

Article

Estimation of the Population Dynamics of *Taxus cuspidata* by Using a Static Life Table for Its Conservation

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Abstract: *Taxus cuspidata* is a rare and endangered plant species with an extremely small population which is endemic to China. This study focused on the natural *T. cuspidata* population in Jilin Province in China. Conventional population ecology survey methods were used to describe its population structure characteristics. Then, we chose diameter structure instead of temporal structure to establish a static population life table, draw a population survival curve, and quantify the future development trend by using population dynamic analysis and time sequence prediction. The results showed that: (1) the static life table suggested that the population of *T. cuspidata* was stable overall. The population survival curve tended to be Deevey II, with a high early seedling mortality, and the later population growth tended to be stable; (2) the survival curve suggested that the population initially experienced higher mortality rates in the early stage. However, as time progressed and the population aged, the mortality rates decreased, resulting in a more stable population in the middle and late stages; (3) The diameter class structure of *T. cuspidata* was stable overall, and the dynamic indices showed that the population was fluctuating. The population was influenced by external disturbances and showed some resistance to human disturbance; (4) time sequence prediction analysis showed that the mortality rate of young individuals was high, natural renewal could be maintained, and the population size would remain at a certain amount in the future. The result shows that the Jilin region is a highly suitable area for the growth of *Taxus cuspidata*'s population in Northeast China. We recommend in situ conservation of remaining wild populations, relocation of germplasm resources, and reduction of human activities; these actions will be beneficial to *Taxus cuspidata*'s long-term survival.

Keywords: *Taxus cuspidata*; plant species with extremely small populations (PSESPs); static life table; quantity dynamic; time sequence prediction



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1. Introduction

Taxus cuspidata Siebold et Zucc. is an arboreal species of Taxaceae. It is also a national first-level protected medicinal plant species. It is dioecious, with a height of up to 20 m and a DBH (diameter at breast height) up to 1 m [1]. It is a national first-level rare and endangered plant which is listed in the *Red Book of China's Rare and Endangered Plants and Plant Species with Extremely Small Populations (PSESP)* [2]. Its natural distribution range is relatively narrow, as it is only distributed at an altitude of 600–1200 m. Due to excessive logging and human disturbance, the population of *T. cuspidata* has decreased rapidly. *Taxus cuspidata* have strict requirements for their growth environment: they grow slowly in shady places when they are young and in open and sunny places when they mature. This special ecological adaptability allows them to survive in habitats that are sparse and severely fragmented [1–3]. Seeds can only be collected where there is a mix of males and females, and fruiting is sparse. During the fruiting period, a large number of seeds are eaten by some animals due to the slightly sweet and edible aril of the seeds, and their seeds are

physiologically dormant, requiring at least two winters and a summer to germinate under natural conditions. Additionally, the number of seeds is further reduced due to loss of vigor caused by rotting, drying out, and loss of water [1,4]. Therefore, it is difficult to obtain seedling renewal data for the population, and, due to the long-lived and slow-growing nature, we chose to investigate population dynamics by using diameter structure to represent age structure. In fact, population structure (age or size) of tree species is a well-known way to provide valuable information about population dynamics [5,6].

A population is the unit of reproduction and evolution of a species in nature, a collection of individuals of the same species capable of reproducing under certain spatial and temporal conditions. Population is not only a collection of individuals distributed within a certain geological period and geographic space [7], but also has its own unique structure. In most of the existing population dynamics studies, the future trend of population dynamics is judged by the magnitude of the population growth rate, which is determined by the vital rates such as survival probability, growth probability, and reproduction probability of individuals [8–10]. Due to the long-lived nature of *T. cuspidata* and seeds with notable dormancy, their natural reproduction probability and growth probability cannot be determined [1]. Since its extremely small populations are few in nature, each surviving individual is very precious. Harvesting the tree requires time and human and material resources and is not conducive to the ecological balance of the forest and restoration. So we chose to use diameter class structure to investigate the population structure and dynamics of *T. cuspidata*. We used diameter class structure to represent a time sequence for the woody plants to construct a static life table. This is particularly appropriate for endangered species. The static life table reflects the allocation of individuals with different sizes in a population to reflect the population dynamics and evolution [11]. Through the analysis of the static life table, we can see the population's diameter structure, and, by quantifying the dynamics, it is possible to grasp the status of the population, predict its development, and provide a scientific reference for the management of the population. Constructing a life table is another important way to understand the characteristics of population structure. The compilation of a static life table can measure the current status of the population and infer the past structure and possible historical structure [12] so as to predict the future trend of the population and the possibility of survival and reproduction under specific conditions. Demographic parameters are a function of the individual's age, size, and developmental state, or a combination of any of those. Knowledge about the contributions of different stages of the life cycle to population growth rate enhances our understanding of the life histories of species [13]. Population dynamics are the historical results of the interaction between population viability and environmental factors.

