



Article Exploring the Species Richness Pattern and Areas of Endemism of Tenebrionidae (Coleoptera) in Xinjiang, China

Yalin Li, Yujie Wang, Hui Zhang, Shaoyang Li, Zichao Chai ២ and Guodong Ren *

The Key Laboratory of Zoological Systematics and Application, College of Life Sciences, Institute of Life Science and Green Development, Hebei University, Baoding 071002, China; 13903120974@139.com (Y.L.); wangyujie@163.com (Y.W.); kidancer2008@163.com (H.Z.); shaoyang3808@163.com (S.L.); chaizichao0329@163.com (Z.C.)

* Correspondence: gdren@hbu.edu.cn; Tel.: +86-13503380558

Abstract: Species richness and areas of endemicity (AOE) are the basis of biogeography, which is of great significance for understanding the evolution of species and making conservation plans. The present study aimed to investigate the species richness pattern and AOEs of Tenebrionidae in Xinjiang, China. We collected information on the geographical distribution of 556 species from several sources and obtained 2226 distribution records for the analyses. The AOEs were detected using the parsimony analysis of endemicity (PAE) and endemicity analysis (EA) at 0.5°, 1°, and 1.5° grid sizes, respectively. A total of six AOEs were found, including three mountain ranges (Altai Mountains, Tianshan Mountains, and Kunlun Mountains) and one basin (Junggar Basin), which was largely congruent with the species richness pattern. The results indicated that the complex terrain and stable climate in the mountainous area played an important role in the formation of tenebrionid species diversity and their endemic areas in Xinjiang.

Keywords: Tenebrionidae; species richness; parsimony analysis of endemicity; endemicity analysis; area of endemism; mountainous areas

1. Introduction

Xinjiang (~166 km²), a section of Central Asia, is the largest autonomous region located in northwest China, with a complex terrain environment of two basins and three mountains [1,2]. The Tarim Basin and Kunlun Mountains are located in southern and western Xinjiang, Altai Mountain and Junggar Basin are in the north and east areas, and the Tianshan Mountains run through the central areas [3,4]. Xinjiang is surrounded by mountains, which not only leads to complex and varied topography but also to drought and a low-rain climate scenario [5–7]. A large part of the region is arid and semiarid, and one-sixth is covered by desert [3]. Due to the characteristics of its special geographical location, Xinjiang has become a biodiversity hotspot with high species richness and endemic levels, especially breeding many drought-tolerant insect species [8–11]. For example, 84 species of Meloidae Gyllenhal, 1810, accounting for approximately 40% of the total number of species occurring in China, were recorded in Xinjiang [12]. More remarkably, a total of 422 species of Tenebrionidae from Xinjiang were listed [13–16].

Tenebrionidae, a large insect group, is widely distributed in Central Asia [17,18]. It has a high level of species richness in a variety of environments [19,20]. There are approximately 20,000 known species of Tenebrionidae in the world [21,22]. It has been reported that more than 600 species are distributed in the desert and semidesert areas of China, with Xinjiang as the most important geographical component [23]. However, in recent years, a series of events have affected the natural environment in Xinjiang, which may have changed the habitat of some insects, such as the increase in rainfall [24,25], the rise in temperature [5], the prevention and control of desertification [26], and the trend of homogenizing the landscape [20]. As a classic indicator of the desert ecological environment, Tenebrionidae



Citation: Li, Y.; Wang, Y.; Zhang, H.; Li, S.; Chai, Z.; Ren, G. Exploring the Species Richness Pattern and Areas of Endemism of Tenebrionidae (Coleoptera) in Xinjiang, China. *Diversity* 2022, 14, 558. https:// doi.org/10.3390/d14070558

Academic Editor: Hongzhang Zhou

Received: 17 June 2022 Accepted: 10 July 2022 Published: 11 July 2022

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). may be affected by the above events to varying degrees in this area [20]. Therefore, it is of significance to study the distribution pattern of Tenebrionidae in the area to identify priority areas for biodiversity conservation [23,27–30]. Some studies have focused on the taxonomy [31,32], adaptive characteristics [17,33], diversity analysis [34], and molecular level [35–37] of tenebrionid beetles, but no formal biogeographical methods have been used to analyse Tenebrionidae in Xinjiang.

The study of the distribution patterns of species richness and endemic areas have always been central issues in biogeography [38–44], which not only represent the highest degree of the historical and ecological imprint of all biological entities [45] but also play an important role in the exploration of biodiversity hotspots [46–51].

Species richness, defined as the number of species in each cell, is the most basic and important indicator for identifying biodiversity hotspots and endemic areas [52,53]. Several studies have been undertaken using species richness as a measure to understand what affects biodiversity [10,48,51,53,54]. However, because species richness is easily affected by external factors (e.g., anthropogenic factors, geographic isolation, environmental change, etc.), simple species richness alone cannot reflect the true level of species spatial distribution [55], and the highest level of species richness does not necessarily mean the highest level of endemism [51]. Therefore, a new index called the areas of endemism (AOEs) has also been used to explore the distribution patterns of organisms [51,53,56].

Recently, a number of biogeographic and evolutionary studies have implemented and applied AOEs to resolve practical issues [48,49,51–53,56–58]. Although the definition of AOEs has been controversial in the past, it is now generally accepted as the distribution of two or more endemic species [59–62]. AOEs are entities compared in terms of ascertaining earth history based on biological patterns [62,63]. AOEs are important because they represent the operational unit of historical biogeography, which constitutes the first step for biogeographical regionalization [59,64]. In addition, AOEs show unique biota and are considered significant priority areas for biodiversity conservation [65]. Numerous biogeographers and ecologists have shown interest in evaluating the causes for the presence of AOEs [57,58,64,66].

Currently, although a number of alternative protocols and algorithms have been proposed to delimit AOEs [62,67–70], no consensus has emerged on the most appropriate method for detecting endemism. In some cases, more complete results can be obtained by combining two or multiple methods [62,71]. Among several approaches, two of them are most commonly used in exploring AOEs [51,53,56,72,73]. One approach is parsimony analysis of endemicity (PAE), which identifies ecological affinities between species by analysing species composition at local or regional scales [74,75]. The branching diagram is obtained through analysis, and the distance on the branching represents the size of ecological affinity [76]. The other approach is endemicity analysis (EA). Species distribution is analysed in different grid sizes by using the latitude–longitude information of species [72]. The algorithm assigns a score to each species by comparing the composition of species in each grid. The grid score is the sum of the scores of all species in a grid and is used to determine the common area [74].

In the present study, we collected the distribution information for a total of 556 Tenebrionidae species in Xinjiang based on both the literature and the examined material and then analysed the data with an analysis of endemicity using two different methods, aiming to (i) identify the centre of species richness and AOEs; (ii) explore the consistency of species richness and endemic patterns; and (iii) test the hypothesis of whether the centre of species diversity in previous studies is located in mountainous areas [53,56].

