal of Oceanology and Limnology*, 38(4), 930-961.

Appendix

List of tables

Table A1: Species included in the study with information on elevational distribution.....30

Table of figures

Figure 1: An example for the derivation of the potential area of occupancy for *Acantholimon autostrianicus*.....10

Figure 2: Bivariate density kernel (= climatic niche) of *Cousina calocephala* and the climatic space in the associated PA's under current climate.....11

Figure 3: Climatic representation within associated PAs across all endemic species analysed.....13

Figure 4: Representation of species' climatic niches within PAs in relation to mean elevation.....14

Figure 5: Climatic representation within associated PAs for widespread and range-restricted species separately.....15

Figure A1: The study area.....28

Figure A2: Mean annual precipitation (mm) of each GCM.....28

Figure A3: Relative change in climatic representation in associated PAs for each species.....29

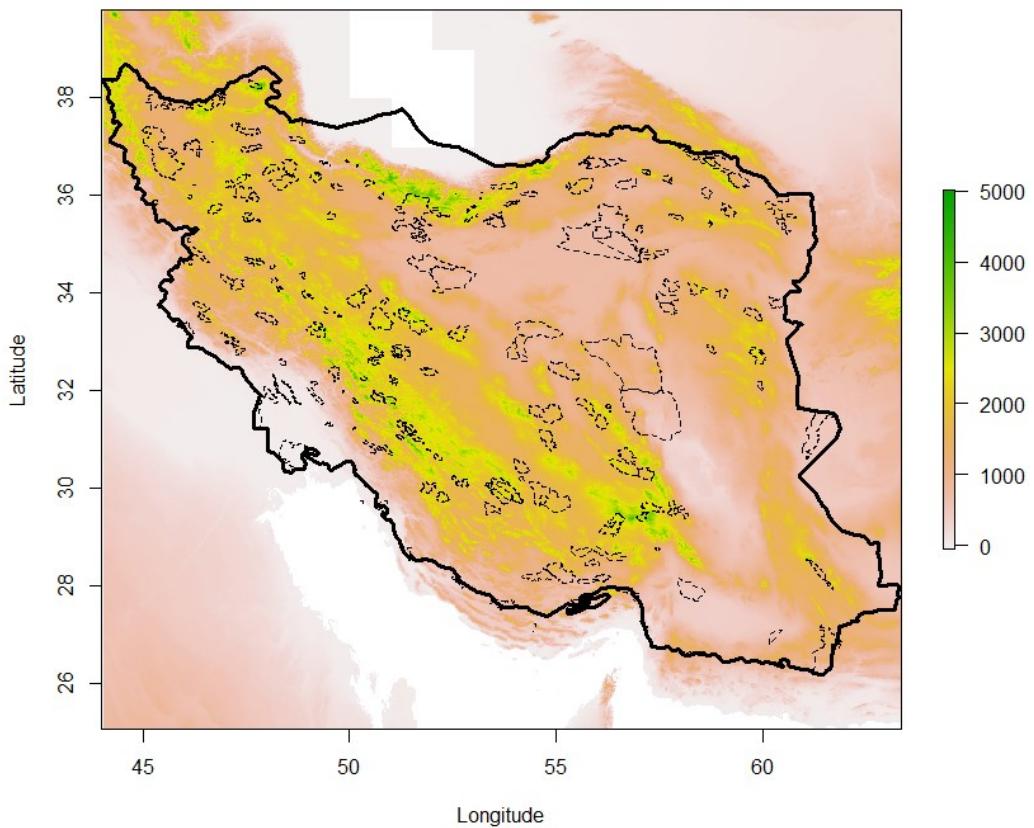


Fig. A1: Borders of Iran (solid line) and the Protected Area network (dashed lines). Background map is a digital elevation model (DEM).

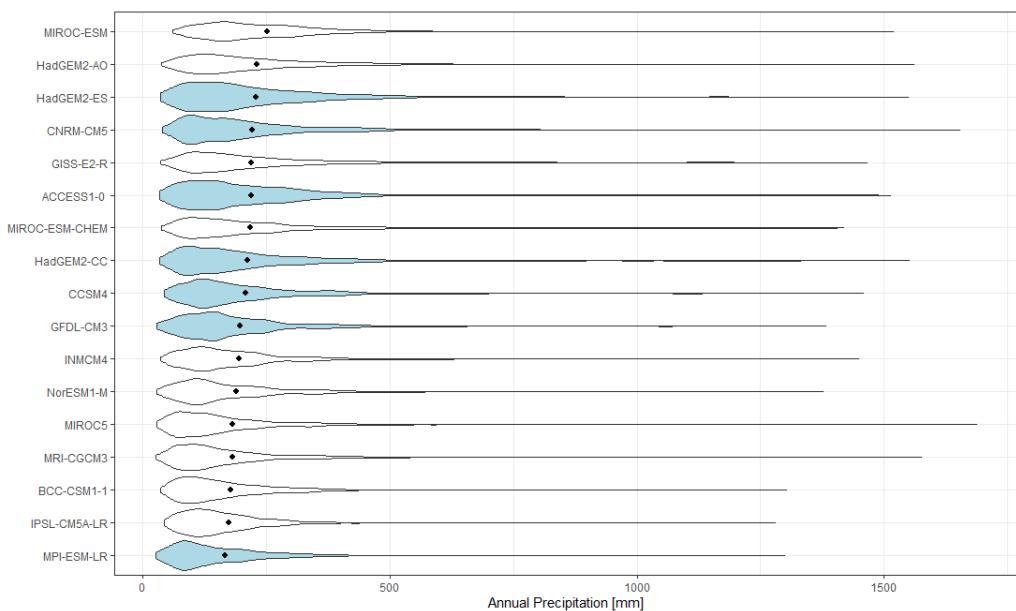


Fig. A2: Mean annual precipitation (mm) predicted by the GCMs of CMIP5 with 30" resolution which are available on WorldClim (www.worldclim.org) for the year 2070 under the climate scenario RCP 8.5. Coloured GCMs indicate good performance in McSweeney et al., 2015. Mean annual precipitation is shown on the y-axis, common abbreviations of GCMs on the x-axis.

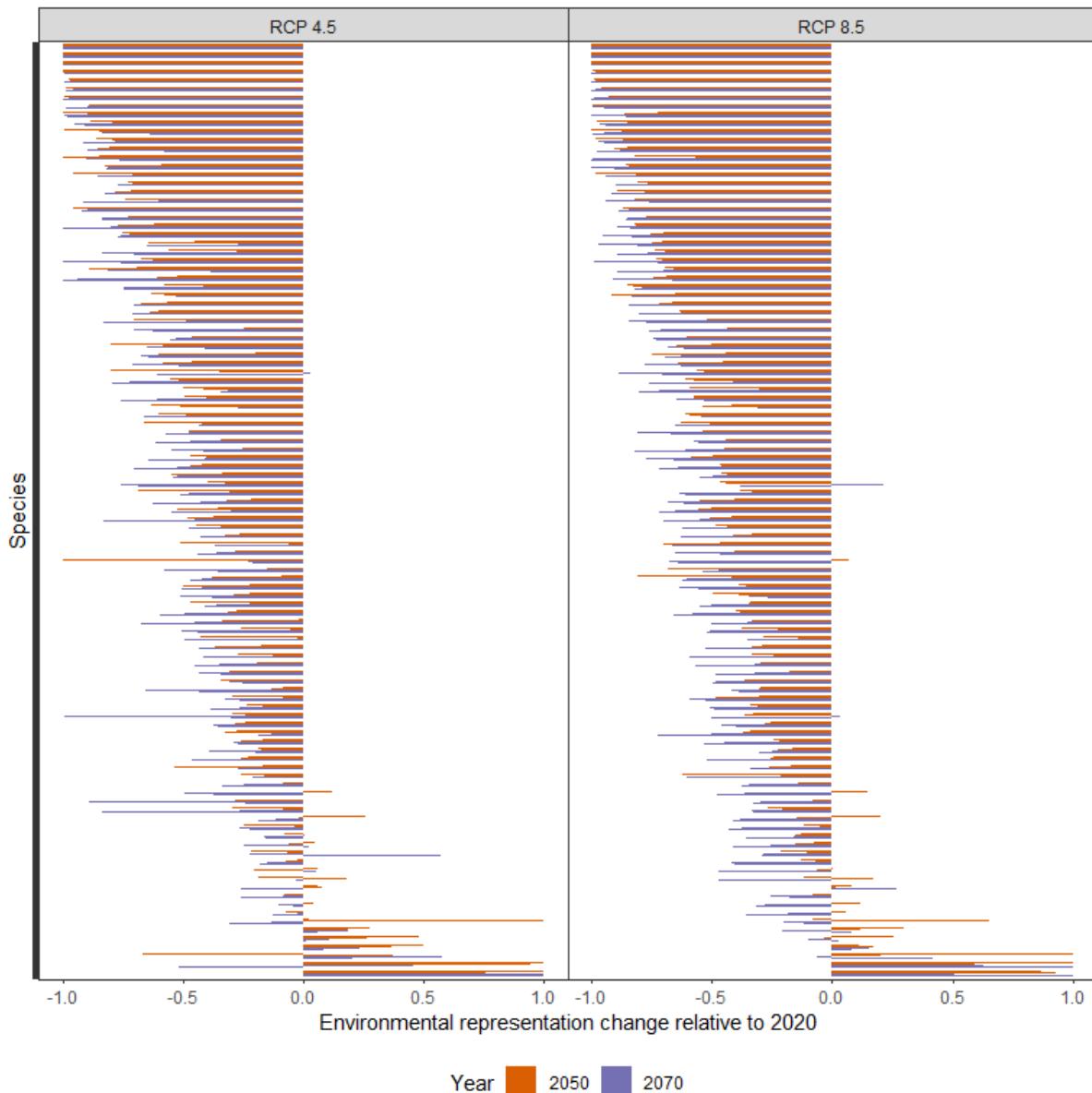


Fig. A3: Relative change in climatic representation in associated PAs for each species ($n= 1192$) separately. Shown is the difference between 2020 and 2070 for RCP4.5 (left panel) and RCP8.5 (right panel). Relative change values are displayed on the x-axis and individual species on the y-axis (species names omitted). Values range between -1 (complete loss of climatic representation) and 1 (doubling of climatic representation). Species with values higher than 1 were set to 1 for clearer display.

Table A1: Species with sufficient elevational range (>800m) to be included in the study. Max and min describe highest and lowest elevation of documented records per species. Elev. range is the elevational range resulting from max. – min.

Species	min.	max.	elev. range
<i>Acantholimon acmostegium</i> Boiss. & Buhse	800	2600	1800
<i>Acantholimon aspadanum</i> Bunge	1200	2882	1682
<i>Acantholimon asphodelinum</i> Mobayen	1611	3000	1389
<i>Acantholimon atropatanum</i> Bunge	1300	2200	900
<i>Acantholimon austro-iranicus</i> Rech.f. & Schiman-Czeika	900	1900	1000
<i>Acantholimon bodeanum</i> Bunge	1250	3000	1750
<i>Acantholimon brachystachyum</i> Boiss. ex Bunge	2250	3500	1250
<i>Acantholimon bromifolium</i> Boiss. ex Bunge	1620	2800	1180
<i>Acantholimon cephalotoides</i> Rech.f. in DC.	2100	3750	1650
<i>Acantholimon chlorostegium</i> Rech.f. & Schiman-Czeika	2000	3661	1661
<i>Acantholimon collare</i> Koeie & Rech.f.	1100	2438	1338
<i>Acantholimon curviflorum</i> Bunge	850	1920	1070
<i>Acantholimon cymosum</i> Bunge	1100	2500	1400
<i>Acantholimon demavendicum</i> Bornm.	3300	4200	900
<i>Acantholimon eschkerense</i> Boiss. & Hausskn.	1850	3300	1450
<i>Acantholimon festucaceum</i> (Jaub. & Spach) Boiss.	1100	3300	2200
<i>Acantholimon flabellum</i> Assadi	1500	2500	1000
<i>Acantholimon flexuosum</i> Boiss. & Hausskn. ex Bunge	1600	3200	1600
<i>Acantholimon gilliati</i> Turrill	500	2700	2200
<i>Acantholimon glabratum</i> Assadi	2100	2950	850
<i>Acantholimon incomptum</i> Boiss. & Buhse	1210	2700	1490
<i>Acantholimon kermanense</i> Assadi & Mirtadz.	2601.86	3440.05	838.19
<i>Acantholimon melananthum</i> Boiss.	2500	3900	1400
<i>Acantholimon modestum</i> Bornm. ex Rech.f. & Schiman-Czeika	1836	4012	2176
<i>Acantholimon nigricans</i> Mobayen	2480	3900	1420
<i>Acantholimon oliganthum</i> Boiss.	1600	3507	1907
<i>Acantholimon olivieri</i> (Jaub. & Spach) Boiss.	1290	3100	1810
<i>Acantholimon ophiocladus</i> Rech.f. & Schiman-Czeika	1652	3000	1348
<i>Acantholimon pterostegium</i> Bunge	1000	2500	1500
<i>Acantholimon quinquelobum</i> Bunge	1000	1900	900
<i>Acantholimon restiaceum</i> Bunge	900	1975	1075
<i>Acantholimon rhodopolius</i> Rech.f. & Schiman-Czeika	1420	2350	930
<i>Acantholimon scabrellum</i> Boiss. & Hausskn.	2000	4100	2100
<i>Acantholimon schahrudicum</i> Bunge	1060	2600	1540
<i>Acantholimon scirpinum</i> Bunge	1200	3500	2300
<i>Acantholimon scorpius</i> (Jaub. & Spach) Boiss.	600	2700	2100
<i>Acantholimon serotinus</i> Rech. f. & Schiman-Czeika	1408	2857	1449
<i>Acantholimon senganense</i> Bunge	1600	3000	1400
<i>Acantholimon sirchense</i> Assadi & Mirtadzadini	2007.12	3200	1192.88
<i>Acantholimon spinicalyx</i> Koeie & Rech.f.	1150	2465	1315
<i>Acantholimon talagonicum</i> Boiss. in DC.	1250	3000	1750
<i>Acantholimon tomentellum</i> Boiss.	3000	3900	900
<i>Acantholimon tragacanthinum</i> (Jaub. & Spach) Boiss.	1300	2600	1300