The endangered situation of *T. cuspidata* has attracted great attention from scholars, leading to studies of its genetic diversity [14], habitat adaptability evaluation [14], introductory suitability [15], and related nursery technology [16–18]. Studies on the ecological aspects of *T. cuspidata* populations have focused on population reproductive characteristics, survival environment [2], interspecific competition [19], population survival community biodiversity, and interspecific associations [6]. The survival and growth of the remaining population have been seriously threatened and urgently need to be protected [3]. *T. cuspidata* is mainly distributed in the three provinces of Heilongjiang, Jilin, and Liaoning, most of which comprise unsuitable habitats. Of a total area of 702,200 km², the area of suitable distribution is 104,100 km², which accounts for only 12.9% of the total area of the three provinces. Most of the suitable habitat area is located in Jilin Province, with 49,500 km² of low-suitability area and 33,800 km² of high-suitability area. Heilongjiang Province includes 12,900 km² of low-suitability area and 7500 km² of high-suitability area [3]. There have been studies on the growth status of *T. cuspidata* populations in Heilongjiang [2,20], but there is a lack of studies on the population status in Jilin, which is the most suitable area for *T. cuspidata*.

This study is based on a survey of the structural characteristics and population dynamics, which have been declining in recent years, of *T. cuspidata* in Northeast China.

In this paper, our focus is on revealing population dynamics and choosing appropriate mathematical analysis tools to reduce errors between age and diameter, including log-linear analysis [21,22] and algorithms to identify the number of size classes that minimize error [7,23]. In addition, the results are specific to the observed population dynamics pattern of variation [24]. Understanding the structure characteristics of the *T. cuspidata* population is a prerequisite for understanding population dynamics and survival and for constructing suitable habitats that are consistent with the species characteristics, providing a theoretical basis for in situ conservation, and promoting artificial population restoration and reconstruction [25]. The results of the analysis are presented in the following sections: (1) the current distribution of the population and the process of population formation; (2) predicting population growth and providing scientific advice for the conservation of the *T. cuspidata* population.

2. Materials and Methods

2.1. Sample Site Survey

On the basis of the recommendations of the Forestry Bureau and field surveys conducted in 2016 and 2017 in the distribution areas of *T. cuspidata* populations, the natural distribution of *T. cuspidata* was generally grasped, and representative sites in the protected areas of Helong Forestry Bureau and Wangqing Forestry Bureau were selected for community surveys. The region has a temperate continental monsoon climate with short summers and long winters. The average annual temperature is around 3.9 °C, the average annual rainfall is about 580 mm, the precipitation is mainly concentrated in May–August, the frost-free period is 110–141 days, and the annual sunshine hours are 2700 h. The soil is mainly dark brown loam with pH between 4.5 and 5.7, and the moisture content ranges from 10% to 40%. The types of forests are *Pinus koraiensis*, *Pinus densiflora*, *Abies holophylla* mixed forests, *Pinus koraiensis* mixed forests, *Abies holophylla*, *Pinus koraiensis*, and *Pinus tabulaeformis* mixed forests [26].

The habitat factors of altitude and inclination were investigated by setting up 20 plots, each measuring 40 m × 40 m. To collect data on the tree species present in these plots, a 5 m × 5 m sample area was designated using the adjacent grid method (Figure 1). Within each sample plot, the following information was recorded for each individual tree: species name, tree height, and diameter at breast height (DBH). Additionally, the number of seedlings and saplings below 4 cm in diameter at breast height was also noted. Details of the sample plots can be found in Table 1.