2. Materials and Methods

2.1. Species Distribution Data

In the present study, the georeferenced records of 556 species of Tenebrionidae were obtained from published papers [12,18,19,77–86], books [13–15], museum specimens (The Museum of Hebei University), and iNaturalist (https://www.inaturalist.org/ (accessed

on 15 June 2022)), and a total of 2322 distribution records were compiled in a geographic database. The distribution records without locations were excluded, and those with a lack of or imprecise coordinates were supplemented and standardized via Google Maps. Ultimately, the distribution information of 433 species with 2226 geographical records with robust coordinates was retained in the following analysis.

2.2. Assessing Sampling Bias and Mapping Species Richness

ArcGIS 10.8 was used to process the latitude and longitude geographic information of the species. In a $1^{\circ} \times 1^{\circ}$ cell grid, 115 grid cells with information were obtained, and different colours were used to replace the species richness of different degrees. The incidentbased bootstrap estimators were used to construct the species accumulation curve, which was designed to assess species inventory integrity in the study area [51]. EstimateS v9.1 was used to perform 100 randomized matrix analyses, where a matrix was created for the presence (1) or absence (0) of each species in a 1° grid size [87]. In addition, the number of records and the richness of the 1° grid were converted using square roots. Then, a linear regression was fitted to explore the completeness of richness, following previous methods [56].

2.3. Identifying Areas of Endemism

Two methods were used to explore the AOEs of Tenebrionidae: parsimony analysis of endemicity (PAE) and endemicity analysis (EA). In addition, three different grid sizes were used: $0.5^{\circ} \times 0.5^{\circ}$, $1^{\circ} \times 1^{\circ}$, and $1.5^{\circ} \times 1.5^{\circ}$.

For the PAE, matrices were created for the presence (1) or absence (0) of each species in three different grid sizes. Under the New Technical algorithms, TNT v1.5 was used to analyse all matrices, which added all zeros "Root" as the hypothetical outgroup of the tree [88]. The branches with relatively moderate bootstrap values (\geq 50%) were the candidates for AOEs [53]. Then, AOEs (clades of cells), comprising two or more endemic species restricted to these areas and at least two continuous cells, were mapped using ArcGIS to obtain the final results [52].

NDM/VNDM v3.1 was used for EA analysis under three grid sizes [88]. Due to the incompatibility of the input files, GeX was used to convert the latitude and longitude geographic information of the 433 species into XYD format [51,68,88]. The temporary set was saved with the current score in the 0.99 range. Sets were preserved with two or more endemics species with scores above 3.0. The search was repeated 100 times. Overlapping subsets were maintained when 50% of species were unique [74]. Species with a minimum score of 0.4 were selected in the obtained subsets [89]. Based on strict rules, the consensus area was calculated with a cutoff of 100% similarity in species. Other parameters were applied by using the default value. Finally, consensus areas were overlapped in different grid sizes to obtain the AOEs, and ArcGIS was used to draw the final results [51,53,72,90].

3. Results

3.1. Species Richness Pattern

Inadequate collection is a potential problem in biogeographic research, which may lead to the misidentification of biodiversity hotspots [30,91]. In this study, the 1° grid size species accumulation curves did not show inadequate collection (bootstrap mean approximately 505), with the data integrity for analysis as 86.1% (Figure 1A). The ratio of observed species richness to those predicted by the linear regression models for each grid cell was >69.3% (Figure 1B). This indicates that the data collected were sufficient. The variation in the number of species was well explained by changes in the number of species collected.



Figure 1. (A) Species accumulation curves for Tenebrionidae in Xinjiang, China. (B) Linear regression (y = 0.8846x + 0.2327) for the number of records and species richness in a 1° grid. The observed richness in the cells below the regression line was >69.3% of the predicted richness.

Within Xinjiang, most species were found in the Altai Mountains, Tianshan Mountains, and Kunlun Mountains (Figure 2). Among them, 118 species were found in the Altai Mountains, 340 in the Tianshan Mountains, and 80 in the Kunlun Mountains (Table S1).



Figure 2. (**A**) The general distribution pattern of Tenebrionidae in Xinjiang, China. (**B**) The species richness pattern identified by a 1° grid size in Xinjiang, China.

3.2. Parsimony Analysis of Endemicity

The four most parsimonious trees for AOE identification were obtained under three different grid cells. The optimal tree at a 1.5° grid size is a candidate for AOE identification (Figure 3). The branches with at least two consecutive cells were considered AOEs. Finally, 10 branches met the criteria and were selected, two of which belonged to Altay Prefecture–Akxoki Prefecture, two to Bortala Mongolian Autonomous Prefecture–Akxoki Prefecture, two to Kizilsu Kirgiz Autonomous Prefecture–Aksu Prefecture, two to Kashi Prefecture, and one to Hotan prefecture.



Figure 3. The optimal tree obtained by PAE using TNT v1.5 under a 1.5° grid size. Different coloured shadows represent different AOEs.

Under the criteria of identification, six AOEs were obtained (Table S2 and Figure 4): (i) Altay–Akxoki (AA), which shared 135 species; (ii) Bortala–Akxoki–Changji–Ili (BACI), which shared 264 species; (iii) Hami Prefecture (HP), which shared 21 species; (iv) Kizilsu Kirgiz–Aksu (KKA), which shared 73 species; (v) South of Kashgar (SK), which shared 41 species; and (vi) South of Hotan (SH), which shared 7 species.

3.3. Endemicity Analysis

In the EA analysis, three sizes were analysed by NDM/VNDM v3.1 under 0.5° , 1° , and 1.5° grid cells, and 3, 12, and 25 consensus areas were obtained, respectively. The consensus areas under the 0.5° grid size were only related to the Tianshan Mountains (TM) (Figure S1), and those under a $1^{\circ} \times 1^{\circ}$ cell grid (Figures S2 and S3) were similar to those under the 1.5° grid (Figures S4–S7); both were related to the Altai Mountains (AM), Tianshan Mountains (TM), Kunlun Mountains (KM), and Junggar Basin (JB) (Figure 5). By overlapping the consensus areas with different grid sizes, a total of four AOEs were finally detected.



Figure 4. AOEs obtained for Tenebrionidae by PAE using three different cells in Xinjiang, China: (A) 0.5° grid size, (B) 1° grid size, and (C) 1.5° grid size.



Figure 5. AOEs obtained for Tenebrionidae by EA analysis using three different cells in Xinjiang, China: (**A**) 0.5° grid size (red shadows); (**B**) 1° grid size (green shadows); and (**C**) 1.5° grid size (blue shadows).

(i) Altai Mountains (AM): The AM included consensus areas 6, 8, 9, 10, 36, and 40. The consensus regions scored 3.250000, 3.916667, 4.071429, 3.357143, 6.966667, and 3.000000 (Table S3 and Figures S2, S3, S6 and S7). One hundred and eleven species were found in both areas, including *Scythis altaicus*, *S. affinis*, *S. arenarius arenarius*, *S. vitorovianus*, *S. intermedia intemedia*, *Anatolica dashidorzsi subalpina*, *A. subpolita*, etc.