<i>Acantholimon wendelboi</i> Rech.f. & Schiman-Czeika	1509	2800	1291
<i>Acantholimon zaeifii</i> Assadi	1800	3551	1751
<i>Acanthophyllum crassifolium</i> Boiss.	1120	3500	2380
<i>Acanthophyllum leucostegium</i> Schiman-Czeika	650	2300	1650
<i>Acanthophyllum pachycephalum</i> Schiman-Czeika	1160	2033	873
<i>Achillea aucheri</i> Boiss.	2500	4220	1720
<i>Achillea eriophora</i> DC.	746	2850	2104
<i>Achillea oxydonta</i> Boiss.	1700	2900	1200
<i>Achillea pachycephala</i> Rech.f.	1200	3000	1800
<i>Achillea talagonica</i> Boiss.	1300	3000	1700
<i>Aethionema semnanensis</i> Mozaffarian	1900	3300	1400
<i>Aethionema stenopterum</i> Boiss.	1600	3352.05	1752.05
<i>Aethionema umbellatum</i> (Boiss.) Bornm.	3300	4200	900
<i>Agropyron brachyphyllum</i> Boiss. & Hausskn. ex Boiss.	2110	3300	1190
<i>Ajuga chamaecistus</i> Ging. ex Benth.	1000	3000	2000
<i>Albraunia foveopilosa</i> Speta	100	1500	1400
<i>Alcea arbelensis</i> Boiss. & Hausskn.	500	2900	2400
<i>Alcea glabrata</i> Alef.	700	2800	2100
<i>Alcea gorganica</i> (Rech.f.	100	2050	1950
<i>Alcea koelzii</i> I.Riedl	1020	2800	1780
<i>Alcea kurdica</i> (Schlecht.) Alef.	1000	2200	1200
<i>Alcea lenkoranica</i> Iljin	20	1600	1580
<i>Alcea loftusii</i> (Baker) Zohary	300	2000	1700
<i>Alcea schirazana</i> Alef	889	2850	1961
<i>Alcea tarica</i> Pakravan & Ghahreman	1000	2900	1900
<i>Alcea wilhelminae</i> I. Riedl	250	2500	2250
<i>Alchemilla amardica</i> Rothm.	2000	2900	900
<i>Alchemilla citrina</i> S. E. Fr'hner	2000	3682.01	1682.01
<i>Alchemilla fluminea</i> S. E. Fr'hner	1800	3750	1950
<i>Alchemilla melancholica</i> S. E. Fr'hner	1400	2950	1550
<i>Alchemilla microscopica</i> S. E. Fr'hner	1302.05	2200	897.95
<i>Alchemilla pectiniloba</i> S. E. Fr'hner	2000	3200	1200
<i>Alchemilla plicatissima</i> S. E. Fr'hner	1302.05	2600	1297.95
<i>Alchemilla surculosa</i> S. E. Fr'hner	1800	3000	1200
<i>Alkanna bracteosa</i> Boiss.	1500	3000	1500
<i>Alkanna frigida</i> Boiss.	1300	3400	2100
<i>Allium akaka</i> S.G. Gmel. ex Schult. & Schult.f.	1300	2810.97	1510.97
<i>Allium aladaghense</i> Memariani & Joharchi	1700	2800	1100
<i>Allium alamutense</i> Razyfard	1250	3100	1850
<i>Allium austroiranicum</i> R.M. Fritsch	1700	3300	1600
<i>Allium bakhtiaricum</i> Regel	1700	3000	1300
<i>Allium brachyodon</i> Boiss.	1441.96	3835.01	2393.05
<i>Allium breviscapum</i> Stapf	1875.08	3202.02	1326.94
<i>Allium capitellatum</i> Boiss.	3100	3900	800
<i>Allium caspium</i> (Pall.) M. Bieb.	500	1900	1400
<i>Allium cathodicarpum</i> Wendelbo	1537.32	3800	2262.68

Allium chrysantherum Boiss. & Reut.	1500	2300	800
Allium cristophii Trautv.	100	2200	2100
Allium derderianum Regel	1200	3400	2200
Allium egorovae M.V. Agab. & Ogan.	1200	3000	1800
Allium elburzense Wendelbo	1600	3500	1900
Allium ellisii Hook.f.	1000	2400	1400
Allium fedtschenkoi N b lek	1509	3068.74	1559.74
Allium grande Lipsky	650	3100	2450
Allium graveolens (R.M. Fritsch) R.M. Fritsch	950	3000	2050
Allium haemanthoides Boiss. & Reut. ex Regel	1400	3700	2300
Allium helicophyllum Vved.	1200	2500	1300
Allium jesdianum Boiss. & Buhse	1550	3420	1870
Allium kazerouni Parsa	1660	2860	1200
Allium keusgenii R.M. Fritsch	1450	2500	1050
Allium koelzii (Wendelbo) K. Perss. & Wendelbo	1312.24	3108.61	1796.37
Allium kuhsorkhense R.M. Fritsch & Joharchi	900	2500	1600
Allium lalesaricum Freyn & Bornm.	3200	4000	800
Allium latifolium Jaub. & Spach	1384.69	3000	1615.31
Allium longivaginatum Wendelbo	1800	4140	2340
Allium materculae Bordz.	950	2400	1450
Allium minutiflorum Regel	1545.02	2431.88	886.86
Allium moderense R.M. Fritsch	1930	2880	950
Allium monophyllum Vved.	1527.98	2697.02	1169.04
Allium montelburzense R.M. Fritsch	2280	3620	1340
Allium orientiranicum Neshati	1300	2300	1000
Allium pseudobodeanum R.M. Fritsch & Matin	1422.89	2600	1177.11
Allium pseudohollandicum R.M. Fritsch	1400	2700	1300
Allium remediorum (R.M. Fritsch) R.M. Fritsch	1454.95	2800	1345.05
Allium sabalense R.M. Fritsch	1800	3000	1200
Allium sahandicum R.M. Fritsch	1350	3000	1650
Allium saralicum R.M. Fritsch	800	2221.85	1421.85
Allium scotostemon Wendelbo	950	3400	2450
Allium shelkovnikovii Grossh.	800	3200	2400
Allium subakaka Razyfard & Zarre	1430	2900	1470
Allium ubipetrense R.M. Fritsch	1100	3100	2000
Allium zagricum R.M. Fritsch	1454.95	2795.33	1340.38
Alnus dolichocarpa Zare	-14	1200	1214
Alrawia bellii (Baker) K. Persson & Wendelbo	1100	2800	1700
Alyssum bracteatum Boiss. & Buhse	1000	2023	1023
Alyssum lanigerum DC.	1400	2800	1400
Alyssum mozaffarianii Kavousi	2000	2900	900
Alyssum polycladum Rech. F.	1650	4000	2350
Amygdalus eburnea Spach	200	2614	2414
Amygdalus elaeagnifolia Spach	1300	3500	2200
Amygdalus haussknechtii (C.K.Schneider.) Bornm.	1330	3100	1770
Anabasis calcarea (Charif & Aellen) Bokhari & Wendelbo	850	2250	1400

<i>Anabasis haussknechtii</i> Bunge ex Boiss.	1065	2502	1437
<i>Anchonium elichrysifolium</i> (DC.) Boiss.	2100	3800	1700
<i>Andrachne fruticulosa</i> Boiss.	1000	2000	1000
<i>Anthemis atropatana</i> Iranshahr	760	2000	1240
<i>Anthemis austroiranica</i> Rech.f. Aell. & Esfand.	1	2127	2126
<i>Anthemis brachystephana</i> Bornm. & Gauba	909	2060	1151
<i>Anthemis gayana</i> Boiss.	900	2650	1750
<i>Anthemis gilanica</i> Bornm. & Gauba	246	2200	1954
<i>Anthemis lorestanica</i> Iranshahr	251	2060	1809
<i>Anthemis moghanica</i> Iranshahr	80	2702	2622
<i>Anthemis persica</i> Boiss.	234	1900	1666
<i>Anthemis schizostephana</i> Boiss. & Hausskn.	300	2028	1728
<i>Anthemis susiana</i> NAB.	18	889	871
<i>Anthemis talyschensis</i> A. Fedor.	1250	2084	834
<i>Anthochlamys multinervis</i> Rech. f.	795	1800	1005
<i>Aphanopleura breviseta</i> (Boiss.) Heywood & Jruy	750	2500	1750
<i>Arabis ottonis-schulzii</i> Bornm. & Gauba	1450	2700	1250
<i>Arabis rimarum</i> Rech. F.	2200	3200	1000
<i>Arenaria persica</i> Boiss.	2500	3850	1350
<i>Arenaria polycnemifolia</i> Boiss.	1500	3250	1750
<i>Arenaria szowitsii</i> Boiss.	1385	2300	915
<i>Arenaria zargariana</i> Parsa	1190	2500	1310
<i>Argyrolobium trigonelloides</i> Jaub. & Spach	122	2300	2178
<i>Aristolochia hyrcana</i> Davis & M.S.Khan	200	1700	1500
<i>Aristolochia olivieri</i> Collengo in Boiss.	700	2500	1800
<i>Artemisia kermanensis</i> Podl.	1254	2475	1221
<i>Artemisia khorassanica</i> Podl.	700	2258	1558
<i>Artemisia melanolepis</i> Boiss.	3000	4000	1000
<i>Artemisia splendens</i> Willd.	2200	3700	1500
<i>Arum giganteum</i> Ghebreman	1275	2651	1376
<i>Asparagus touranensis</i> Hamdi & Assadi	36	1724	1688
<i>Asperula fragillima</i> Boiss. & Hausskn. ex Boiss.	1800	3610.94	1810.94
<i>Asperula gorganica</i> Schonb.-Tem. & Ehrend.	600	2080	1480
<i>Asperula mazanderanica</i> Ehrend.	950	2700	1750
<i>Asperula microphylla</i> Boiss.	300	2500	2200
<i>Asperula rechingeri</i> Ehrend. & Schonb.- Tem	2000	3261.67	1261.67
<i>Asperula seticornis</i> Boiss.	1900	3000	1100
<i>Astragalus abnormalis</i> Rech.f.	1125	2489	1364
<i>Astragalus ahmad-parsae</i> Maassoumi	1500	2350	850
<i>Astragalus albispinus</i>	1715	2650	935
<i>Astragalus alyssiformis</i> Maassoumi	1875	3000	1125
<i>Astragalus angurianensis</i> Podlech & Maassoumi	1950	2800	850
<i>Astragalus angustistipulatus</i> Podlech	2800	4000	1200
<i>Astragalus anserinaefolius</i> Boiss.	100	2230	2130
<i>Astragalus argyrostachys</i> Boiss.	1650	2500	850
<i>Astragalus askius</i> Bunge	1370	3800	2430

<i>Astragalus aspadanus</i> Bunge	1010	3013	2003
<i>Astragalus atricapillus</i> Bornm.	2900	3800	900
<i>Astragalus avicennicus</i> Parsa	1730	3200	1470
<i>Astragalus baba-alliar</i> Parsa	139	2130	1991
<i>Astragalus babakhanloui</i> Maassoumi & Podl.	1500	3600	2100
<i>Astragalus bazarganii</i> Podlech & Zarre	867	1919	1052
<i>Astragalus bazmanicus</i> Podlech	2000	3000	1000
<i>Astragalus beckii</i> Bornm.	1800	3700	1900
<i>Astragalus belgheisicoides</i> Podlech & Maassoumi	1900	2900	1000
<i>Astragalus belgheisicus</i> Maassoumi	600	2900	2300
<i>Astragalus birangae</i> Maassoumi	160	1650	1490
<i>Astragalus biserrula</i> Bunge	850	2000	1150
<i>Astragalus bodeanus</i> Fisch.	1600	2705	1105
<i>Astragalus bojnurdensis</i> Podlech	1082	1938	856
<i>Astragalus borujenensis</i> Ranjbar & Maassoumi	1100	1950	850
<i>Astragalus bounophilus</i> Boiss. & Hohen.	1567.22	3241.68	1674.46
<i>Astragalus brachyodontus</i> Boiss.	200	2600	2400
<i>Astragalus bradosticus</i> Maassoumi & Podlech	700	2700	2000
<i>Astragalus brevirhachis</i> Tietz & Zarre	1800	2700	900
<i>Astragalus brunianus</i> Bornm.	1350	2200	850
<i>Astragalus callainus</i> Podlech	938	2100	1162
<i>Astragalus calliphysa</i> Bunge	750	2708	1958
<i>Astragalus campylanthus</i> Boiss.	1100	3114	2014
<i>Astragalus capax</i> Maassoumi	1500	3800	2300
<i>Astragalus carmanicus</i> Bornm.	2000	3700	1700
<i>Astragalus catacamptus</i> Bunge	800	2577	1777
<i>Astragalus cemerinus</i> Beck	1500	2500	1000
<i>Astragalus cephalanthus</i> DC.	1100	3400	2300
<i>Astragalus chalaranthus</i> Boiss. & Hausskn. in Boiss.	2148	3200	1052
<i>Astragalus chartostegius</i> Boiss. & Hausskn.	2300	4000	1700
<i>Astragalus chrysanthus</i> Boiss. & Hohen	2100	3532.35	1432.35
<i>Astragalus chrysotrichus</i> Boiss.	1200	3081	1881
<i>Astragalus circumlacustris</i> Podlech & Sytin	1300	2122	822
<i>Astragalus clivicola</i> Podlech & Maassoumi	1500	2350	850
<i>Astragalus confusus</i> Bunge.	2300	3500	1200
<i>Astragalus curviflorus</i> Boiss.	200	2700	2500
<i>Astragalus cyclophyllon</i> Beck	1000	3000	2000
<i>Astragalus daenensis</i> Boiss.	3300	4200	900
<i>Astragalus darlingtonii</i> Podl.	1600	2751	1151
<i>Astragalus demavendicola</i> Bornm. & Gauba	1483	3300	1817
<i>Astragalus demavendicus</i> Boiss. & Buhse	900	2450	1550
<i>Astragalus dictyolobus</i> Bunge	1100	2900	1800
<i>Astragalus doghrunensis</i> Maassoumi & Podlech	1488	3550	2062
<i>Astragalus dschuparensis</i> Freyn & Bornm.	1762	3500	1738
<i>Astragalus durandianus</i> Aitch. & Baker	1473	3500	2027
<i>Astragalus ebenoides</i> Boiss.	1160	2700	1540