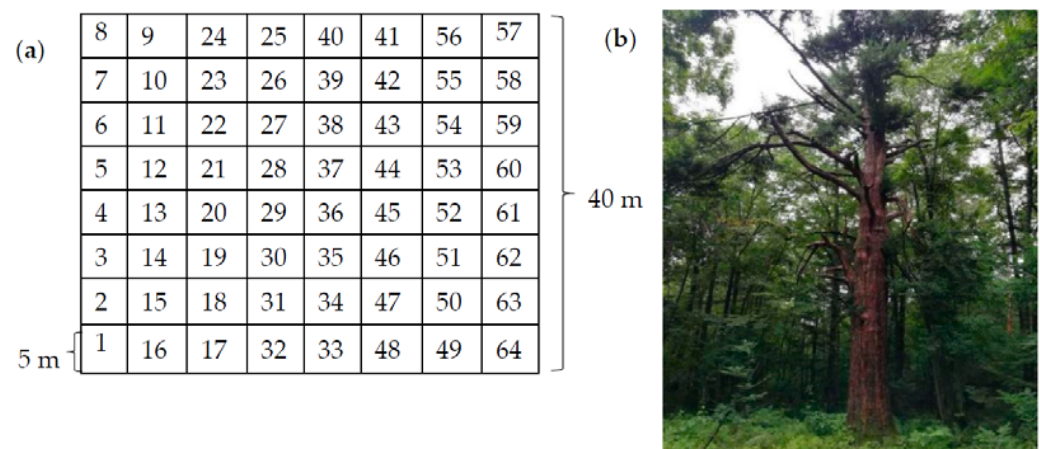


Figure 1. Spots-setting method (a) and photo of *Taxus cuspidata* (b).

Table 1. Basic information in the plots of the communities in which *Taxus cuspidata* populations are located.