(ii) Tianshan Mountains (TM): These AOEs included consensus areas 1, 2, 3, 4, 5, 7, 11, 13, 14, 15, 16, 18, 20, 22, 24, 25, 26, 27, 28, 29, 30, 32, 33, 34, 37, 38, and 39 (Table S3 and Figures S1–S6). The consensus area scores were 3.70000, 3.250000, 3.000000, 3.666667, 4.375000, 3.928571, 3.000000, 3.458974, 3.000000, 3.333333, 5.000000, 9.869374, 7.947520, 10.089686, 3.000000, 6.558036, 5.207238,

7.365209, 3.333333, 6.672132, 3.230769, 6.400000, 7.418164, 10.898350, 5.583333, 4.641865, and 3.000000. Three hundred seventy-five species were counted in both regions, for example, *Colposphena brevlcollis, Scythis banghaasi, S. bulganicus, S. tatarica pseudoscythis, S. vitorovianus, S. angusticollis angusticollis, S. intermedia intemedia, Anatolica gobioltaica altaica, and A. hoboksarana.*

(iii) Kunlun Mountains (KM): The consensus areas 12, 31, and 35 were covered by KM, scoring 3.541667, 3.871429, and 5.250000, respectively (Table S3 and Figures S3, S5 and S6). Eighty-eight species were supported in these AOEs, including *Anatolica sternalis*, *Ascelosodis concinnus*, *Microdera* (*Microdera*) parvicollis, M. (*Microdera*) laticollis laticollis, M. (Dordanea) elegans, M. (Dordanea) mongolica, Colposcelis (Scelocolpis) damone, and C. (Scelocolpis) forsteri.

(iv) Junggar Basin (JB): This region contained consensus areas 17, 19, 20, 21, 23, 27, and 33 (Table S3 and Figures S4–S6). The scores were 4.783780, 5.188492, 7.947520, 3.500000, 6.387500, 7.365209, and 7.418164. A total of two hundred and twenty-one species were supported in these consensus areas, e.g., *Scythis altaicus, S. banghaasi, S. sculptilis, S. affinis, S. arenarius arenarius, S. vitorovianus, Anatolica dashidorzsi subalpina*, and *A. subpolita*.

4. Discussion

4.1. Congruent Patterns between Species Richness and Endemism Areas

In general, the species richness centres and endemism areas in this study were mainly found in the Altai, Tianshan, and Kunlun Mountains and Junggar Basin, which indicated that the species richness pattern of Tenebrionidae was basically consistent with the AOEs. This condition is found not only in insects [48,51,53,58,73] but also in other groups, such as plants [91], mammals [72], and birds [92]. This supports the hypothesis that AOEs have historically served as speciation centres [51,53], because AOEs characterized by stable climates and diverse habitats could maintain the long-term existence of organisms [51,53,72,93]. It is worth noting that there is an AOE in the basin area. In the centre of the basin, the vegetation coverage is lower than that around the basin, resulting in a more arid environment [94]. However, the Tenebrionidae there have evolved and adapted to the arid and semiarid environment in morphology, biology, and behaviour, which contributed to the high species richness in the middle of the basin [17,23,28,95].

4.2. AOEs of Tenebrionidae

Here, Tenebrionidae beetles were selected as the subjects to detect AOEs in Xinjiang for the first time, and a total of four AOEs were detected by two different methods, including three mountain regions (AM, TM, and KM) and one basin (JB).

4.2.1. AOEs in Montane Areas

Mountains often harbour high biodiversity due to their complex topography and stable climate. The formation of mountains is geological uplift, which is usually caused by the collision of continental plates [96,97]. The complex terrain and diverse environments in montane areas hinder the exchange of species, but they also provide the basis for the emergence of new species [98,99]. In addition, the complex terrain increases the number of ecological niches, which also promotes the formation of new species [99–101].

(i) AA belongs to the Altai Mountains. There was no significant difference in the AOEs between the PAE and EA under a 1.5° grid size, while the PAE had a larger consensus region than the EA under a 0.5° grid size. Under a $1^{\circ} \times 1^{\circ}$ cell grid, the EA detected larger AOEs, which were mainly located in the southern Altai Mountains. The Altai Mountains have a northwest to southeast trend and cross China diagonally. The Altai Mountains first appeared during the Caledonian movement, and the Himalayan movement caused the Altai Mountains to rise along the NW-trending fault block displacement, which provided shelters for several species and contributed to the high biodiversity of the areas [102,103].

(ii) BACI, HP, and KKA belong to the Tianshan Mountains. The EA detected more consensus areas in the region. Compared to PAE, the AOEs detected by EA were more widely distributed, mainly located in the southeastern and southwestern Tianshan Mountains. Significant differences could be observed in the 1.5° grid size. This may be caused by the different height gradients of the crustal changes in the late Cenozoic [104]. Tianshan

Mountain is located in the hinterland of Eurasia running from east to west and accounts for approximately one-third of the entire area of Xinjiang. The annual precipitation gradually decreases from west to the east on the same slope, which enhanced the adaptation of Tenebrionidae to the special climate and provided a certain guarantee for their survival, reproduction, and evolution [105,106].

(iii) The Kunlun Mountains include SK and SH. Within the 1° and 1.5° cells, the consensus area was mostly distributed in the western part of the northern slope of the Kunlun Mountains. Although the consensus areas of the EA and PAE were similar, the AOEs identified by the PAE were more widely distributed in this area, which is mainly located in the middle of the northern slope of the Kunlun Mountains. The neotectonic movement of the Kunlun Mountains is extremely strong, showing high values in the west and low values in the east [107]. The northern slope belongs to the Tarim Desert and Qaidam Desert in the warm temperate zone with low precipitation. With the increase in altitude, its terrain transitions from warm temperate desert into alpine desert, and the precipitation in this area also increases [25,99]. The higher topography of the west provides greater opportunities for the geographical isolation of species, which may be one reason why endemic areas are widespread in the west [108–110].

In addition, the high level of AOEs is closely related to the geological events experienced by the region [56]. The collision of the Indian plate with the Asian plate caused the Kunlun Mountains and the Qinghai-Tibet Plateau uplift, which is known as 'the roof of the world' [51,99,111,112]. Because the Qinghai–Tibet Plateau has affected the northwards movement of westerlies and the blocking of water vapour; hence, warm and wet air cannot reach Xinjiang across the Qinghai–Tibet Plateau, which leads to the arid climate in Xinjiang [4,5,24,99,106,113]. Tenebrionidae has adapted to arid environments, which contributes to the high species richness in this region [17,23].

4.2.2. AOE in the Basin

Basins with high surroundings (mountains or plateaus) and low central areas (plains or hills) can be divided into two types according to the influence of special geology and surface external force changes [114,115]. The Junggar Basin was formed as a result of plate movement [100,103,116,117]. Although the vegetation coverage of the basin is low, the surrounding mountains or plateaus have higher plant coverage than the central region [94], which provides certain environmental conditions for the existence of organisms.

(iv) The Junggar Basin (JB) is the second largest inland basin in China located in the northern part of Xinjiang. The basin is located between the Altai Mountains and the Tianshan Mountains, with the north being slightly higher than the south [94,103]. In addition, special geographical conditions prevent water vapour from moving northwards, which causes climatic changes and produces many arid and semiarid regions [1]. The Gurbantunggut Desert lies in the middle of the basin. The flourishing of the Tenebrionidae in this region is facilitated by adapting to the environment in arid and semiarid regions. In terms of morphology, beetles have adapted to merged anterior wings, degeneration of posterior wings, variable legs, formation of a subelytral cavity, and a well-developed tarsus [17]. Meanwhile, in biology and behaviour, they have suspended animation and selfdefence through the release of smelly fluids, gregariousness, and diurnal activity [17,23].