<i>Astragalus eburneus</i> Bornm. & Gauba	300	1920	1620
<i>Astragalus ecbatanus</i> Bunge	342	2130	1788
<i>Astragalus echidna</i> Bunge	1115	2300	1185
<i>Astragalus elwendicus</i> Bornm.	1480	2500	1020
<i>Astragalus eriopodus</i> Boiss.	1000	2724	1724
<i>Astragalus eriostomus</i> Bornm.	1675	2500	825
<i>Astragalus erubescens</i> Podl.	1018	1997	979
<i>Astragalus erwini-gaubae</i> Sirj. & Rech.f.	895	2120	1225
<i>Astragalus erythrolepis</i> Boiss.	1600	2500	900
<i>Astragalus esferayenicus</i> Podl. & Maassoumi	1400	3000	1600
<i>Astragalus evanensis</i> Maassoumi & Podl.	300	2000	1700
<i>Astragalus expetitus</i> Maassoumi	1761	2665	904
<i>Astragalus exspectatus</i> Maassoumi	1720	3900	2180
<i>Astragalus fagh-soleimanensis</i> Maassoumi & Podlech	1325	2130	805
<i>Astragalus farsicus</i> Sirj & Rech.f.	15	2224	2209
<i>Astragalus fasciculifolius</i> Boiss.	50	2230	2180
<i>Astragalus filifoliolatus</i> Maassoumi	772	1811	1039
<i>Astragalus fischeri</i> Fisch.	500	2432	1932
<i>Astragalus floccosus</i> Boiss	900	3100	2200
<i>Astragalus fragiferus</i> Bunge	1700	3600	1900
<i>Astragalus fridae</i> Rech.f.	1150	2800	1650
<i>Astragalus fuliginosus</i> Beck	280	1594	1314
<i>Astragalus gaubae</i> Bornm.	684	1530	846
<i>Astragalus ghahremanii</i> Maassoumi & Podl.	89	1280	1191
<i>Astragalus ghashghaicus</i> Tietz & Zarre	2550	3700	1150
<i>Astragalus gigantirostratus</i> Maassoumi et al.	800	1800	1000
<i>Astragalus glaucacanthus</i> Fisch.	107	2500	2393
<i>Astragalus glaucops</i> Bornm.	1780	3100	1320
<i>Astragalus glumaceus</i> Boiss.	1100	2500	1400
<i>Astragalus griseus</i> Boiss.	1300	3950	2650
<i>Astragalus gueldenstaedtiae</i> Bunge	1290	2500	1210
<i>Astragalus gulul-saranii</i> Podlech	800	2900	2100
<i>Astragalus gypsaceus</i> Beck	750	2160	1410
<i>Astragalus hamadanus</i> Boiss.	1200	2500	1300
<i>Astragalus harazensis</i> Zarre & Podlech	1464	2500	1036
<i>Astragalus herbertii</i> Maassoumi	1900	3000	1100
<i>Astragalus heterodoxus</i> Bunge.	2000	3600	1600
<i>Astragalus hirticalyx</i> Bunge	1601.11	3200	1598.89
<i>Astragalus holopsis</i> Bunge	1420	2700	1280
<i>Astragalus holosemius</i> Bunge	1450	2900	1450
<i>Astragalus horridus</i> Boiss.	2300	3950	1650
<i>Astragalus huthianus</i> Freyn & Bornm.	1600	3081	1481
<i>Astragalus hymenocalyx</i> Boiss.	1500	2750	1250
<i>Astragalus hymenostegis</i> Fisch. & C.A. Mey.	1400	2450	1050
<i>Astragalus hypsogeton</i> Bonge	1393	3000	1607
<i>Astragalus ibicinus</i> Boiss. & Haussk. in Boiss.	1500	3250	1750

<i>Astragalus icmadophilus</i> Hand.-Mazz.	1940	3500	1560
<i>Astragalus impexus</i> Podl.	1500	3000	1500
<i>Astragalus indistinctus</i> Podl. & Maassoumi	931	2900	1969
<i>Astragalus iodotropis</i> Boiss. & Hohen.	2500	3852.3	1352.3
<i>Astragalus iranicus</i> Bunge	950	3000	2050
<i>Astragalus ischredensis</i> Bunge	900	3100	2200
<i>Astragalus jacobsii</i> Podlech	1300	2200	900
<i>Astragalus jesdianus</i> Boiss. & Buhse	569	2484	1915
<i>Astragalus jessenii</i> Bunge	600	2500	1900
<i>Astragalus johannis</i> Boiss.	1257	3700	2443
<i>Astragalus joharchii</i> Ghahremani nejad & J. F. Gaskin	680	2100	1420
<i>Astragalus kashafensis</i> Podl.	907	1854	947
<i>Astragalus kashmarense</i> Maassoumi & Podl.	1000	1950	950
<i>Astragalus kentrophyllus</i> Podlech	300	1300	1000
<i>Astragalus keredjensis</i> Podlech	850	2500	1650
<i>Astragalus khoshjailensis</i> Sirj. & Rech.f.	600	2170	1570
<i>Astragalus kordloricus</i> Zarre	1560	2550	990
<i>Astragalus kuhidashtehensis</i> Podlech	2100	3050	950
<i>Astragalus lacus-valashti</i> Maassoumi	430	2600	2170
<i>Astragalus laristanicus</i> Bornm. & Gauba	200	1755	1555
<i>Astragalus lateritiiformis</i> Zarre	2100	3100	1000
<i>Astragalus lateritius</i> Boiss & Hausskn. In Boiss.	600	2600	2000
<i>Astragalus ledinghamii</i> Barneby	650	2900	2250
<i>Astragalus leonardii</i> Maassoumi	1647	3000	1353
<i>Astragalus lepidus</i> Podl.	1700	3000	1300
<i>Astragalus leptyticus</i> Maassoumi	1350	3050	1700
<i>Astragalus lilacinus</i> Boiss.	1350	3100	1750
<i>Astragalus longicuspis</i> Bunge	1000	2150	1150
<i>Astragalus longirostratus</i> Pau	1340	2800	1460
<i>Astragalus longistylus</i> Bunge	1200	3500	2300
<i>Astragalus lycioides</i> Boiss.	1200	3900	2700
<i>Astragalus macrosemius</i> Boiss. & Hohen.	3300	4100	800
<i>Astragalus magistratus</i> Maassoumi	1050	2300	1250
<i>Astragalus megalocystis</i> Bunge	1380	2400	1020
<i>Astragalus melanocalyx</i> Boiss. & Buhse	1750	3500	1750
<i>Astragalus melanodon</i> Boiss.	1600	3900	2300
<i>Astragalus membranostipulus</i> Maassoumi	1457	2300	843
<i>Astragalus microphysa</i> Boiss.	1860	3800	1940
<i>Astragalus modestus</i> Boiss. & Hohen.	2500	3500	1000
<i>Astragalus monanthemus</i> Boiss.	1700	3700	2000
<i>Astragalus mucronifolius</i> Boiss.	921	2700	1779
<i>Astragalus murinus</i> Boiss.	1760	3740	1980
<i>Astragalus myriacanthus</i> Boiss.	1200	3750	2550
<i>Astragalus naftabensis</i> Sirj. & Rech.f.	2300	3200	900
<i>Astragalus neochalderanicus</i> Podlech & Maassoumi	1680	2800	1120
<i>Astragalus nigrohirsutus</i> (Tietz & Zarre) Borjan	1750	3600	1850

<i>Astragalus nubicola</i> Podlech	1450	3200	1750
<i>Astragalus nurensis</i> Boiss. & Buhse	150	2800	2650
<i>Astragalus ochreatus</i> Bunge	1300	2131	831
<i>Astragalus ochrochlorus</i> Boiss. & Hohen.	1700	3900	2200
<i>Astragalus ovigerus</i> Boiss.	2200	3600	1400
<i>Astragalus ovoideus</i> Sirj. & Rech.f.	550	2757	2207
<i>Astragalus paralurges</i> Bunge	1500	2600	1100
<i>Astragalus parvulus</i> Bornm.	1586	3630	2044
<i>Astragalus patrius</i> Maassoumi	1400	3400	2000
<i>Astragalus paucifoliolatus</i> Podlech	1225	2539	1314
<i>Astragalus pauperiflorus</i> Bornm.	1600	3550	1950
<i>Astragalus pauxillis</i> Maassoumi & F.Ghahremani	1850	2650	800
<i>Astragalus pellitus</i> Bunge	720	2300	1580
<i>Astragalus penetratus</i> Maassoumi	1800	3200	1400
<i>Astragalus pentanthus</i> Boiss.	1000	2500	1500
<i>Astragalus perdurans</i> Podl.	2100	3950	1850
<i>Astragalus persicus</i> (DC.) Fisch. & C.A. Mey.	1600	2513	913
<i>Astragalus piranshahricus</i> Maassoumi & Podl.	1105	2327	1222
<i>Astragalus plagiophacos</i> Maassoumi & Podlech	2200	3900	1700
<i>Astragalus platysematus</i> Bunge	1600	3800	2200
<i>Astragalus plebejus</i> Boiss.	1800	3650	1850
<i>Astragalus podosphaerus</i> Boiss. & Housskn.	2200	3340	1140
<i>Astragalus porphyrophysa</i> Bornm. & Gauba	670	2070	1400
<i>Astragalus pseudocyclophyllus</i> Rech.f.	1200	2500	1300
<i>Astragalus pseudoibicinus</i> Maassoumi & Podlech	1500	2800	1300
<i>Astragalus pseudoindurascens</i> Sirj. & Rech. f.	1200	2100	900
<i>Astragalus pseudopersicus</i> Maassoumi & Podlech	1695	2850	1155
<i>Astragalus pseudorobustus</i> Podlech & Maassoumi	1375	2400	1025
<i>Astragalus pseudoshebarensis</i> Podlech	1500	2500	1000
<i>Astragalus psilostylus</i> Bunge	1663	2620	957
<i>Astragalus ptychophyllus</i> Boiss. in Kotschy	1500	3000	1500
<i>Astragalus raswendicus</i> Hausskn. & Bornm.	1570	2600	1030
<i>Astragalus recognitus</i> Fisch.	1120	2600	1480
<i>Astragalus remotiflorus</i> Boiss.	1600	3600	2000
<i>Astragalus remotijugus</i> Boiss. & Hohen. in Boiss. Diagn. Pl. Or. Nov. Ser. 1	1250	2700	1450
<i>Astragalus renzianus</i> Podlech	1300	3000	1700
<i>Astragalus repentinus</i> Ekici & Podlech	1000	2500	1500
<i>Astragalus reuterianus</i> Boiss.	1100	2580	1480
<i>Astragalus rhabdophorus</i> Bornm.	1300	2600	1300
<i>Astragalus rhodosemius</i> Boiss. & Hausskn.	1200	3700	2500
<i>Astragalus rollovii</i> Grossh	1360	2300	940
<i>Astragalus rosellus</i> Sirj. & Rech.f.	1500	3000	1500
<i>Astragalus rubriflorus</i> Bunge	1700	2600	900
<i>Astragalus rubrocyclinus</i> Maassoumi & Podl.	1075	2700	1625
<i>Astragalus rubrostriatus</i> Bunge	1450	2300	850
<i>Astragalus rudimentus</i> Maassoumi	2000	3900	1900

<i>Astragalus rufescens</i> Freyn	2350	3850	1500
<i>Astragalus ruscifolius</i> Boiss.	1120	2300	1180
<i>Astragalus saccatus</i> Boiss.	1600	2514	914
<i>Astragalus sahendi</i> Fisch.	2500	3600	1100
<i>Astragalus saremii</i> Maassoumi	1000	1870	870
<i>Astragalus savellanicus</i> Podlech	1400	3600	2200
<i>Astragalus scapiger</i> Ranjbar & Maassoumi	1250	2400	1150
<i>Astragalus schmidii</i> Podl.	933	2300	1367
<i>Astragalus sciureus</i> Boiss. & Hohen. in Boiss.	1300	3000	1700
<i>Astragalus sclerocladus</i> Bunge	1636	2550	914
<i>Astragalus semiromensis</i> Podlech & Maassoumi	1500	2725	1225
<i>Astragalus senilis</i> Bornm.	1200	2900	1700
<i>Astragalus shahbazanicus</i> Podlech	1008	2500	1492
<i>Astragalus sitiens</i> Bunge	500	1893	1393
<i>Astragalus sojakii</i> Podl.	1604	3000	1396
<i>Astragalus spachianus</i> Boiss. & Buhse	1200	3300	2100
<i>Astragalus speciosus</i> Boiss. & Hohen.	1800	3400	1600
<i>Astragalus sphaeranthus</i> Boiss.	2162	3800	1638
<i>Astragalus stenolepis</i> Fisch.	1200	2600	1400
<i>Astragalus stenostegius</i> Boiss. & Hausskn.	2100	3950	1850
<i>Astragalus straussii</i> Bornm.	1250	2800	1550
<i>Astragalus subalpinus</i> Boiss. Buhse	1800	3500	1700
<i>Astragalus submittis</i> Boiss. & Hohen.	500	2750	2250
<i>Astragalus subsecundus</i> Boiss. & Hohen.	1700	3600	1900
<i>Astragalus susianus</i> Boiss.	1210	3355	2145
<i>Astragalus sympileicarpus</i> Rech.f.	932	2000	1068
<i>Astragalus tabrizianus</i> Fisch.	1133	2900	1767
<i>Astragalus talimansurensis</i> Sirj. & Rech.f.	219	1500	1281
<i>Astragalus tarumensis</i> Sirj. & Rech.f.	781	3200	2419
<i>Astragalus teheranicus</i> Boiss & Hohen. In Boiss.	920	2700	1780
<i>Astragalus tenellus</i> Bunge	1839	2700	861
<i>Astragalus tenuiramosus</i> Podlech & Zarre	65	2300	2235
<i>Astragalus tenuiscapus</i> Freyn & Bornm.	2460	3915	1455
<i>Astragalus touranicus</i> Freitag & Podl.	1200	3500	2300
<i>Astragalus trachyacanthos</i> Fischer	950	3200	2250
<i>Astragalus tricholobus</i> DC.	1110	2600	1490
<i>Astragalus turgidus</i> Podlech	2700	3900	1200
<i>Astragalus vanillae</i> Boiss.	819	2500	1681
<i>Astragalus vegetus</i> Bunge	1009	3000	1991
<i>Astragalus vereskensis</i> Maassoumi & Podl.	200	2750	2550
<i>Astragalus xiphidiopsis</i> Bornm.	1300	2450	1150
<i>Astragalus yazdii</i> (Vassilcz.) Podlech & Maassoumi	2503	4000	1497
<i>Astragalus yushensis</i> T. Sabaii	500	2500	2000
<i>Astragalus zoshkensis</i> Ghahremaninejad	1700	3000	1300
<i>Astrodaucus persicus</i> (Boiss.) Drude in Engler & Prantl	550	2750	2200
<i>Asyneuma multicaule</i> (Boiss.) Rech. f. Schiman-Czeina	2100	3300	1200