4.3. Limitations of Dataset and Methods

Although the taxonomy of the Tenebrionidae from Xinjiang, China, is well studied and documented with rich geographical data, under-collecting remains a potential problem in biogeographical research. Due to the harsh environment and technological limitations, we were unable to make a systematic survey for the species distribution of each cell. However, the data integrity for analysis showed an adequate collection. This suggests that the geographic data we collected were sufficient, and the variation in numbers of species was explained well by the variation in the numbers of collections [51].

In general, although slight differences were shown in the two approaches, both of them detected similar AOEs. The results of the EA had one more AOE than those of the PAE method, which might be explained by the different algorithms of the two methods [51]. There is no widely accepted answer as to which method can identify AOEs more accurately [62,118], and more accurate AOEs cannot be obtained by using a fixed method [53,56,93,119]. Thus, we adopted the PAE and EA results to provide more comprehensive AOE information.

Moreover, the grid sizes are also an important factor affecting the results of AOEs [72,93]. The smaller grid sizes will produce narrow and accurate consensus areas, but too small may also lead to the fragmentation of consensus areas [51], such as 0.5° in the present study. However, a grid size that is too large may also cause inaccurate consensus areas, such as a 1.5° grid size. Therefore, in the present study, we overlapped the consensus areas by different methods and three grid sizes to obtain more comprehensive endemism areas, following previous studies [51,56].

5. Conclusions

The geographical patterns of species richness and endemic areas of Tenebrionidae in Xinjiang, China, were analysed using the PAE and EA. We found that the species richness pattern was consistent with the AOEs of Tenebrionidae. In addition, the AOEs of Tenebrionidae in Xinjiang were mostly located in mountainous areas and basins, such as the Altai Mountains, Tianshan Mountains, Kunlun Mountains, and Junggar Basin. This is mainly due to the complex terrain and stable climate of the mountainous area, which promote long-term persistence, speciation, and species accumulation. Our findings indicate that greater conservation efforts should be expended in montane areas. Future studies should explore the relation between the AOEs and evolution histories at the molecular level.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/d14070558/s1, Table S1: A list of Tenebrionidae distributed in Altai Mountains, Tianshan Mountains and Kunlun Mountains; Table S2: Summary information of areas of endemism (AOEs) of Tenebrionidae using Parsimony analysis of endemicity (PAE) based on 1.5° cell grid; Table S3: Summary information on the consensus areas of Tenebrionidae using endemicity analysis (EA), with their respective score, number of cells for each consensus area, the maximum scores and grid size of each consensus areas; Figure S1: Consensus areas 1–3 detected by endemicity analysis (EA) using 0.5° grid size; Figure S2: Consensus areas 4–9 detected by endemicity analysis (EA) using 1° grid size; Figure S3: Consensus areas 10–15 detected by endemicity analysis (EA) using 1° grid size; Figure S4: Consensus areas 16–23 detected by endemicity analysis (EA) using 1.5° grid size; Figure S4: Consensus areas 24–31 detected by endemicity analysis (EA) using 1.5° grid size; Figure S6: Consensus areas 32–39 detected by endemicity analysis (EA) using 1.5° grid size; Figure S7: Consensus areas 40 detected by endemicity analysis (EA) using 1.5° grid size;

Author Contributions: Conceptualization, Y.L.; methodology, Y.W.; software, Y.W.; formal analysis, Y.L. and Y.W.; investigation, G.R. and Y.L.; resources, Y.L. and H.Z.; data curation, S.L. and Z.C.; writing, Y.L.; supervision, G.R. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by the National Natural Science Foundation of China (Grant No. 31970452) and the Key Project of Science-Technology Basic Condition Platform from The Ministry of Science and Technology of the People's Republic of China (Grant No. 2005DKA21402).

Institutional Review Board Statement: Not applicable.

Data Availability Statement: The original contributions presented in the study are included in the article and the Supplementary Files.

Acknowledgments: We are grateful to Tong Liu (Hebei University, Baoding) for his valuable help in the PAE and EA. We highly appreciate the help we received from Yuxia Yang (Hebei University, Baoding). We also thank the anonymous reviewers for their constructive comments.

Conflicts of Interest: The authors declare no conflict of interest.

References

- 1. Li, Z.; Chen, Y.N.; Shen, Y.J.; Liu, Y.B.; Zhang, S.H. Analysis of changing pan evaporation in the arid region of Northwest China. *Water Resour. Res.* **2013**, *49*, 2205–2212. [CrossRef]
- 2. Pi, H.; Sharratt, B.; Lei, J. Atmospheric dust events in central Asia: Relationship to wind, soil type, and land use. *J. Geophys. Res. Atmos.* **2017**, *122*, 6652–6671. [CrossRef]
- 3. Yuan, F.C.; Yang, F.X. The basic geomorphologic characteristics of Xinjiang, China. Arid Land Geogr. 1990, 13, 1–5. [CrossRef]
- 4. Zhang, X.J.; Wang, J.; Huang, G.; Chen, Y.H.; Yang, L.M.; Li, H.J.; Li, M.; Zheng, N. Spatiotemporal distribution of cloud liquid water volume over three main mountains in Xinjiang. *Arid Zone Res.* **2018**, *35*, 846–854. [CrossRef]
- 5. Liu, B.; Feng, J.M.; Ma, Z.G.; Wei, R.Q. Characteristics of climate changes in Xinjiang from 1960 to 2005. *Clim Environ. Res.* 2009, 14, 414–426.
- 6. Zhang, Z.W.; Yang, F.X.; Wu, J.L.; Zhou, J.; Yin, H.Y. Spatial Distribution Patterns and Type Structure of the Deserts in Xinjiang. *Arid Zone Res.* **2014**, *31*, 763–770. [CrossRef]
- 7. Hu, D.G. Multivariate Spatiotemporal Kriging Interpolation and Spatiotemporal Analysis of Precipitation in Xinjiang. Ph.D. Thesis, Wuhan University, Wuhan, China, 2019.
- 8. Huang, J.; Liu, C.; Guo, Z.; Ma, K.; Zang, R.; Ding, Y.; Lu, X.; Wang, J.; Yu, R. Seed plant features, distribution patterns, diversity hotspots, and conservation gaps in Xinjiang, China. *Nat. Conserv.* **2018**, *27*, 1–15. [CrossRef]
- 9. Feng, L.J.; Chu, J.Y.; Gao, L.; Zhou, L.; Wang, J. *Cerasus tianschanica* community characteristics and flora comparison in Ili and Tacheng regions of Xinjiang. *Non-Wood For. Res.* **2021**, *39*, 156–164. [CrossRef]
- 10. Li, J.J.; Li, Q.; Wu, Y.X.; Ye, L.Q.; Liu, H.H.; Wei, J.F.; Huang, X.L. Mountains act as museums and cradles for hemipteran insects in China: Evidence from patterns of richness and phylogenetic structure. *Global. Ecol. Biogeogr.* **2021**, *30*, 1070–1085. [CrossRef]
- 11. Wu, J.; Li, H.; Wan, H.; Wang, Y.; Sun, C.; Zhou, H. Analyzing the Relationship between Animal Diversity and the Remote Sensing Vegetation Parameters: The Case of Xinjiang, China. *Sustainability* **2021**, *13*, 9897. [CrossRef]
- 12. Li, X.M.; Li, J.; Pan, Z. New species and new faunistic records of the family Meloidae Gyllenhal, 1810 (Coleoptera: Tenebrionoidea) from China, with a list of meloid species from Xinjiang. *J. Asia-Pac. Entomol.* **2020**, *23*, 1144–1150. [CrossRef]
- 13. Ren, G.D.; Yang, X.J. Classificatory Account. In *Fauna of Soil Darkling Beetles in China*; Lin, L., Zhang, Y., Eds.; Science Press: Beijing, China, 2006; Volume 1, pp. 36–201.
- 14. Ren, G.D.; Ba, Y.B. Classificatory Account. In *Fauna of Soil Darkling Beetles in China*; Han, X.Z., Chen, L.Z., Eds.; Science Press: Beijing, China, 2010; Volume 2, pp. 26–188.
- 15. Ren, G.D.; Ba, Y.B.; Liu, H.Y.; Niu, Y.P.; Zhu, X.C. Classificatory Account. In *Fauna Sinica Insecta*; Wang, J., Zheng, J.H., Eds.; Science Press: Beijing, China, 2016; Volume 63, pp. 84–442.
- 16. Iwan, D.; Lobl, I. Tenebrionidae. In *Catalogue of Palaearctic Coleoptera*; Koninklijke Brill NV Press: Leiden, The Netherlands, 2020; Volume 5, pp. 104–474.
- 17. Ren, G.D.; Yu, Y.Z.; Ma, F. Desert Environment and Adaptation of Darkling Beetles. J. Agric. Sci. 1993, 14, 85–92.
- 18. Xu, J. Taxonomic Study on the Genus *Bioramix* Bates from China (Coleoptera: Tenebrionoidea: Platyscelidini). Master's Thesis, China West Normal University, Nanchong, China, 2017.
- 19. Zhang, C.L. Faunistic and Evolution on Tenebrionidaeof the Arid and Semi-Arid Region of China. Master's Thesis, Hebei University, Baoding, China, 2010.
- Lescano, M.N.; Elizalde, L.; Werenkraut, V.; Pirk, G.I.; Flores, G.E. Ant and tenebrionid beetle assemblages in arid lands: Their associations with vegetation types in the Patagonian steppe. J. Arid Environ. 2017, 138, 51–57. [CrossRef]
- 21. Li, X. Classification of Stored Products Darkling Beetles in China (Coleoptera: Tenebrionidae). Master's Thesis, Hebei University, Baoding, China, 2020.
- 22. Bouchard, P.; Bousquet, Y.; Davies, A.E.; Alonso-Zarazaga, M.A.; Lawrence, J.F.; Lyal, C.H.C.; Newton, A.F.; Reid, C.A.M.; Schmitt, M.; Slipinski, S.A.; et al. Family-group names in Coleoptera (Insecta). *Zookeys* **2011**, *88*, 1–972. [CrossRef]
- 23. Yang, G.J.; Wang, Y.; Wang, M.; Jia, L. Niche and interspecific association of darkling beetles in a desert grassland of alluvial fans in Helan Mountain, northwestern China. *Acta Entomol. Sin.* **2021**, *64*, 840–850. [CrossRef]
- 24. Li, H.H.; Min, Y.; Li, A.B.; Li, R.Q. Comparative analysis of on water vapor characteristics of two extreme rainstorms in the north slope of Kulun Mountains. *Arid Land Geogr.* 2022, 45, 715–724. [CrossRef]
- 25. Zhang, J.L.; Li, W.; Zhang, Y.L. Weather Classification and Radar Echo Characteristics of Short-term Heavy Precipitation in the Northern Kunlun Mountains. *Desert Oasis Meteorol.* 2022, *16*, 1–9. [CrossRef]
- Cui, C.; Guo, Y.; Shen, Y.J. Spatio-temporal variation in and the driving factors of desert vegetation in Xinjiang. *Chin. J. Eco-Agric.* 2021, 29, 1668–1678. [CrossRef]
- 27. Ren, G.D.; Yu, Y.Z. *The Darkling Beetles from Deserts and Semideserts of China (Coleoptera: Tenebrionidae);* Hebei University Press: Baoding, China, 1999; pp. 37–346.
- 28. Ayal, Y. Trophic structure and the role of predation in shaping hot desert communities. J. Arid Environ. 2007, 68, 171–187. [CrossRef]
- 29. Sun, X.J.; Ren, G.D. Analysis of Diversities and Faunal Composition of Darkling Beetles (Tenebrionidae, Coleoptera) in Inner Mongolia. J. Inn. Mong. Univ. (Nat. Sci. Ed.) 2015, 46, 541–547. [CrossRef]
- 30. Liu, J.L.; Ba, Y.B.; Niu, R.X.; Li, F.R.; Zhao, W.Z. Ground beetle diversity and their value as bioindicators for desertification in a natural desert of the middle of the Hexi Corridor, Northwest China. *Acta Ecol. Sin.* **2021**, *41*, 5435–5445. [CrossRef]