Atraphaxis aucheri Jaub. & Spach	300	1225	925
Azilia eryngioides (Pau) Hedge & Lamond	400	2400	2000
Ballota platyloma Rech. f.	400	2750	2350
Bellevalia decolorans Bornm.	1000	2693	1693
Bellevalia shirazana Parsa	2085	3020	935
Biarum platyspathum Bornm.	895	2000	1105
Biarum straussii Engl.	800	2450	1650
Bromus frigidus Boiss. & Hausskn.	2700	3800	1100
Brossardia papyracea Boiss.	1700	2600	900
Bufonia enervis Boiss.	1400	3042	1642
Bufonia koelzii Rech. f.	1900	3300	1400
Bufonia kotschyana Boiss.	1400	3200	1800
Bufonia macrocarpa Ser. in DC.	1300	3300	2000
Bunium luristanicum Rech. f.	1200	2450	1250
Bunium wolffii Kluykov	980	2500	1520
Bupleurum flexile Bornm. ex Gauba	1200	2700	1500
Bupleurum ghahremanii Mozaffarian	1850	2950	1100
Caccinia kotschyi Boiss.	888	2500	1612
Caccinia strigosa Boiss.	630	2440	1810
Calligonum bungei Boiss.	570	1616	1046
Calligonum denticulatum Bunge ex Boiss.	403	1829	1426
Calligonum persicum (Boiss. & Buhse) Boiss.	200	2700	2500
Calligonum schizopterum Rech.f. & Schiman-Czeika	450	1449	999
Campanula candida DC.	1334	2650	1316
Campanula humillima DC.	1200	3684.69	2484.69
Campanula kermanica (Rech.f.	1020	2713	1693
Campanula kurdistanica Advay & Maroofi	1800	3000	1200
Campanula lourica Boiss.	1500	3300	1800
Campanula luristanica Freyn	2000	2800	800
Campanula persepolitana Ky. in Boiss.	1340	3000	1660
Centaurea albonitens Turrill	1500	2780	1280
Centaurea amadanensis Schultz-Bip.	1200	2500	1300
Centaurea aucheri Wagenitz	1300	2780	1480
Centaurea aziziana Rech.f.	1070	2100	1030
Centaurea congesta Wagenitz	1200	3000	1800
Centaurea gabrielae (Bornm.) Wagenitz	1000	1870	870
Centaurea galactochroa Rech.f.	1100	2250	1150
Centaurea gaubae (Bornm.) Wagenitz	1346	2800	1454
Centaurea gilanica Bornm.	1300	2695.06	1395.06
Centaurea hyrcanica Bornm.	400	2750	2350
Centaurea imperialis Hausskn. ex Bornm.	1100	2160	1060
Centaurea incanescens (DC.) Sch. Bip.	1204.61	2820	1615.39
Centaurea intricata Boiss.	100	1966.52	1866.52
Centaurea irritans Wagenitz	979.15	2000	1020.85
Centaurea ispahanica Boiss.	1223	3300	2077
Centaurea kandavanensis Wagenitz	270	2696.33	2426.33

Centaurea leuzeoides (Jaub. & Spach) Walp.	1210	2589	1379
Centaurea liristanica Rech.f.	900	2150	1250
Centaurea microlonchoides Boiss.	1202.19	2161.33	959.14
Centaurea persica Boiss.	1550	3000	1450
Centaurea sintenisiana Gand.	400	2220	1820
Centaurea sosnovskyi Grossh.	550	2670	2120
Centaurea urvillei DC.	700	2450	1750
Centaurea ustulata DC.	1263.19	3000	1736.81
Centaurea xeranthemoides Rech.f.	1467.35	2700	1232.65
Centaurea zuvandica (SOSN.) SOSN.	900	2780	1880
Cephalaria juncea Boiss.	1465	3100	1635
Cephalorrhynchus brassicifolius (Boiss.) Tuisl	1500	2730	1230
Cephalorrhynchus gorganicus (Rech.f. & Esfand.) Tuisl	1900	2700	800
Cephalorrhynchus kossinskyi (Krasch.) Krip.	1074	2500	1426
Cephalorrhynchus microcephalus (DC.) Schchian	563	2733	2170
Cerasus brachypetala Boiss.	2100	3600	1500
Cerasus chorassanica Pojark.	1134	3000	1866
Chaenorhinum grossecostatum Speta	2192	4000	1808
Chaerophyllum khorassanicum Czern. ex Schischk.	1400	3300	1900
Chaerophyllum nivale Hedge & Lamond	3200	4100	900
Chesneya kotschy Boiss.	77	2500	2423
Cicer spiroceras Jaub. & Spach	1347	3800	2453
Cicer staphianum Rech. f.	2900	3800	900
Cicer subaphyllum Boiss.	1750	2850	1100
Cicer tragacanthoides Jaub. & Spach	2700	3800	1100
Cirsium bracteosum DC.	1389.68	2339.49	949.81
Cirsium lappaceum M. B.	2000	3400	1400
Cirsium spectabile DC.	1700	3000	1300
Cistanche eremodoxa Bornm.	1200	2046	846
Clastopus erubescens Hausskn.	2400	3300	900
Clastopus vestitus (Desv.) Boiss.	1830	3352.05	1522.05
Cleome foliolosa DC.	1100	2287	1187
Colchicum varians (Freyen & Bornm.) Czernjak.	1300	2850	1550
Colchicum wendelboi K. Persson	850	2951	2101
Colpodium violaceum (Boiss.) Griseb.	3200	4000	800
Colutea persica Boiss.	1000	3500	2500
Colutea porphyrogramma Rech. f.	730	2000	1270
Colutea uniflora C. Back in Stapf	300	1250	950
Consolida linarioides (Boiss.) Munz	1600	2700	1100
Consolida teheranica (Boiss.) Rech. f.	300	2800	2500
Convolvulus argyracanthus Rech.f.	419	1900	1481
Convolvulus cephalophorus Boiss.	21	1000	979
Convolvulus eremophilus Boiss. et Buhse	750	1803	1053
Convolvulus gonocladus Boiss.	15	1492	1477
Convolvulus oxysepalus Boiss.	260	2700	2440
Convolvulus schirazianus Boiss.	1000	2900	1900

<i>Convolvulus stapfii</i> Rech.f.	450	1789	1339
<i>Convolvulus turrillianus</i> Parsa	160	2300	2140
<i>Convolvulus urosepalus</i> Pau	2518	3600	1082
<i>Cotoneaster assadii</i> Khatamsaz	1040	2250	1210
<i>Cotoneaster esfandiarii</i> Khatamsaz	2500	3400	900
<i>Cotoneaster persicus</i> Pojark.	1000	3300	2300
<i>Cousinia adenosticta</i> Bornm.	2200	3500	1300
<i>Cousinia aggregata</i> DC.	1100	2600	1500
<i>Cousinia akredii</i> Bornm. & Gauba	1500	2782.81	1282.81
<i>Cousinia alexeenkoana</i> Bornm.	1250	2550	1300
<i>Cousinia amplissima</i> (Boiss.) Boiss.	1000	2523.23	1523.23
<i>Cousinia antonowii</i> C. Winkl.	247	2170.27	1923.27
<i>Cousinia arakensis</i> Attar & Djavadi	1635	2500	865
<i>Cousinia araneosa</i> DC.	1641.34	3700	2058.66
<i>Cousinia archibaldii</i> Rech.f.	1800	3600	1800
<i>Cousinia arctotidifolia</i> Bunge	885.29	2107.73	1222.44
<i>Cousinia assyriaca</i> Jaub. & Spech	1026.4	2500	1473.6
<i>Cousinia bachtiarica</i> Boiss. & Hausskn.	2301.67	3419.53	1117.86
<i>Cousinia barbeyi</i> C.Winkl.	1100	2400	1300
<i>Cousinia belangeri</i> DC.	976.9	2000	1023.1
<i>Cousinia calcitrapa</i> Boiss.	1459.01	3000	1540.99
<i>Cousinia calocephala</i> Jaub. & Spech	1031	3207	2176
<i>Cousinia calolepis</i> Boiss.	1500	2500	1000
<i>Cousinia chaetocephala</i> Kult.	1154.19	2000	845.81
<i>Cousinia chamaepeuce</i> Boiss.	2100	3252.79	1152.79
<i>Cousinia chlorosphaera</i> Bornm.	1459	3400	1941
<i>Cousinia chrysacantha</i> Jaub. & Spech	900	2169.3	1269.3
<i>Cousinia commutata</i> Bunge	1400	3400	2000
<i>Cousinia concinna</i> Boiss. & Hausskn.	1100	3000	1900
<i>Cousinia concolor</i> Bunge	1446.88	2300	853.12
<i>Cousinia crispia</i> Jaub. & Spech	2200	3800	1600
<i>Cousinia cylindracea</i> Boiss.	1200	2962.75	1762.75
<i>Cousinia decipiens</i> Boiss. & Hohen.	1200	2810	1610
<i>Cousinia denaensis</i> Attar & Djavadi	1810	3170	1360
<i>Cousinia edmondsonii</i> Rech.f.	1750	2810	1060
<i>Cousinia ekbatanensis</i> Bornm.	1830	2962.75	1132.75
<i>Cousinia elata</i> Boiss. & Buhse	277	3000	2723
<i>Cousinia erinacea</i> Jaub. & Spach	247.96	1545.27	1297.31
<i>Cousinia eriobasis</i> Bunge	1700	2600	900
<i>Cousinia eryngioides</i> Boiss.	900	2250	1350
<i>Cousinia esfandiarii</i> Rech.f. & Aell.	750	2426.31	1676.31
<i>Cousinia fabrorum</i> Rech.f.	1056	2300	1244
<i>Cousinia firuzkuhensis</i> Rech.f.	1800	2800	1000
<i>Cousinia freynii</i> Bornm.	1000	2213.7	1213.7
<i>Cousinia gabrielae</i> Bornm.	1600	2700	1100
<i>Cousinia gatchsaranica</i> Mehregan	1460	2800	1340

<i>Cousinia gaubae</i> Bornm.	1400	2700.04	1300.04
<i>Cousinia gedrosiaca</i> Bornm.&Gauba	1100	2700	1600
<i>Cousinia gilanica</i> Bornm.	2000	3100	1100
<i>Cousinia glaucopsis</i> Bornm. & Rech.f.	1302.05	2400	1097.95
<i>Cousinia gmelini</i> C. Winkl.	2600	3400	800
<i>Cousinia gracilis</i> Boiss.	2300	3100	800
<i>Cousinia grandis</i> C. A. Mey.	1250	2700	1450
<i>Cousinia harazensis</i> Rech.f.	1850	3600	1750
<i>Cousinia heliantha</i> Bunge	1030.08	2193.82	1163.74
<i>Cousinia hergtiana</i> Bornm.	1890	2700	810
<i>Cousinia hypoleuca</i> Boiss.	1300	2900	1600
<i>Cousinia hypopolia</i> Bornm.& Sint.	700	2260.43	1560.43
<i>Cousinia irritans</i> Rech.f.	1900	3650	1750
<i>Cousinia kandavanensis</i> Attar	1263	2733	1470
<i>Cousinia komarovii</i> (O. Kuntze) C. Winkl.	879.15	1850	970.85
<i>Cousinia kornhuberi</i> Heimerl	1700	3300	1600
<i>Cousinia kotschyi</i> Boiss.	1100	3300	2200
<i>Cousinia lepida</i> Bunge ex Boiss.	880.85	1750	869.15
<i>Cousinia longifolia</i> C. Winkl. & Bornm.	2394.7	3690.23	1295.53
<i>Cousinia lurorum</i> (Bornm.) Bornm.	1200	2750	1550
<i>Cousinia maassoumii</i> Assadi	1400	2350	950
<i>Cousinia macrocephala</i> C. A. Mey.	614	2664	2050
<i>Cousinia meshhedensis</i> Bornm. & Rech.f.	1300	2880	1580
<i>Cousinia neurocentra</i> Bunge	1000	2300	1300
<i>Cousinia noeana</i> Boiss.	1490	2673.3	1183.3
<i>Cousinia onopordioides</i> Ledeb.	944	2881	1937
<i>Cousinia oreodoxa</i> Bornm. & Sint.	1000	2801.93	1801.93
<i>Cousinia orthoclada</i> Hausskn. & Bornm.	2100	2900	800
<i>Cousinia pergamacea</i>	1200	2300	1100
<i>Cousinia pinarocephala</i> boiss	1800	3090	1290
<i>Cousinia piptocephala</i> Bunge	947.2	2625.44	1678.24
<i>Cousinia pterocaulos</i> (C. A. Mey.) Rech.f.	1500	3350	1850
<i>Cousinia pugionifera</i> Jaub. & Spech	1551.21	2548.91	997.7
<i>Cousinia rechingerae</i> Bornm.	464.14	1300	835.86
<i>Cousinia recurvata</i> DC.	1800	3000	1200
<i>Cousinia sagittata</i>	1285	2970	1685
<i>Cousinia seidlitzii</i> Bunge	1000	1979.92	979.92
<i>Cousinia shahvarica</i> Rech.f.	1750	3400	1650
<i>Cousinia sicigera</i> C. Winkl. & Bornm.	2700	4150	1450
<i>Cousinia silyboides</i>	1300	2600	1300
<i>Cousinia smirnowii</i> Trautv	1350	3340	1990
<i>Cousinia sphaerocephala</i> Jaub. & Spech	2200	3300	1100
<i>Cousinia stahliana</i> Bornm. & Gauba	300	2500	2200
<i>Cousinia straussii</i> Hausskn. & Winkl. ex Winkl.	1433	2371.35	938.35
<i>Cousinia tabriziana</i> Bunge	1000	1860.15	860.15
<i>Cousinia tenuifolia</i> C. A. Mey.	1272.72	2250	977.28