- 31. Ferrer, J.; Yvinec, J.H. Revision of the tribe Lachnogyini Reitter, 1904 sensu nov. and description of a new genus and species from the desert of Taklamakan, China. Coleoptera: Tenebrionidae, Pimeliinae. *Ann. De La Soc. Entomol. De Fr.* **2004**, 40, 41–49. [CrossRef]
- 32. Bai, X.L.; Ren, G.D. Revision of the genus *Bioramix* Bates, 1879 (Coleoptera: Tenebrionidae: Platyscelidini) from China. *Zootaxa* 2020, 4815, 1–102. [CrossRef] [PubMed]
- Shao, X.L.; Niu, Y.P.; Ren, G.D. Morphological basis and biological significance of the healing of the elytra and the formation of subelytral cavity in desert beetles. *J. Environ. Entomol.* 2019. Available online: http://kns.cnki.net/kcms/detail/44.1640.Q.201902 21.1612.002.html (accessed on 16 June 2022).
- Bai, X.L.; Ren, G.D. The diversity and faunal composition of the Darkling beetles in Qinghai-Xizang Plateau and adjacent areas. J. Environ. Entomol. 2015, 37, 475–482. [CrossRef]
- 35. Liu, X.L. Molecular Systematic Study on Partial Darkling Beetles Basedon Sequences of Mitoehondrial 16S rDNA and 18S rDNA. Master's Thesis, Hebei University, Baoding, China, 2004.
- Li, J.Q.; Lu, X.Y.; Liu, X.N.; Ma, J. Cloning and expression profiling of an attacin gene in response to coldstress in the desert beetle Microdera punctipennis (Coleoptera: Tenebrionidae). Acta Entomol. Sin. 2013, 56, 1381–1390. [CrossRef]
- Ruan, M.G.; Li, J.Q.; Meng, S.S.; Ma, J. Cloning and expression profiling in response to low temperature of Ras GTPase-activating protein gene *MpRasGAP* in the desert beetle *Microdera punctipennis* (Coleoptera: Tenebrionidae). *Acta Entomol. Sin.* 2015, 58, 367–374. [CrossRef]
- 38. Laffan, S.W.; Crisp, M.D. Assessing endemism at multiple spatial scales, with an example from the Australian vascular flora. *J. Biogeogr.* **2003**, *30*, 511–520. [CrossRef]
- Orme, C.D.; Davies, R.G.; Burgess, M.; Eigenbrod, F.; Pickup, N.; Olson, V.A.; Webster, A.J.; Ding, T.; Rasmussen, P.C.; Ridgely, R.S.; et al. Global hotspots of species richness are not congruent with endemism or threat. *Nature* 2005, 436, 1016–1019. [CrossRef]
- Posadas, P.; Crisci, J.; Katinas, L. Historical biogeography: A review of its basic concepts and critical issues. J. Arid. Environ. 2006, 66, 389–403. [CrossRef]
- Sandel, B.; Arge, L.; Dalsgaard, B.; Davies, R.G.; Gaston, K.J.; Sutherland, W.J.; Svenning, J.C. The influence of Late Quaternary climate change velocity on species endemism. *Science* 2011, 334, 660–664. [CrossRef] [PubMed]
- 42. Huang, J.H.; Chen, B.; Liu, C.R.; Lai, J.S.; Zhang, J.L.; Ma, K.P. Identifying hotspots of endemic woody seed plant diversity in China. *Divers. Distrib.* 2012, *18*, 673–688. [CrossRef]
- 43. Feng, G.; Mao, L.F.; Sandel, B.; Swenson, N.G.; Svenning, J.C. High plant endemism in China is partially linked to reduced glacial-interglacial climate change. *J. Biogeogr.* 2016, 43, 145–154. [CrossRef]
- 44. Li, Y.; Chen, J.; Jiang, L.Y.; Qiao, G.X. Islands conserve high species richness and areas of endemism of Hormaphidinae aphids. *Curr. Zool.* **2017**, *63*, 623–632. [CrossRef]
- 45. Casazza, G.; Zappa, E.; Mariotti, M.; Medail, F.; Minuto, L. Ecological and historical factors affecting distribution pattern and richness of endemic plant species: The case of the Maritime and Ligurian Alps hotspot. *Divers. Distrib.* 2008, 14, 47–58. [CrossRef]
- 46. Myers, N.; Mittermeier, R.A.; Mittermeier, C.G.; Da Fonseca, G.A.B.; Kent, J. Biodiversity hotspots for conservation priorities. *Nature* **2000**, 403, 853–858. [CrossRef]
- Lamoreux, J.F.; Morrison, J.C.; Ricketts, T.H.; Olson, D.M.; Dinerstein, E.; McKnight, M.W.; Shugart, H.H. Global tests of biodiversity concordance and the importance of endemism. *Nature* 2006, 440, 212–214. [CrossRef] [PubMed]
- Huang, X.L.; Qiao, G.X.; Lei, F.M. Diversity and distribution of aphids in the Qinghai-Tibetan Plateau-Himalayas. *Ecol. Entomol.* 2006, *31*, 608–615. [CrossRef]
- 49. Huang, X.L.; Qiao, G.X.; Lei, F.M. Use of parsimony analysis to identify areas of endemism of Chinese birds: Implications for conservation and biogeography. *Int. J. Mol. Sci.* 2010, *11*, 2097–2108. [CrossRef]
- 50. Gomes-da-Silva, J.; Amorim, A.M.; Forzza, R.C. Distribution of the xeric clade species of pitcairnioideae (Bromeliaceae) in south America: A perspective based on areas of endemism. *J. Biogeogr.* **2017**, *44*, 1994–2006. [CrossRef]
- 51. Liu, T.; Liu, H.Y.; Wang, Y.N.; Xi, H.C.; Yang, Y.X. Assessing the Diversity and Distribution Pattern of the Speciose Genus *Lycocerus* (Coleoptera: Cantharidae) by the Global-Scale Data. *Front. Ecol.* **2022**, *10*, 794750. [CrossRef]
- 52. Huang, X.L.; Lei, F.M.; Qiao, G.X. Areas of endemism and patterns of diversity for aphids of the Qinghai-Tibetan Plateau and the Himalayas. *J. Biogeogr.* 2008, *35*, 230–240. [CrossRef]
- Gao, C.; Chen, J.; Li, Y.; Jiang, L.Y.; Qiao, G.X. Congruent patterns between species richness and areas of endemism of the Greenideinae aphids (*Hemiptera: Aphididae*) revealed by global-scale data. *Zool. J. Linn. Soc.-Lond.* 2018, 183, 791–807. [CrossRef]
- Zhao, Z.X.; Yang, L.; Long, J.K.; Chang, Z.M.; Zhou, Z.X.; Zhi, Y.; Yang, L.J.; Li, H.X.; Sui, Y.J.; Gong, N.; et al. Testing Seven Hypotheses to Determine What Explains the Current Planthopper (Fulgoridae) Geographical and Species Richness Patterns in China. *Insects* 2020, *11*, 892. [CrossRef] [PubMed]
- 55. Yao, C.; Wang, J.C.; Xi, C.B.; Qian, T.L.; Sheng, C.Y. Spatial pattern of species richness among terrestrial mammals in China. *Diversity* **2020**, *12*, 96. [CrossRef]
- 56. Zhao, Z.X.; Yang, L.; Long, J.K.; Chang, Z.M.; Zhou, Z.X.; Zhi, Y.; Yang, L.J.; Li, H.X.; Sui, Y.J.; Gong, N.; et al. Endemism patterns of Planthoppers (Fulgoroidea) in China. *Front. Ecol. Evol.* **2021**, *9*, 683722. [CrossRef]
- López-Pujol, J.; Zhang, F.M.; Sun, H.Q.; Ying, T.S.; Ge, S. Centres of plant endemism in China: Places for survival or for speciation? J. Biogeogr. 2011, 38, 1267–1280. [CrossRef]

[PubMed]

- 58. Yuan, S.; Huang, M.; Wang, X.S.; Ji, L.Q.; Zhang, Y.L. Centers of endemism and diversity patterns for Typhlocybine leafhoppers (Hemiptera: Cicadellidae: Typhlocybinae) in China. *Insect Sci.* 2014, 21, 523–526. [CrossRef]
- 59. Cracraft, J. Historical biogeography and patterns of differentiation within the South American avifauna: Areas of endemism. *Ornith. Monogr.* **1985**, *36*, 49–84. [CrossRef]
- 60. Harold, A.S.; Mooi, R.D. Areas of endemism: Definition and recognition criteria. Syst. Biol. 1994, 43, 261–266. [CrossRef]
- Morrone, J.J. Society of systematic biologists on the identification of areas of endemism. *Syst. Biol.* 1994, 43, 438–441. [CrossRef]
 Linder, H.P. On areas of endemism, with an example from the African Restionaceae. *Syst. Biol.* 2001, *50*, 892–912. [CrossRef]
- 63. Henderson, I.M. Biogeography without area? *Aust. Sust. Bot.* **1991**, *4*, 59–71. [CrossRef]
- 64. Tribsch, A. Areas of endemism of vascular plants in the Eastern Alps in relation to Pleistocene glaciation. *J. Biogeogr.* 2004, 31, 747–760. [CrossRef]
- 65. Silva, J.M.C.; Sousa, M.C.; Castelletti, C.H.M. Areas of endemism for passerine birds in the Atlantic forest, South America. *Glob. Ecol. Biogeogr.* **2004**, *13*, 85–92. [CrossRef]
- 66. Anderson, S. Area and endemism. Q. Rev. Biol. 1994, 69, 451-471. [CrossRef]
- Shi, G.R. Multivariate data analysis in palaeoecology and palaeobiogeography-a review. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 1993, 105, 199–234. [CrossRef]
- Szumik, C.A.; Cuezzo, F.; Goloboff, P.A.; Chalup, A.E. An optimality criterion to determine areas of endemism. *Syst. Biol.* 2002, 51, 806–816. [CrossRef] [PubMed]
- 69. Hausdorf, B.; Hennig, C. Biotic element analysis in biogeography. Syst. Biol. 2003, 52, 717-723. [CrossRef]
- 70. Morrone, J.J. Parsimony analysis of endemicity (PAE) revisited. J. Biogeogr. 2014, 41, 842–854. [CrossRef]
- 71. Carine, M.A.; Humphries, C.J.; Guma, I.R.; Reyes-Betancort, J.A.; Santos, G.A. Areas and algorithms: Evaluating numerical approaches for the delimitation of areas of endemism in the Canary Islands archipelago. *J. Biogeogr.* 2009, *36*, 593–611. [CrossRef]
- 72. Prado, J.R.; Brennand, P.G.G.; Godoy, L.P.; Libardi, G.S.; Abreu-Júnior, E.F.; Roth, P.R.O.; Chiquito, E.; Percequillo, A.R. Species richness and areas of endemism of oryzomyine rodents (Cricetidae, Sigmodontinae) in South America: Anndm/vndmapproach. *J. Biogeogr.* **2015**, *42*, 540–551. [CrossRef]
- Zhuang, H.; Yago, M.; Settele, J.; Li, X.; Ueshima, R.; Grishin, N.V.; Wang, M. Species richness of Eurasian Zephyrus hairstreaks (Lepidoptera: Lycaenidae: Theclini) with implications on historical biogeography: An NDM/VNDM approach. PLoS ONE 2018, 13, e0191049. [CrossRef] [PubMed]
- 74. Szumik, C.; Goloboff, P. Areas of endemism: An improved optimality criterion. Syst. Biol. 2004, 53, 968–977. [CrossRef]
- 75. Nihei, S.S. Misconceptions about parsimony analysis of endemicity. J. Biogeogr. 2006, 33, 2099–2106. [CrossRef]
- 76. Zheng, W. Improvement and Optimization of Algorithms for Constructing Phylogenetic Trees based on Maximum Parsimony. Master's Thesis, University of Chinese Academy of Sciences, Guangzhou, China, 2013.
- 77. Ren, G.D.; Wang, X.M. Three new species of the genus *Adesmia* Fisch. from China (Coleoptera, Tenebrionidae, Adesmini). *J. Agric. Sci.* **1993**, *14*, 1–8.
- 78. Ren, G.D.; Zheng, Z.M. Six new species of the genus *Crypticus* Latr. from northwest of China (Coleoptera: Tenebrionidae). J. Agric. Sci. **1993**, 14, 9–23.
- 79. Ren, G.D.; Yu, Y.Z. Larvae-Monograph of the Genus *Cyphogenia* Solier, 1836 in China (Coleoptera: Tenebrionidae). *J. Hebei Univ.* **2000**, *20*, 52–57.
- Wu, W.; Fan, Z.T.; Huang, R.X. Three New Species of *Platyope* (Coleoptera: Tenebrionidae) from Xinjiang, China. *Entomotaxonomia* 2005, 27, 283–288.
- 81. An, W.T. Systematics of the Tribe Platyopini in China (Coleoptera:Tenebrionidae). Master's Thesis, Hebei University, Baoding, China, 2010.
- 82. Zhou, Y. Taxonomy of Lagriina from China (Coleoptera, Tenebrionidae, Lagriini). M.S. Thesis, Hebei University, Baoding, China, 2011.
- 83. Ba, Y.B. Systematics of Pimeliinae and its Distribution in China (Coleoptera: Tenebrionidae). Ph.D. Thesis, Hebei University, Baoding, China, 2012.
- Yang, X.J.; Ren, G.D. A New Record Genus and Species of Lachnogyini from China (Coleoptera:Tenebrionidae). Acta Zootax Onomica Sin. 2012, 37, 453–455.
- Bai, X.L. Systematic Taxonomy of Platyscelidini from China (Coleoptera: Tenebrionidae). Ph.D. Thesis, Hebei University, Baoding, China, 2020.
- 86. Ba, Y.B.; Ren, G.D.; Liu, J.L. Two new nocturnal species of the genus *Anatolica* (Coleoptera: Tenebrionidae: Pimeliinae) from the deserts of northwest China. *Entomotaxonomia* **2021**, *43*, 130–137. [CrossRef]
- 87. Colwell, R.K.; Elsensohn, J.E. EstimateS turns 20: Statistical estimation of species richness and shared species from samples, with non-parametric extrapolation. *Ecography* **2014**, *37*, 609–613. [CrossRef]
- Santos, C.M.D.; Fuhlendorf, M. GeX: An automated tool for generating XYD files for analysis of endemicity using VNDM. *Cladistics* 2019, 35, 125–129. [CrossRef] [PubMed]
- 89. Escalante, T.; Morrone, J.J.; Rodríguez-Tapia, G. Biogeographic regions of North American mammals based on endemism. *Biol. J. Linn. Soc.* 2013, 110, 485–499. [CrossRef]