<i>Cousinia trachylepis</i> Bunge	993.6	2000	1006.4
<i>Cousinia turcomanica</i> C. Winkl.	1100	2250	1150
<i>Cousinia urumiensis</i> Bornm.	1000	2400	1400
<i>Cousinia verbascifolia</i> Bunge	284	2500	2216
<i>Cousinia xipholepis</i> Boiss.	2113.37	3500	1386.63
<i>Crepis asadbarensis</i> Bornm. ex Rech.f.	2250	3600	1350
<i>Crepis ciliata</i> C. Koch	600	2800	2200
<i>Crepis elbrusensis</i> Boiss.	2800	3900	1100
<i>Crepis elymaitica</i> Bornm.	2400	3467	1067
<i>Crepis gaubae</i> Bornm.	1360	2200	840
<i>Crepis heterotricha</i> DC.	3000	4100	1100
<i>Crepis khorassanica</i> Boiss.	1500	2500	1000
<i>Crepis papposissima</i> Babcock	2500	3500	1000
<i>Crepis quercifolia</i> Bornm. & Gauba	600	1754	1154
<i>Crepis sahendi</i> Boiss. & Buhse	2500	3400	900
<i>Crepis straussii</i> Bornm.	570	2343	1773
<i>Crepis willemetoides</i> Boiss.	50	2900	2850
<i>Crocus gilanicus</i> Mathew	1560	2412	852
<i>Crucianella gilanica</i> Trin.	1400	3500	2100
<i>Crucianella platyphylla</i> Ehrend. & Schonb.-Tem.	1000	2500	1500
<i>Cyclotrichium depauperatum</i> (Bunge) Manden. & Scheng.	910	2700	1790
<i>Cyclotrichium haussknechtii</i> (Bunge) Manden. & Scheng.	1500	2300	800
<i>Cyclotrichium straussii</i> (Bornm.) Rech.f.	1130	2400	1270
<i>Daucus littoralis</i> Smith in Sibth. & Smith	-25	1370	1395
<i>Delphinium aquilegifolium</i> (Boiss.) Bornm.	1200	3000	1800
<i>Delphinium elbursense</i> Rech. f.	1200	3155.19	1955.19
<i>Delphinium lalesaricum</i> Iranshahr	2650	3453.69	803.69
<i>Delphinium lanigerum</i> Boiss. & Hohen.	1500	3600	2100
<i>Delphinium saniculifolium</i> Boiss.	1800	3550	1750
<i>Delphinium tuberosum</i> Auch. ex Boiss.	1000	2900	1900
<i>Delphinium ursinum</i> Rech. f.	870	3100	2230
<i>Demavendia pastinacifolia</i> (Boiss. & Hausskn.) Pimenov	1000	2600	1600
<i>Deyeuxia parsana</i> Bor	2300	3350	1050
<i>Dianthus agrostolepis</i> Rech. f.	420	2000	1580
<i>Dianthus austroiranicus</i> Lempert in Rech.f.	200	2600	2400
<i>Dianthus diversifolius</i> Assadi	1550	2900	1350
<i>Dianthus erythrocoleus</i> Boiss.	2300	4100	1800
<i>Dianthus hyrcanicus</i> Rech. f.	450	2250	1800
<i>Dianthus macranthoides</i> Hausskn. ex Bornm.	200	2700	2500
<i>Dianthus subaphyllus</i> (Lempert) Rech. f.	97	2700	2603
<i>Dianthus szowitsianus</i> Boiss.	965	2700	1735
<i>Dianthus tabrizianus</i> Bienert ex Boiss.	1020	2338	1318
<i>Diceratella canescens</i> (Boiss.) Boiss.	70	1200	1130
<i>Dicyclophora persica</i> Boiss.	45	2150	2105
<i>Didymophysa aucheri</i> Boiss.	3200	4800	1600
<i>Dielsiocharis kotschy</i> (Boiss.) O. E. Schulz.	2000	4100	2100

<i>Dionysia archibaldii</i> Wendelbo	2250	4150	1900
<i>Dionysia aretioides</i> (Lehm.) Boiss.	600	3200	2600
<i>Dionysia bryoides</i> Boiss.	1700	3220	1520
<i>Dionysia caespitosa</i> (Duby) Boiss.	2100	3300	1200
<i>Dionysia diapensiifolia</i> Boiss.	1000	2700	1700
<i>Dionysia gaubae</i> Bornm.	1100	2430	1330
<i>Dionysia haussknechtii</i> Bornm. & Strauss	2100	3200	1100
<i>Dionysia iranica</i> Jamzad	1500	2350	850
<i>Dionysia lamingtonii</i> Stapf	1700	3200	1500
<i>Dionysia leucotricha</i> Bornm.	1200	2900	1700
<i>Dionysia oreodoxa</i> Bornm.	2000	3400	1400
<i>Dionysia revoluta</i> Boiss.	1600	3700	2100
<i>Dionysia sawyeri</i> (Watt) Wendelbo	1460	2900	1440
<i>Dionysia termeana</i> Wendelbo	2680	3500	820
<i>Dionysia zagrica</i> Grey-Wilson	1500	2850	1350
<i>Diplotaenia cachrydifolia</i> Boiss.	2200	3100	900
<i>Dorema ammoniacum</i> D. Don	1000	3250	2250
<i>Dorema aucheri</i> Boiss.	1700	2550	850
<i> Doronicum bracteatum</i> Edmondson	2248	3300	1052
<i>Doronicum wendelboi</i> Edmondson	1513.47	2600	1086.53
<i>Draba pulchella</i> Willd. ex DC.	3200	4000	800
<i>Dracocephalum aucheri</i> Boiss.	3350	4400	1050
<i>Dracocephalum ghahremani</i> Jamzad	2200	3000	800
<i>Dracocephalum kotschy</i> Boiss.	1290	3400	2110
<i>Dracocephalum polychaetum</i> Bornm.	2900	4000	1100
<i>Dracocephalum surmandinum</i> Rech. f.	3000	3900	900
<i>Ducrosia assadii</i> Alava	1300	2700	1400
<i>Echinophora cinerea</i> (Boiss.) Hedge & Lamond	2000	3050	1050
<i>Echinophora platyloba</i> DC.	1200	3700	2500
<i>Echinops austro-iranicus</i> Mozaff.	650	1460	810
<i>Echinops bakhtiaricus</i> Rech.f.	459.26	1320	860.74
<i>Echinops cephalotes</i> DC.	900	2450	1550
<i>Echinops ceratophorus</i> Boiss.	1520	2780	1260
<i>Echinops chorassanicus</i> Bunge	800	2700	1900
<i>Echinops cyanocephalus</i> Boiss.	1100	3000	1900
<i>Echinops dichrous</i> Boiss. & Hausskh.	100	2000	1900
<i>Echinops ecbatanus</i> Bornm.	1370	2962.75	1592.75
<i>Echinops elburensis</i> Rech.f.	1950	3260	1310
<i>Echinops elymaiticus</i> Bornm.	1750	2962.75	1212.75
<i>Echinops farsicus</i> Rech. F	950	2300	1350
<i>Echinops gedrosiacus</i> Bornm. ўўўў ў	600	1656.2	1056.2
<i>Echinops glandulos-punctatus</i> Rech.f.	852.44	1670.74	818.3
<i>Echinops haussknechtii</i> Boiss.	600	2520.52	1920.52
<i>Echinops heteromorphus</i> Bunge	1043.78	1900	856.22
<i>Echinops ilicifolius</i> Bunge	1000	2025.9	1025.9
<i>Echinops jesdianus</i> Boiss.	1800	2700	900

<i>Echinops keredjensis</i> Rech.f.	1139	2500	1361
<i>Echinops koelzii</i> Rech.f.	400	3000	2600
<i>Echinops kotschyi</i> Boiss.	2119.17	3200	1080.83
<i>Echinops lalesarensis</i> Bornm.	1900	3000	1100
<i>Echinops longipeticillatus</i> Mozaff. & Ghahr.	100	1500	1400
<i>Echinops macrophyllus</i> Boiss. & Hausskn.	70	2600	2530
<i>Echinops mosulensis</i> Rech.f.	527	2600	2073
<i>Echinops persepolitanus</i> Rech.f.	940	2100	1160
<i>Echinops polygamus</i> Bunge	1000	2100	1100
<i>Echinops robustus</i> Bunge	997.49	2000	1002.51
<i>Echinops sabzevarensis</i> Mozaff.	1150	2000	850
<i>Echinops shulabadensis</i> Mozaff.	1000	1850	850
<i>Echinops sojakii</i> Rech . F.	1000	1900	900
<i>Echinops tenuisectus</i> Rech.f.	1000	3000	2000
<i>Echinops viscidulus</i> Mozaff.	1430	3200	1770
<i>Elaeosticta nodosa</i> (Boiss.) Boiss.	1250	3100	1850
<i>Elburzia fenestrata</i> (Boiss.) Hedge	1820	3000	1180
<i>Epipactis reichingeri</i> Renz	750	2200	1450
<i>Erigeron hyrcanicus</i> Bornm. & Vierh.	2700	4300	1600
<i>Eriocycla ghafooriana</i> Akhani	550	1600	1050
<i>Eriocycla olivieri</i> (Boiss.) Wolff	750	3000	2250
<i>Eritrichium gracillimum</i> Rech. f.	550	3000	2450
<i>Erysimum caespitosum</i> DC.	1800	4300	2500
<i>Erysimum elbrusense</i> Boiss.	1800	3700	1900
<i>Erysimum nanum</i> Boiss. & Hohen. in Boiss.	1900	4000	2100
<i>Erysimum nasturtioides</i> Boiss. & Hausskn.	2800	3800	1000
<i>Euphorbia connata</i> Boiss. in DC.	1210	2098	888
<i>Euphorbia decipiens</i> Boiss. & Buhse	1000	3100	2100
<i>Euphorbia erythradenia</i> Boiss.	1760	3900	2140
<i>Euphorbia hebecarpa</i> Boiss.	3000	3800	800
<i>Euphorbia iranshahri</i> Pahlevani	1878	3000	1122
<i>Euphorbia mazandaranica</i> Pahlevani	110	1470.82	1360.82
<i>Euphorbia plebeia</i> Boiss.	2000	2816.46	816.46
<i>Euphorbia teheranica</i> Boiss. in DC.	1200	2300	1100
<i>Euphrasia juzepczukii</i> Deniss.	2100	4200	2100
<i>Ferula alliacea</i> Boiss.	1550	2500	950
<i>Ferula behboudiana</i> (Rech.f. & Esfand.) Chamberlain	300	1800	1500
<i>Ferula flabelliloba</i> Rech. f. & Aell.	1700	2550	850
<i>Ferula hezarlalehzarica</i> Ajani	2400	3300	900
<i>Ferula hirtella</i> Boiss.	800	2100	1300
<i>Ferula macrocolea</i> Boiss.	570	2500	1930
<i>Ferula microcolea</i> (Boiss.) Boiss.	1600	3000	1400
<i>Ferula persica</i> Willd.	1300	2820	1520
<i>Ferula serpentinica</i> Rech.f.	1700	2700	1000
<i>Ferula sharifii</i> Rech. f. & Esfand.	1120	2000	880
<i>Ferula stenocarpa</i> Boiss. & Hausskn. in Boiss.	200	2300	2100