- 90. Du, C.C.; Chen, J.; Jiang, L.Y.; Qiao, G.X. High correlation of species diversity patterns between specialist herbivorous insects and their specific hosts. *J. Biogeogr.* 2020, 47, 1232–1245. [CrossRef]
- 91. Moline, P.M.; Linder, H.P. Input data, analytical methods and biogeography of *Elegia* (Restionaceae). J. Biogeogr. 2006, 33, 47–62. [CrossRef]
- Jetz, W.; Rahbek, C.; Colwell, R.K. The coincidence of rarity and richness and the potential signature of history in centres of endemism. *Ecol. Lett.* 2004, 7, 1180–1191. [CrossRef]
- 93. Lago-Barcia, D.; DaSilva, M.B.; Conti, L.A.; Carbayo, F. Areas of endemism of land planarians (Platyhelminthes: Tricladida) in the Southern Atlantic Forest. *PLoS ONE* 2020, *15*, e0235949. [CrossRef]
- Zhang, Q.L.; Zhou, J.Y.; Dou, P.J.; Yue, S.R. Remote Sensing Monitoring of Spatiotemporal Changes of Vegetation Coverage in Junggar Basinduring 2000–2018. *Heilongjiang Sci.* 2020, 11, 8–10.
- 95. Ren, G.D.; Ji, Q.Q. Species diversity of the fungus-beetle and its relationship with fungi. *J. Hebei Univ. (Nat. Sci. Ed.)* **2019**, *39*, 166–174. [CrossRef]
- 96. Jansson, R. Global patterns in endemism explained by past climatic change. *Proc. Roy. Soc. B Biol. Sci.* 2003, 270, 583–590. [CrossRef]
- Gillespie, R.G.; Roderick, G.K. Evolution: Geology and climate drive diversifification. *Nature* 2014, 509, 297–298. [CrossRef] [PubMed]
- 98. Zhou, Y.D.; Chen, S.C.; Hu, G.W.; Mwachala, G.; Yan, X.; Wang, Q.F. Species richness and phylogenetic diversity of seed plants across vegetation zones of Mount Kenya, East Africa. *Ecol. Evol.* 2018, *8*, 8930–8939. [CrossRef] [PubMed]
- 99. Du, W.B. Patterns of Plant Diversity and Formation Mechanism in the Kunlun Mountains. Ph.D. Thesis, Lanzhou University, Lanzhou, China, 2021.
- 100. Tang, W.B.; Zhang, Y.Y.; Pe-Piper, G.; Piper, D.J.W.; Guo, Z.J.; Li, W. Permian rifting processes in the NW Junggar Basin, China: Implications for the post-accretionary successor basins. *Gondwana Res.* **2021**, *98*, 107–124. [CrossRef]
- 101. Du, W.B.; Jia, P.; Du, G.Z. Current biogeographical roles of the Kunlun Mountains. Ecol. Evol. 2022, 12, e8493. [CrossRef]
- Han, F.F.; Yan, J.J.; Ling, H.B. Variance of vegetation coverage and its sensitivity to climatic factors in the Irtysh River basin. *PeerJ* 2013, 9, e11334. [CrossRef]
- 103. Jia, C.Z.; Li, B.L.; Lei, Y.L.; Chen, Z.X. The structure of Circum-Tibetan Plateau Basin-Range System and the large gas provinces. *Sci. China Earth Sci.* 2013, *56*, 1853–1863. [CrossRef]
- 104. Fang, Y.A.; Wu, C.D.; Wang, Y.Z.; Hou, K.J.; Guo, Z.J. Topographic evolution of the Tianshan Mountains and their relation to the Junggar and Turpan Basins, Central Asia, from the Permian to the Neogene. *Gondwana Res.* 2019, 75, 47–67. [CrossRef]
- 105. Heng, T.; He, X.; Yang, L.; Yu, J.; Yang, Y.; Li, M. The Spatiotemporal Patterns and Interrelationships of Snow Cover and Climate Change in Tianshan Mountains. *Water* **2021**, *13*, 404. [CrossRef]
- 106. Guan, X.F.; Yao, J.Q.; Schneider, C. Variability of the precipitation over the Tianshan Mountains, Central Asia. Part II: Multidecadal precipitation trends and their association with atmospheric circulation in both the winter and summer seasons. *Int. J. Climatol.* **2022**, *42*, 139–156. [CrossRef]
- 107. Yu, X.J.; Guo, Z.J.; Chen, Y.X.; Du, W.; Wang, Z.D.; Bian, Q. River system reformed by the Eastern Kunlun Fault: Implications from geomorphological features in the Eastern Kunlun Mountains, Northern Tibetan Plateau. *Geomorphology* 2020, 350, 106876. [CrossRef]
- Leneveu, J.C.A.; Wahlberg, N. Varying rates of diversification in the genus *Melitaea* (Lepidoptera:Nymphalidae) during the past 20 million years. *Biol. J. Linn. Soc.* 2009, 97, 346–361. [CrossRef]
- 109. Condamine, F.L.; Rolland, J.; Hohna, S.; Sperling, F.A.H.; Sanmartin, I. Testing the role of the Red Queen and Court Jester asdrivers of the macroevolution of Apollo butterflies. *Syst. Biol.* **2018**, *67*, 940–964. [CrossRef] [PubMed]
- 110. Li, X.M. Phylogenetic and Historical Biogeography of Partial Genera of the Tribe Blaptini from the Qinghai-Xizang Plateau. Ph.D. Thesis, Hebei University, Baoding, China, 2020.
- 111. Zhang, Q.; Kong, D.D.; Shi, P.J.; Singh, V.P.; Sun, P. Vegetation phenology on the Qinghai-Tibetan Plateau and its response to climate change (1982–2013). *Agric. For. Meteorol.* **2018**, 248, 408–417. [CrossRef]
- 112. Duan, H.C.; Xue, X.; Wang, T.; Kang, W.P.; Liao, J.; Liu, S.L. Spatial and Temporal Differences in Alpine Meadow, Alpine Steppe and All Vegetation of the Qinghai-Tibetan Plateau and Their Responses to Climate Change. *Remote Sens.* **2021**, *13*, 669. [CrossRef]
- 113. Li, L.; Wang, Z.; Zerbe, S.; Abdusalih, N.; Tang, Z.; Ma, M.; Yin, L.K.; Mohammat, A.; Han, W.X.; Fang, J.Y. Species Richness Patterns and Water-Energy Dynamics in the Drylands of Northwest China. *PLoS ONE* **2013**, *8*, e66450. [CrossRef]
- 114. Wang, X.Y.; Lu, H.Y.; Vandenberghe, J.; Chen, Z.Y.; Li, L.P. Distribution and Forming Model of Fluvial Terrace in the Huangshui Catchment and its Tectonic Indication. *Acta Geolo. Sin-Engl.* **2010**, *84*, 415–423. [CrossRef]
- Heller, P.L.; McMillan, M.E.; Humphrey, N. Climate-induced formation of a closed basin: Great Divide Basin, Wyoming. *Geol. Soc. Am. Bull.* 2011, 123, 150–157. [CrossRef]
- 116. Yan, D.Z.; Xu, H.M.; Xu, Z.H.; Tian, M.; Lei, Z.C. Mesozoic multistage structural deformation along the Ke-Bai fault zone in the north-west Junggar Basin, NW China. *Geol. J.* **2019**, *55*, 4097–4111. [CrossRef]
- 117. Imin, A.; Zha, M.; Ding, X.J.; Bian, B.L.; Liu, Y.; Zheng, M.L.; Han, C. Identification of a Permian foreland basin in the western Junggar Basin (NW China) and its impact on hydrocarbon accumulation. *J. Geophys. Eng.* **2020**, *187*, 106810. [CrossRef]

- 118. Escalante, T. Parsimony analysis of endemicity and analysis of endemicity: A fair comparison. *Syst. Biodivers.* **2015**, *13*, 413–418. [CrossRef]
- 119. Garrafoni, A.R.S.; Moura, F.R.; Lourenco, A.P. Areas of endemism in the Atlantic Forest: Quantitative biogeography insights from orchid bees (Apidae: Euglossini). *Apidologie* 2017, 48, 513. [CrossRef]