<i>Ferula tabasensis</i> Rech. f.	850	1950	1100
<i>Ferulago angulata</i> (schlecht.) Boiss.	800	3000	2200
<i>Ferulago carduchorum</i> Boiss. & Haisskn.	2200	3990	1790
<i>Ferulago contracta</i> Boiss. & Hausskn.	1700	3100	1400
<i>Fibigia multicaulis</i> (Boiss. & Hohen.) Boiss.	2000	3400	1400
<i>Fibigia umbellata</i> (Boiss.) Boiss.	1900	3900	2000
<i>Fortuynia garcinii</i> (Burm.) Shuttlew.	10	1519	1509
<i>Fritillaria chlorantha</i> Hausskn. & Bornm.	1300	2864.63	1564.63
<i>Fritillaria kotschyana</i> Herbert	800	3300	2500
<i>Fritillaria olivieri</i> Baker	1600	3560	1960
<i>Fritillaria straussii</i> Bornm.	1509	2700	1191
<i>Fritillaria zagrica</i> Stapf	1500	3032	1532
<i>Gagea iranica</i> Zarrei & Zarre	438	1435	997
<i>Gagea uliginosa</i> Siehe & Pascher	3000	3800	800
<i>Gagea wendelboi</i> Rech. f.	1000	2508	1508
<i>Gaillonia eriantha</i> Jaub. & Spach	1347	2600	1253
<i>Galium aucheri</i> Boiss.	3000	4300	1300
<i>Galium decumbens</i> (Ehrend.) Ehrend. & Schonb.-Tem.	1500	3700	2200
<i>Galium delicatulum</i> Boiss. & Hohen.	2300	4000	1700
<i>Galium diplopion</i> Boiss. & Hohen. in Boiss.	2000	3400	1400
<i>Galium problematicum</i> (Ehrend.) Ehrend. & Schonb.-Tem.	1800	2900	1100
<i>Galium pseudokurdicum</i> (Ehrend.) Sch"nb.-Tem.	1609.86	3900	2290.14
<i>Galium schoenbeck-Temesiae</i> Ehrend.	2400	3400	1000
<i>Geranium persicum</i> Sch"nbeck-Temesy	1404.31	2800	1395.69
<i>Geum iranicum</i> Khatamsaz	1800	3000	1200
<i>Gladiolus persicus</i> Boiss.	1000	3200	2200
<i>Glaucium contortuplicatum</i> Boiss.	300	2600	2300
<i>Graellsia integrifolia</i> (Rech.f.) Rech.f.	1800	2745	945
<i>Graellsia saxifragifolia</i> (DC.) Boiss. ssp. <i>longistyla</i> Poulter	1700	3500	1800
<i>Graellsia stylosa</i> (Boiss. & Hohen.) Poulter	1400	3200	1800
<i>Grantia discoidea</i> Bunge ex Boiss.	425	1313	888
<i>Gypsophila acantholimoides</i> Bornm.	2300	3300	1000
<i>Gypsophila caricifolia</i> Boiss.	1700	2800	1100
<i>Gypsophila elymaitica</i> Mozaffarian	700	1500	800
<i>Gypsophila iranica</i> Barkoudah	1600	3550	1950
<i>Gypsophila lurorum</i> Rech.f.	1200	2300	1100
<i>Gypsophila melampoda</i> Bienert ex Boiss.	150	1714	1564
<i>Gypsophila persica</i> Barkoudah	1070	2700	1630
<i>Gypsophila polyclada</i> Fenzl ex Boiss.	1500	2300	800
<i>Gypsophila pseudomelampoda</i> Gauba & Rech. f. in Rech. f.	1000	2200	1200
<i>Gypsophila rupestris</i> Mozaffarian	1600	2500	900
<i>Gypsophila xanthochlora</i> Rech. f.	1050	2300	1250
<i>Halimocnemis azarbaijanensis</i> Assadi	218	1200	982
<i>Halimocnemis mamamensis</i> (Bge.) Assadi	300	1854	1554
<i>Haplophyllum canaliculatum</i> Boiss.	110	2200	2090
<i>Haplophyllum furfuraceum</i> Bunge ex Boiss.	700	1950	1250

<i>Haplophyllum glaberrimum</i> Bunge ex Boiss.	790	1917	1127
<i>Haplophyllum lissonotum</i> C.C.Townsend	1650	2523	873
<i>Haplophyllum rubro-tinctum</i> C. Townsend	990	2300	1310
<i>Haplophyllum stapfianum</i> Hand.-Mzt.	427	1862	1435
<i>Haplophyllum virgatum</i> Spach	300	2200	1900
<i>Haplophyllum viridulum</i> Sojak	700	1971	1271
<i>Haussknechtia elymaitica</i> Boiss.	1700	2500	800
<i>Hedysarum callithrix</i> Bunge ex Boiss.	1350	2900	1550
<i>Hedysarum criniferum</i> Boiss.	1581	2993	1412
<i>Helianthemum assadii</i> Ghahremaninejad & Gholamian	20	1436	1416
<i>Helichrysum artemisioides</i> Boiss. & Hausskn.	1477	3430	1953
<i>Helichrysum davisianum</i> Rech.f.	2360	3510	1150
<i>Helichrysum globiferum</i> Boiss.	790	2590	1800
<i>Helichrysum leucocephalum</i> Boiss.	520	2650	2130
<i>Helichrysum oligocephalum</i> DC.	1600	3600	2000
<i>Helichrysum oocephalum</i> Boiss.	220	2720	2500
<i>Helichrysum psychophilum</i> Boiss.	2700	4000	1300
<i>Heliocarya monandra</i> Bunge	1124	2000	876
<i>Heliotropium agdense</i> Bunge	180	1950	1770
<i>Heliotropium aucheri</i>	830	2302	1472
<i>Heliotropium denticulatum</i> Boiss. et Hausskn. in Boiss.	900	1930	1030
<i>Heliotropium esfandiarii</i> Akhani & Riedl	830	1950	1120
<i>Heliotropium samoliflorum</i> Bunge	854	2318	1464
<i>Heracleum anisactis</i> Boiss. & Hohen.	2200	3500	1300
<i>Heracleum gorganicum</i> Rech. f.	1500	2500	1000
<i>Heracleum rechingeri</i> Manden.	1250	2950	1700
<i>Hertia angustifolia</i> (DC.) O. Kuntze	1120	3250	2130
<i>Hesperis nivalis</i> Boiss. & Hausskn.	2180	3149.04	969.04
<i>Hieracium azerbaijanense</i> Lack	1191	2400	1209
<i>Hymenocrater incanus</i> Bunge	1150	2650	1500
<i>Hymenocrater oxyodontus</i> Rech. f.	1430	2260	830
<i>Hymenocrater platystegius</i> Rech. f.	850	2600	1750
<i>Hymenocrater yazdianus</i> Rech. f.	2200	3440.05	1240.05
<i>Hyoscyamus kotschyanus</i> pojark.	1700	3000	1300
<i>Hyoscyamus kurdicus</i> Bornm.	1800	3000	1200
<i>Hyoscyamus malekianus</i> Parsa	1900	3800	1900
<i>Hyoscyamus tenuicaulis</i> Schonbeck-Temesy	1	2400	2399
<i>Hypericum dogonbadanicum</i> Assadi	1000	1900	900
<i>Iranecio elbrusensis</i> (Boiss.) B. Nord	2150	3386.1	1236.1
<i>Iranecio oligolepis</i> (Boiss.) B. Nord.	3000	3800	800
<i>Iranecio paucilobus</i> (DC.) B. Nord.	1300	3550	2250
<i>Iris barnumiae</i> Baker & Foster	2000	3600	1600
<i>Iris meda</i> Stapf	1400	2900	1500
<i>Isatis campylocarpa</i> Boiss.	1480	2501	1021
<i>Isatis gaubae</i> Bornm.	570	2700	2130
<i>Isatis koeiei</i> Rech.f.	700	1900	1200

<i>Isatis raphanifolia</i> Boiss.	90	2379	2289
<i>Isatis rugulosa</i> Bunge ex Boiss.	947	1900	953
<i>Isatis zarrei</i> Al-Shehbaz	1800	2800	1000
<i>Johrenia ramosissima</i> Mozaff.	150	1700	1550
<i>Johreniopsis scoparia</i> (Boiss.) Pimenov	2300	3200	900
<i>Johreniopsis stricticaulis</i> (Rech.f.) M. Pimen.	1010	1900	890
<i>Jurinea bungei</i> Boiss.	1500	2500	1000
<i>Jurinea catharinae</i> Iljin.	800	2185.74	1385.74
<i>Jurinea eriobasis</i> DC.	2000	2800	800
<i>Jurinea heterophylla</i> (Jaub. & Spech) Borss.	898.51	2500	1601.49
<i>Jurinea inuloides</i> Boiss. & hausskn	1500	2400	900
<i>Jurinea leptoloba</i> DC.	1350	2334	984
<i>Jurinea meda</i> Bornm.	2200	4100	1900
<i>Jurinea monocephala</i> Aitch. & Hemsl.	1200	2800	1600
<i>Jurinea radians</i> Boiss.	997	2000	1003
<i>Jurinea stenocalathia</i> Rech.f.	1000	2300	1300
<i>Jurinella frigida</i> (Boiss.) Wagenitz	3300	4200	900
<i>Jurinella microcephala</i> (Boiss.) Wagenitz	2100	3900	1800
<i>Kalakia marginata</i> (Boiss.) Alava	600	2500	1900
<i>Karvandarina aphylla</i> Rech. f.	297	1594	1297
<i>Klasea viciifolia</i> (Boiss. & Hausskn.) L. Martins	1713	2742	1029
<i>Lactuca azerbaijanica</i> Rech.f.	1044	1884	840
<i>Lactuca polyclada</i> Boiss.	2405.85	3300	894.15
<i>Lagochilus alutaceus</i> Bunge	1050	2430	1380
<i>Lagochilus aucheri</i> Boiss.	1000	3120	2120
<i>Lagochilus kotschyanus</i> Boiss.	1000	3100	2100
<i>Lagochilus lasiocalyx</i> (Stapf) Jamzad	1500	2700	1200
<i>Lagochilus macracanthus</i> Fisch. & C.A. Mey. in Schrenk	900	3000	2100
<i>Laser rechingeri</i> Akhani	770	2100	1330
<i>Launaea acanthodes</i> (Boiss.) O. Kuntze	550	3000	2450
<i>Launaea peistocarpa</i> (Boiss.) Rech.f.	1490	2798	1308
<i>Lavandula sublepidota</i> Rech. f.	120	1700	1580
<i>Lepechinella persica</i> (Boiss.) H. Riedl	3000	4000	1000
<i>Lepechinella wendelboi</i> H. Riedl in Wendelbo	2000	4100	2100
<i>Leucopoa pseudosclerophylla</i> (Krivot.) Bor	2100	3042	942
<i>Leutea cupularis</i> (Boiss.) M. Pimen.	1800	3700	1900
<i>Leutea gracillima</i> M. Pimen.	750	2000	1250
<i>Leutea nematoloba</i> (Rech.f.) M. Pimen.	1000	2400	1400
<i>Leutea petiolaris</i> (DC.) Pimenov	1000	3350	2350
<i>Leutea polyscias</i> (Boiss.) M. Pimen.	200	1650	1450
<i>Leutea rechingeri</i> (Leute) Pimenov	2300	3300	1000
<i>Ligularia persica</i> Boiss.	2300	3500	1200
<i>Linaria azerbaijanensis</i> Hamdi & Assadi	1050	2900	1850
<i>Linaria elymaitica</i> (Boiss.) Kuprian.	1744	3200	1456
<i>Linaria golestanensis</i> Hamdi & Assadi	670	1827	1157
<i>Linaria karajensis</i> Hamdi & Assadi	1700	3200	1500

<i>Linaria khalkhalensis</i> Hamdi & Assadi	1500	2500	1000
<i>Linaria lineolata</i> Boiss.	1300	3400	2100
<i>Linaria michauxii</i> Chav.	889	2510	1621
<i>Linaria nurensis</i> Miller	1800	3100	1300
<i>Linaria remotiflora</i> Patzak	2012.4	3235	1222.6
<i>Linaria shahroudensis</i> Hamdi & Assadi	2300	3700	1400
<i>Lindelofia kandavanensis</i> Bornm. & Gauba	1304	2900	1596
<i>Linum persicum</i> Ky. ex Boiss.	1400	3113	1713
<i>Malabaila kotschyi</i> Boiss.	320	2200	1880
<i>Malabaila porphyrodiscus</i> Stapf & Wetst.	1100	2600	1500
<i>Marrubium cordatum</i> N b lek	2800	3600	800
<i>Matthiola dumulosa</i> Boiss. & Buhse	948	2000	1052
<i>Matthiola ovatifolia</i> (Boiss.) Boiss.	900	2800	1900
<i>Matthiola revoluta</i> Bunge ex Boiss.	70	2500	2430
<i>Mentha mozaaffarianii</i> Jamzad	500	2000	1500
<i>Michauxia stenophylla</i> Boiss. & Hausskn.	330	1610	1280
<i>Micrantha multicaulis</i> (Boiss.) F. Dvo k	2100	3350	1250
<i>Micromeria hedgei</i> Rech. f.	650	2000	1350
<i>Minuartia acuminata</i> Turrill	1800	3200	1400
<i>Minuartia glandulosa</i> (Boiss. & A. Huet) Bornm.	1400	3500	2100
<i>Minuartia lineata</i> Bornm.	2600	3700	1100
<i>Minuartia litwinowii</i> Schischk.	1800	3600	1800
<i>Minuartia sublineata</i> Rech. f	1650	3300	1650
<i>Muscari pseudomuscari</i> (Boiss. & Buhse) Wendelbo	251	2200	1949
<i>Myopordon hyrcanum</i> (Bornm.) Wagenitz	3000	3900	900
<i>Myopordon persicum</i> Boiss.	3300	4100	800
<i>Myosotis anomala</i> H.Riedl	33	2450	2417
<i>Nanorrhimum campyloceras</i> (Rech. F. & Esfand.) Naanaie	40	1673	1633
<i>Nanorrhimum chasmophyticum</i> (Wendelbo) Naanaie	2250	3300	1050
<i>Nanorrhimum khuzestanicum</i> Naanaie	100	2166	2066
<i>Nectaroscordum koelzii</i> Wendelbo	1500	2800	1300
<i>Nepeta archibaldii</i> Rech. f.	2300	4140	1840
<i>Nepeta assurgens</i> Hausskn. & Bornm.	1700	3000	1300
<i>Nepeta bakhtiarica</i> Rech.f.	1200	2700	1500
<i>Nepeta binaloudensis</i> Jamzad	1560	3000	1440
<i>Nepeta bokhonica</i> Jamzad	1800	2700	900
<i>Nepeta bornmuelleri</i> Hausskn. & Bornm.	1680	3400	1720
<i>Nepeta cephalotes</i> Boiss.	930	2150	1220
<i>Nepeta chionophila</i> Boiss. & Hausskn.	2440	3437.04	997.04
<i>Nepeta crassifolia</i> Boiss. & Buhse	1200	3450	2250
<i>Nepeta crispia</i> Willd.	2100	4140	2040
<i>Nepeta denudata</i> Benth. in DC.	1150	2600	1450
<i>Nepeta depauperata</i> Benth. in DC.	1500	3500	2000
<i>Nepeta dschuparensis</i> Bornm.	2200	3850	1650
<i>Nepeta elymaitica</i> Bornm.	2000	3500	1500
<i>Nepeta eremokosmos</i> Rech.f.	1100	2800	1700

<i>Nepeta gloeocephala</i> Rech.f.	1500	2700	1200
<i>Nepeta hormozganica</i> Jamzad	300	1600	1300
<i>Nepeta kotschy</i> Boiss.	1080	3300	2220
<i>Nepeta lasiocephala</i> Benth.	2300	4100	1800
<i>Nepeta laxiflora</i> Benth.	1400	3000	1600
<i>Nepeta macrosiphon</i> Boiss.	1800	3600	1800
<i>Nepeta menthoides</i> Boiss. Buhse	2024.18	3304.9	1280.72
<i>Nepeta natanzensis</i> Jamzad	2300	4200	1900
<i>Nepeta oxyodonta</i> Boiss.	1500	3300	1800
<i>Nepeta pogonosperma</i> Jamzad & Assadi	2600	3600	1000
<i>Nepeta prostrata</i> Benth.	1150	2600	1450
<i>Nepeta racemosa</i> Lam.	2000	3700	1700
<i>Nepeta rivularis</i> Bornm.	3000	3800	800
<i>Nepeta schiraziana</i> Boiss.	1500	3000	1500
<i>Nepeta sessilifolia</i> Bunge	2000	4000	2000
<i>Nepeta straussii</i> Hausskn. & Bornm.	1400	2850	1450
<i>Nonea anchusoides</i> Boiss. & Buhse	1305	3750	2445
<i>Nonea iranica</i> Falatoury & Pakravan	734	2100	1366
<i>Nonea persica</i> Boiss.	1000	3000	2000
<i>Onobrychis alborzensis</i> Ranjbar & Hajmoradi	1300	2207	907
<i>Onobrychis aucheri</i> Boiss.	1150	2100	950
<i>Onobrychis gaubae</i> Bornm.	1350	2400	1050
<i>Onobrychis iranensis</i> Amirabadizadeh & Ghanavati	913	2100	1187
<i>Onobrychis mazanderanica</i> Rech.f.	900	2400	1500
<i>Onobrychis melanotricha</i> Boiss.	800	3200	2400
<i>Onobrychis psoraleifolia</i> Boiss.	1313	2500	1187
<i>Onobrychis scrobiculata</i> Boiss.	1000	2500	1500
<i>Onopordon caramanicum</i> (Bornm.) Bornm.	700	2760.9	2060.9
<i>Onosma bilabiata</i> Boiss. & Buhse	1300	3500	2200
<i>Onosma chrysochaetum</i> Bornm.	1650	3050	1400
<i>Onosma demavendica</i> H.Riedl	1500	2950	1450
<i>Onosma iranshahrii</i> Ghahreman & Atter	1800	2600	800
<i>Onosma kilouyense</i> Boiss. & Hausskn	1500	3500	2000
<i>Onosma kotschy</i> Boiss.	1220	3100	1880
<i>Onosma pachypoda</i> Boiss.	1050	2540	1490
<i>Onosma platyphylla</i> H.Riedl	1250	3400	2150
<i>Onosma stenosiphon</i>	3000	4000	1000
<i>Onosma straussii</i> (Riedl) Khatamsaz	120	2350	2230
<i>Opsicarpium insignis</i> Mozaff.	1420	2800	1380
<i>Oreophysa microphylla</i> (Jaub. & Spach) Browicz	1300	2100	800
<i>Ornithogalum pycnanthum</i> Wendelbo	2400	3200	800
<i>Orobanche pulchra</i> Gilli	1220	3400	2180
<i>Otostegia michauxii</i> Briq.	824.9	1700	875.1
<i>Oxytropis bicornis</i> Vassilcz	1100	2450	1350
<i>Oxytropis binaludensis</i> Vassilcz.	1600	3700	2100
<i>Oxytropis chrysocarpa</i> Boiss.	1900	3000	1100

Oxytropis cinerea Vassilcz.	1900	3300	1400
Oxytropis hypsophila Bunge & Boiss.	1200	3000	1800
Oxytropis Iranica Vassilcz.	2240	3850	1610
Oxytropis jahherdehi Maassoumi	1800	2700	900
Oxytropis karjaginii Grossh.	2500	3400	900
Oxytropis kermanica Freyn & Bornm	2700	4100	1400
Oxytropis kuchanensis Vassilcz.	1100	2863	1763
Oxytropis neorechingeriana Vassilcz.	1550	2800	1250
Oxytropis persica Boiss.	2500	3500	1000
Oxytropis rechingeri Vassilcz.	1100	2024	924
Oxytropis szovitsii Boiss. & Buhse	200	3000	2800
Papaver armeniacum (L.) DC	2179.51	3200	1020.49
Papaver persicum Lindl.	1400	2250	850
Papaver tenuifolium Boiss. & Hohen. ex Boiss.	200	2200	2000
Paracaryum cyclhyumenium (Boiss.)H.Riedl	700	2250	1550
Paracaryum luristanicum Nab.	350	2031	1681
Paracaryum modestum Boiss. & Hausskn. ex Boiss.	200	2700	2500
Paracaryum persicum (Boiss.) Boiss.	700	2768	2068
Paracaryum pygmaeum (Rech.f.) Heller in D. Heller & C.C. Heyn	2900	3900	1000
Paraquilegia caespitosa	3500	4800	1300
Parlatoria rostrata Boiss.	1600	2700	1100
Paronychia bungei Boiss.	400	2600	2200
Paronychia caespitosa Stapf	1250	2500	1250
Pentanema flexuosum (Boiss. & Hausskn.) Rech.f.	1700	3000	1300
Pentanema multicaule Boiss.	2100	3750	1650
Pentanema pulicariiforme (DC.) Rech.f.	1466	3528	2062
Phagnalon persicum Boiss.	1700	3400	1700
Phlomis anisodonta Boiss.	950	3300	2350
Phlomis aucheri Boiss.	1200	2700	1500
Phlomis elliptica Benth.	700	2850	2150
Phlomis pachyphylla Rech.f.	450	1806	1356
Phlomis persica Boiss.	0	2800	2800
Phlomoides adenantha Jaub. & Spach	250	2900	2650
Phlomoides azerbaijanica Rech.f.	850	2300	1450
Phlomoides hyoscyamoides Boiss. & Buhse	940	2100	1160
Phlomoides lanata Jamzad	1735	2550	815
Phlomoides pulvinaris Jaub. & Spach	1250	2500	1250
Phuopsis stylosa Benth. & Hook.f	200	2640	2440
Physogeton acanthophyllum Jaub. & Spach	313	1900	1587
Physogeton occultus (Bge.) Assadi	742	1752	1010
Physoptychis gnaphalodes Boiss.	3200	4000	800
Pimpinella deverroides (Boiss.) Boiss.	1500	3500	2000
Pimpinella dichotoma (Boiss. et Hausskn.) Wolff in Engler	1500	2750	1250
Pimpinella gedrosiaca Bornm.	2000	3000	1000
Pimpinella khayyamii Mozaff.	1400	2500	1100
Pimpinella khorasanica Engstrand.	500	1850	1350

<i>Pimpinella tragoides</i> (Boiss.) Benth. & Hook.f. ex Drude	1150	2400	1250
<i>Piptatherum molinioides</i> Boiss.	1872.46	2750	877.54
<i>Platychaete aucheri</i> (Boiss.) Boiss.	16	1800	1784
<i>Platychaete mucronifolia</i> (Boiss.) Boiss.	129	1859	1730
<i>Platychaete velutina</i> Boiss. & Hausskn.	67	911	844
<i>Polygala platyptera</i> Bornm & Gauba	-28	2600	2628
<i>Polygonum aridum</i> Boiss. & Hausskn. in Boiss.	1700	3048	1348
<i>Polygonum dumosum</i> Boiss.	1650	3700	2050
<i>Polygonum hyrcanicum</i> Rech.f.	4	1900	1896
<i>Polygonum salicornioides</i> Jaub. & Spach	800	2950	2150
<i>Polygonum spinosum</i> H. Gross	2400	3900	1500
<i>Polylophium involucratum</i> (Pall.) Boiss.	1010	2810	1800
<i>Postia bombycinia</i> Boiss. & Hausskn.	73	1602	1529
<i>Postia puberula</i> Boiss. & Hausskn.	697	2200	1503
<i>Potentilla argyroloma</i> Boiss. & Hohen.	1800	4200	2400
<i>Potentilla aucheriana</i> Th. Wolf	2200	4200	2000
<i>Potentilla elvendensis</i> Boiss. et Hohen.	1922.68	3183.36	1260.68
<i>Potentilla farsistanica</i> Browicz	1211	2202	991
<i>Potentilla flaccida</i> Th. Wolf	2600	4000	1400
<i>Potentilla iranica</i> (Rech.f.) Schiman-Czeika	1500	2500	1000
<i>Potentilla lignosa</i> Willd. ex D. F. K. Schleidl	2000	3150	1150
<i>Potentilla nuda</i> Boiss.	2200	3900	1700
<i>Potentilla nurensis</i> Boiss. & Hausskn.	1650	3350	1700
<i>Potentilla pannosa</i> Boiss. & Hausskn.	1650	3000	1350
<i>Potentilla polschista</i> Boiss.	3300	4479.93	1179.93
<i>Potentilla porphyrantha</i> Juz.	3300	4150	850
<i>Potentilla poteriifolia</i> Boiss.	2000	3500	1500
<i>Prangos cheilanthifolia</i> Boiss.	1000	2600	1600
<i>Prangos crossoptera</i> Herrnst. & Heyn	1200	2400	1200
<i>Prangos tuberculata</i> Boiss. & Hausskn.	1600	3000	1400
<i>Primula gaubaeaana</i> Bornm.	679	2800	2121
<i>Pseudocamelina aphragmodes</i> (Boiss.) N. Busch	1800	2816.46	1016.46
<i>Pseudocamelina camelinae</i> (Boiss.) N. Busch	1600	2600	1000
<i>Pseudocamelina glaucophylla</i> (DC.) N. Busch	1000	2846.96	1846.96
<i>Pseudofortunia esfandiarii</i> Hedge	1500	2430	930
<i>Pseudofortunia leucoclada</i> (Boiss.) Khosravi	1470	3600	2130
<i>Pseudotrichydium kotschyti</i> (Boiss.) Pimenov & Kluykov	1950	3900	1950
<i>Pseudotrichydium pauciradiatum</i> (Boiss. & Hohen.) Pimenov & Kluykov	1900	3650	1750
<i>Psychogeton persicus</i> (Boiss.) Grierson	2200	3828	1628
<i>Pterocephalus lignosus</i> Freyn & Bornm.	1350	2737	1387
<i>Pterocephalus persicus</i> Boiss.	1600	3100	1500
<i>Pycnocycla acanthorhipsis</i> Rech. f.	685	1600	915
<i>Pycnocycla bashagardiana</i> Mozaffarian	20	1150	1130
<i>Pyrus glabra</i> Boiss.	1200	3000	1800
<i>Pyrus kandevanica</i> Ghahreman	950	2330	1380
<i>Pyrus mazanderanica</i> Schonbeck-Temesy	1400	2400	1000

<i>Quercus brantii</i> Lindl.	1270	2319	1049
<i>Ranunculus amblyolobus</i> Boiss. & Hohen. in Boiss.	2000	3300	1300
<i>Ranunculus aucheri</i> Boiss.	1200	3100	1900
<i>Ranunculus bulbilliferus</i> Boiss. & Hohen.	2100	3000	900
<i>Ranunculus cymophilus</i> Boiss. & Hohen.	3100	3900	800
<i>Ranunculus elbursensis</i> Boiss.	1500	3202.02	1702.02
<i>Ranunculus elymaiticus</i> Boiss. & Hausskn.	2200	4200	2000
<i>Ranunculus eriorrhizus</i> Boiss. & Buhse	2700	4350	1650
<i>Ranunculus papyrocarpus</i> Rech. f.	1530	3400	1870
<i>Ranunculus sahendicus</i> Boiss. & Buhse	1502	2600	1098
<i>Ranunculus sojakii</i> Iranshahr & Rech. f.	2600	3400	800
<i>Ranunculus straussii</i> Bornm.	2300	3392.2	1092.2
<i>Ranunculus trichocarpus</i> Boiss.	1700	3800	2100
<i>Reseda buhseana</i> Mull.-Arg.	1170	2396	1226
<i>Reseda bungei</i> Boiss.	900	1900	1000
<i>Reseda macrobotrys</i> Boiss.	1081	2052	971
<i>Rhabdosciadium aucheri</i> Boiss.	2000	3500	1500
<i>Rhabdosciadium petiolare</i> Boiss. & Hausskn.	1650	3050	1400
<i>Rhamnus cornifolia</i> Boiss. & Hohen.	1700	3700	2000
<i>Rheum kordestanicum</i> Taheri & Assadi	1600	2400	800
<i>Rindera bungei</i> (Boiss.) Gurke in Engler & Prantl	800	2500	1700
<i>Rochelia mirheydari</i> Reidl & Esfandiari	350	1641	1291
<i>Rubia albicaulis</i> Boiss.	1300	2804	1504
<i>Rubia pauciflora</i> Boiss.	1800	3350	1550
<i>Rumex elbursensis</i> Boiss.	1500	3416.12	1916.12
<i>Rumex ephedroides</i> Bornm.	250	2026	1776
<i>Salix firouzkuhensis</i> Maassoumi	1730	2900	1170
<i>Salsola persica</i> Bunge ex Boiss.	800	2000	1200
<i>Salsola yazdiana</i> Assadi	700	1793	1093
<i>Salvia aristata</i> Aucher ex Benth. in DC.	1180	2900	1720
<i>Salvia chorassanica</i> Bunge	1550	3000	1450
<i>Salvia eremophila</i> Boiss.	130	2500	2370
<i>Salvia hypoleuca</i> Benth. in DC.	800	3000	2200
<i>Salvia jamzadii</i> Mozaffarian	1150	2350	1200
<i>Salvia lachnocalyx</i> Hedge	2020.73	3000	979.27
<i>Salvia oligophylla</i> Auch. ex Benth. in DC.	280	2000	1720
<i>Salvia persepolitana</i> Boiss.	180	2000	1820
<i>Salvia sahendica</i> Boiss. & Buhse	850	2600	1750
<i>Salvia sclareopsis</i> Bornm. ex Hedge	300	2550	2250
<i>Salvia sharpii</i> Rech.f. & Esfand.	100	2500	2400
<i>Salvia urmiensis</i> Bunge	1500	2500	1000
<i>Saponaria bodeana</i> Boiss.	1000	2600	1600
<i>Satureja bachtiarica</i> Bunge	1500	3000	1500
<i>Satureja edmondi</i> Briquet	1370	3950	2580
<i>Satureja isophylla</i> Rech.f.	900	3300	2400
<i>Satureja sahendica</i> Bornm.	1300	2950	1650

Saxifraga ramsarica Jamzad	2500	3800	1300
Saxifraga wendelboi Sch"nb.-Tem.	1900	2900	1000
Saxifrage iranica Bornm.	2800	4200	1400
Scabiosa schimperiana Boiss. & Buhse	300	1500	1200
Scilla gorganica Speta	211	1500	1289
Scilla greilhuberi Speta	80	1500	1420
Scilla khorassanica Meikle	976	2800	1824
Sclerochorton haussknechtii Boiss.	2734.37	3900	1165.63
Sclerorhachis leptoclada Rech.f.	200	1916	1716
Sclerorhachis platyrachis (Boiss.) Podl ex Rech.f.	990	2200	1210
Scorzonera calyculata Boiss.	250	3000	2750
Scorzonera grossheimii Lipsch. & Vassilcz	1800	3700	1900
Scorzonera intricata Boiss.	1700	3800	2100
Scorzonera ispahanica Boiss.	1303	2976	1673
Scorzonera kandavanica Rech.f.	600	2445	1845
Scorzonera luristanica Rech.f.	1171	2466	1295
Scorzonera microcalathia (Rech.f.) Rech.f.	552	2253	1701
Scorzonera mucida Rech.f.	750	2800	2050
Scorzonera persica Boiss. & Buhse	2000	3200	1200
Scorzonera perspolitana Boiss.	1000	2846	1846
Scorzonera renzii Rech.f.	1315	2600	1285
Scorzonera rupicola Hausskn.	1787	3235	1448
Scorzonera stenocephala Boiss.	2400	3666.8	1266.8
Scrophularia atroglandulosa Grau	2200	3100	900
Scrophularia crassicaulis Boiss.	2200	3500	1300
Scrophularia crassiuscula Grau	1300	3300	2000
Scrophularia elbursensis Bornm.	1000	3200	2200
Scrophularia farinosa Boiss.	1104	2700	1596
Scrophularia frigida Boiss.	1400	3900	2500
Scrophularia gaubae Bornm.	500	2070	1570
Scrophularia glauca Decne. ex Benth. in DC.	573	2200	1627
Scrophularia gorganica Rech. f.	1900	3700	1800
Scrophularia megalantha Rech.f.	40	2400	2360
Scrophularia oxysepala Boiss.	1500	2971	1471
Scrophularia rechingeri Grau	1065	2000	935
Scrophularia rostrata Boiss. & Buhse	28	2300	2272
Scrophularia schiraziana Attar & Hatami	650	2226	1576
Scrophularia subaphylla Boiss.	1800	3990	2190
Scutellaria bornmuelleri Hausskn. ex Bornm.	600	1660	1060
Scutellaria farsistanica Rech.f.	1700	2750	1050
Scutellaria glechomoides Boiss.	3400	4400	1000
Scutellaria multicaulis Boiss.	1100	4000	2900
Scutellaria nepetifolia Benth.	1295	3300	2005
Scutellaria persica Bornm.	1450	2500	1050
Scutellaria pinnatifida A. Ham.	1800	3500	1700
Scutellaria xylorrhiza Bornm.	1600	2400	800

<i>Sedum callichroum</i> Boiss.	1300	3000	1700
<i>Sedum elburzense</i> Akhiani & Assadi	1500	2500	1000
<i>Sedum kotschyanum</i> Boiss.	2100	4000	1900
<i>Semenovia dichotoma</i> (Boiss.) Manden.	2800	3900	1100
<i>Semenovia frigida</i> (Boiss.) Hausskn.	2400	3900	1500
<i>Semenovia suffruticosa</i> (Freyn & Bornm.) Manden.	2300	3600	1300
<i>Semenovia trajoioides</i> (Boiss.) Manden.	1500	3550	2050
<i>Sempervivum atropatanum</i> Parnell	1350	2800	1450
<i>Sempervivum iranicum</i> Bornm. & Gauba	1110	3000	1890
<i>Senecio lipskyi</i> Lomak.	1540	3400	1860
<i>Senecio vulcanicus</i> Boiss.	3300	4100	800
<i>Serratula haussknechtii</i> Boiss.	2100	3340	1240
<i>Serratula viciifolia</i> Boiss. & Hausskn.	1600	3000	1400
<i>Silene albescens</i> Boiss.	1315	3000	1685
<i>Silene avromana</i> Boiss. & Hausskn.	1500	3450	1950
<i>Silene daenensis</i> Melzh.	3000	4100	1100
<i>Silene elymaitica</i> Bornm.	1670	3350	1680
<i>Silene erysimifolia</i> Stapf	1350	2600	1250
<i>Silene farsistanica</i> Melzh.	1800	3000	1200
<i>Silene ferdowsii</i> Joharchi	668	1532	864
<i>Silene gertraudiae</i> Melzh.	1150	2230	1080
<i>Silene goniocaula</i> Boiss.	2400	3200	800
<i>Silene gynodioica</i> Ghaz.	1150	3500	2350
<i>Silene lucida</i> Chowdhuri	2000	3000	1000
<i>Silene meyeri</i> Fenzl ex Boiss. & Buhse	1200	3000	1800
<i>Silene nurensis</i> Boiss. & Hausskn.	3000	4200	1200
<i>Silene palinotricha</i> Fenzl ex Boiss.	1400	3000	1600
<i>Silene persica</i> Boiss.	2400	3500	1100
<i>Silene pseudoacheriana</i> Melzh.	1090	1920	830
<i>Silene sojakii</i> Melzh.	1000	2780	1780
<i>Stachys acerosa</i> Boiss.	1500	3500	2000
<i>Stachys asterocalyx</i> Rech. f.	1255	3170	1915
<i>Stachys aucheri</i> Benth.	1700	2900	1200
<i>Stachys ixodes</i> Boiss. & Hausskn. ex Boiss.	1710	3000	1290
<i>Stachys kermanshahensis</i> Rech.f.	500	2200	1700
<i>Stachys koelzii</i> Rech.f.	2300	3500	1200
<i>Stachys laxa</i> Boiss. & Buhse	500	2810	2310
<i>Stachys obtusicrena</i> Boiss.	1900	4000	2100
<i>Stachys persepolitana</i> Boiss.	800	2750	1950
<i>Stachys pilifera</i> Benth.	1660	3300	1640
<i>Stachys subaphylla</i> Rech.f.	600	1400	800
<i>Stachys veroniciformis</i> Rech.f.	1100	3200	2100
<i>Stenotaenia nudicaulis</i> Boiss.	1800	3252.79	1452.79
<i>Sterigmostemum longistylum</i> (Boiss.) Bornm.	1260	2200	940
<i>Stipa atriseta</i> Stapf ex Bor	1700	3800	2100
<i>Stipa haussknechtii</i> Boiss.	1950	3200	1250

<i>Straussiella purpurea</i> (Bge.) Hausskn.	1500	2550	1050
<i>Tanacetum dumosum</i> Boiss.	2100	3459.61	1359.61
<i>Tanacetum hololeucum</i> (Bornm.) Podl.	1200	3700	2500
<i>Tanacetum khorassanicum</i> (Krasch.) Parsa	1200	3200	2000
<i>Tanacetum lingulatum</i> (Boiss.) Bornm.	1219	2726	1507
<i>Tanacetum persicum</i> (Boiss.) Mozaff.	1700	3800	2100
<i>Tanacetum tenuisectum</i> (Boiss.) Podl.	1300	3300	2000
<i>Tanacetum walteri</i> (C. Winkl.) Tzvel.	1637	2900	1263
<i>Taraxacum azerbaijanicum</i> Soest	1200	2000	800
<i>Taraxacum iranicum</i> Soest	1988	2906	918
<i>Taraxacum koelzii</i> Soest	1780	3083	1303
<i>Taraxacum kotschy</i> Soest	1640	2800	1160
<i>Taraxacum neo-spurium</i> Soest	2600	3660	1060
<i>Taraxacum rechingeri</i> Soest	1500	2750	1250
<i>Taraxacum roseum</i> Bornm	1900	3586	1686
<i>Taraxacum ruberuliforme</i> Soest	1357	3165	1808
<i>Telephium eriglaucum</i> Williama	1550	2350	800
<i>Tetraena lasiopetalum</i> (Boiss.) Manden.	1500	4000	2500
<i>Tetraena nephrophyllum</i> (Leute) Manden.	1700	3000	1300
<i>Teucrium macrum</i> Boiss. & Hausskn. ex Boiss.	1800	3000	1200
<i>Teucrium persicum</i> Boiss.	80	1900	1820
<i>Thecocarpus meifolius</i> Boiss.	1500	3200	1700
<i>Thlaspi stenocarpum</i> (Boiss.) Hedge	2350	3600	1250
<i>Thymus carmanicus</i> Jalas	2000	3600	1600
<i>Thymus daenensis</i> Celak.	1035	3200	2165
<i>Thymus persicus</i> (Ronniger ex Rech.f.) Jalas	1900	2900	1000
<i>Thymus pubescens</i> Boiss. & Kotschy ex Celak.	1200	3600	2400
<i>Trachydium depressum</i> Boiss.	3000	3900	900
<i>Trachydium kotschy</i> (Boiss.) Boiss.	2000	3900	1900
<i>Trachydium pauciradiatum</i> (Boiss. & Hohen.) Rech. f.	2400	3650	1250
<i>Tragopogon acanthocarpus</i> Boiss.	1300	3690	2390
<i>Tragopogon caricifolius</i> Boiss.	1000	3361	2361
<i>Tragopogon gongylorrhizus</i> Rech.f.	31	900	869
<i>Tragopogon jezdianus</i> Boiss. & Buhse	1300	3352.05	2052.05
<i>Tragopogon kotschy</i> Boiss.	1200	3800	2600
<i>Tragopogon porphyrocephalus</i> Rech.f.	1281	2650	1369
<i>Tragopogon rezaiyensis</i> Rech.f.	1380	3400	2020
<i>Trichodesma aucheri</i> DC.	1428	3027	1599
<i>Trifolium radicosum</i> Boiss. & Hohen.	2800	3800	1000
<i>Trigonella disperma</i> Bornm. ex Vassilcz.	1029	2700	1671
<i>Trigonella elliptica</i> Boiss.	732	2150	1418
<i>Trigonella persica</i> Boiss.	33	2321	2288
<i>Trigonella stenocarpa</i> Rech. f.	1130	2150	1020
<i>Trigonella teheranica</i> Bornm.	1500	2600	1100
<i>Trigonosciadium brachytænium</i> (Boiss.) Alava	1650	2600	950
<i>Trisetum bungei</i> Boiss.	1000	2700	1700

<i>Tulipa montana</i> Lindl.	250	3150	2900
<i>Tulipa ulophylla</i> Wendelbo	600	2500	1900
<i>Ulmus boissieri</i> Graudz	364	2619	2255
<i>Ungernia flava</i> Boiss. & Hausskn. in Boiss.	1416	2381	965
<i>Verbascum aucheri</i> (Boiss.) Hub.-Mor.	1070	2800	1730
<i>Verbascum bornmuellerianum</i> Hub.- Mor.	1313.67	2790.67	1477
<i>Verbascum carmanicum</i> (Bornm.) Hub.-Mor.	2448.22	3400	951.78
<i>Verbascum farsistanicum</i> (Murb.) Hub.- Mor	1000	2400	1400
<i>Verbascum gabrieliae</i> (Bronm.) Hub.-Mor.	2000	3146	1146
<i>Verbascum hasarense</i> Freyn & Bornm. in Freyn	2412	3601	1189
<i>Verbascum intricatum</i> (Benth.) O. Kuntze	1500	3600	2100
<i>Verbascum kermanense</i> Hub.-Mor.	1700	3124	1424
<i>Verbascum lyprocarpum</i> (Murb.) Hub.-Mor.	1273	2509	1236
<i>Verbascum scoparium</i> Mozaffarian	500	2000	1500
<i>Verbascum straussii</i> (Bornm.) Hub.- Mor	1570	2500	930
<i>Verbascum sublobatum</i> Murb.	600	2100	1500
<i>Veronica acrotheca</i> Bornm. & Gauba	1480	2400	920
<i>Veronica aucheri</i> Boiss.	2000	4800	2800
<i>Veronica farinosa</i> Hausskn.	1282	3000	1718
<i>Veronica fragilis</i> Boiss. & Hausskn.	1800	3200	1400
<i>Veronica francispetae</i> M. A. Fischer	-18	1200	1218
<i>Veronica kurdica</i> Benth.	2700	3750	1050
<i>Veronica mazanderanae</i> Wendelbo	70	2400	2330
<i>Veronica mirabilis</i> Wendelbo	2900	3700	800
<i>Veronica paederotae</i> Boiss.	3500	4300	800
<i>Veronica rechingeri</i> M. A. Fischer	1800	3400	1600
<i>Veronica rubrifolia</i> Boiss.	1800	3400	1600
<i>Vicia ciceroidea</i> Boiss.	2563.48	3600.71	1037.23
<i>Vicia kotschyana</i> Boiss.	2400	4110	1710
<i>Vicia persica</i> Boiss.	800	3000	2200
<i>Vicia sojakii</i> Chrtkova-Zertova	2000	3600	1600
<i>Viola pachyrhiza</i> Boiss. & Hohen.	1500	3200	1700
<i>Viola spathulata</i> Willd.	800	3150	2350
<i>Zeravschania aucheri</i> (Boiss.) Pimenov	1300	3300	2000
<i>Zerdana anchonioides</i> Boiss.	2500	4100	1600
<i>Zeugandra iranica</i> P.H.Davis	1020	2154	1134
<i>Zhumeria majdae</i> Rech.f. & Wendelbo	250	2000	1750
<i>Zosima radians</i> Boiss. & Hohen.	1200	2500	1300