Plot	Elevation (m)	Inclination	Location	Latitude and Longitude N E		Dominant Tree	Population Density (/ha)
1	866	26°	Huanggou, Helong	42°24'23"	128°38'4"	<i>Abies nephrolepis</i> , <i>Picea jezoensis</i> , <i>Tilia amurensis</i> , <i>Pinus koraiensis</i>	2.5
2	813	30°	Huanggou, Helong	42°24'24.9"	128°40'5.9"	<i>Tilia amurensis</i> , <i>Acer pseudosieboldianum</i> , <i>Abies nephrolepis</i> , <i>Acer ukurunduense</i>	1.94
3	1021	30°	Huanggou, Helong	42°24'27.9"	128°40'7.8"	<i>Acer ukurunduense</i> , <i>Abies nephrolepis</i> , <i>Acer tegmentosum</i> , <i>Tilia amurensis</i>	1.25
4	998	29°	Huanggou, Helong	42°24'28.5"	128°40'10.8"	<i>Acer ukurunduense</i> , <i>Abies nephrolepis</i> , <i>Pinus koraiensis</i> , <i>Tilia amurensis</i> , <i>Taxus cuspidata</i>	1.69
5	979	32°	Huanggou, Helong	42°24'06.7"	128°39'54.0"	<i>Tilia amurensis</i> , <i>Acer pseudosieboldianum</i> , <i>Abies nephrolepis</i> , <i>Acer ukurunduense</i>	3.56
6	1130	12°	Huanggou, Helong	42°23'58.8"	128°40'0.9"	<i>Pinus koraiensis</i> , <i>Acer ukurunduense</i> , <i>Abies nephrolepis</i> , <i>Tilia amurensis</i> , <i>Abies holophylla</i>	1.81
7	1030	23°	Huanggou, Helong	42°23'45"	128°39'56"	<i>Acer ukurunduense</i> , <i>Pinus koraiensis</i> , <i>Abies nephrolepis</i> , <i>Acer pseudosieboldianum</i> , <i>Tilia amurensis</i>	3.38
8	873	13°	Jingouling, Wangqing	43°22'29"	130°9'33.9"	<i>Abies nephrolepis</i> , <i>Taxus cuspidata</i> , <i>Tilia amurensis</i> , <i>Pinus koraiensis</i> , <i>Betula costata</i>	17.56
9	694	17°	Jingouling, Wangqing	43°21'37"	130°10'12.8"	<i>Abies nephrolepis</i> , <i>Acer pseudosieboldianum</i> , <i>Taxus cuspidata</i> , <i>Acer mono</i> , <i>Pinus koraiensis</i>	6.38
10	910	14°	Duhuangzi, Wangqing	43°12'24"	130°36'35.3"	<i>Abies nephrolepis</i> , <i>Acer ukurunduense</i> , <i>Tilia amurensis</i> , <i>Pinus koraiensis</i> , <i>Acer tegmentosum</i>	2.06
11	907	13°	Duhuangzi, Wangqing	43°12'22.7"	130°36'35.7"	<i>Abies nephrolepis</i> , <i>Tilia amurensis</i> , <i>Acer tegmentosum</i> , <i>Acer barbinerve</i> , <i>Acer ukurunduense</i>	1.5
12	856	21°	Duhuangzi, Wangqing	43°12'27.7"	130°36'36.7"	<i>Abies nephrolepis</i> , <i>Acer barbinerve</i> , <i>Pinus koraiensis</i> , <i>Tilia amurensis</i> , <i>Abies holophylla</i>	2.38
13	808	18°	Duhuangzi, Wangqing	43°12'23.0"	130°36'25.8"	<i>Abies nephrolepis</i> , <i>Acer barbinerve</i> , <i>Tilia amurensis</i> , <i>Acer tegmentosum</i> , <i>Acer ukurunduense</i>	0.81
14	818	18°	Duhuangzi, Wangqing	43°12'28.6"	130°36'25.3"	<i>Abies nephrolepis</i> , <i>Acer barbinerve</i> , <i>Tilia amurensis</i> , <i>Acer ukurunduense</i> , <i>Acer tegmentosum</i>	1.19
15	936	24°	Duhuangzi, Wangqing	43°11'51.3"	130°36'25.7"	<i>Abies nephrolepis</i> , <i>Tilia amurensis</i> , <i>Acer barbinerve</i> , <i>Acer ukurunduense</i> , <i>Acer tegmentosum</i>	1.75
16	900	3°	Huanggou, Wangqing	43°18'45.8"	130°19'31"	<i>Abies nephrolepis</i> , <i>Acer barbinerve</i> , <i>Acer ukurunduense</i> , <i>Tilia amurensis</i> , <i>Taxus cuspidata</i>	4.94
17	700	12°	Lanjia, Wangqing	43°26'19"	130°57'25"	<i>Acer barbinerve</i> , <i>Abies nephrolepis</i> , <i>Pinus koraiensis</i> , <i>Acer ukurunduense</i> , <i>Acer tegmentosum</i>	4
18	789	7°	Lanjia, Wangqing	43°26'19.4"	130°57'15.8"	<i>Acer barbinerve</i> , <i>Tilia amurensis</i> , <i>Abies nephrolepis</i> , <i>Acer ukurunduense</i> , <i>Pinus koraiensis</i>	0.75
19	766	27°	Lanjia, Wangqing	43°27'11.6"	130°56'12.4"	<i>Abies nephrolepis</i> , <i>Tilia amurensis</i> , <i>Acer tegmentosum</i> , <i>Acer barbinerve</i> , <i>Abies holophylla</i>	3.25
20	745	20°	Madida, Wangqing	43°9'35.5"	130°41'23.5"	<i>Abies nephrolepis</i> , <i>Acer tegmentosum</i> , <i>Tilia amurensis</i> , <i>Acer ukurunduense</i> , <i>Abies holophylla</i>	2.31

2.2. Diameter Classification

Age structure is an important characteristic of population, and many scholars have used size structure analysis [27–29] in their studies of population structure and dynamics; sometimes, size may be a better predictor of reproductive output than age under similar forest growth conditions of compared trees [30]. As an endangered species, due to the lack of data on wood analysis and the difficulty of drilling cores to obtain tree age, the age and diameter response of the same species in the same environment is characterized by the consistency of age and diameter [31]. Considering the life history characteristics and survey data of *T. cuspidata* and referring to other classification methods [32–35], we chose diameter to represent age in order to study the population dynamics in practical research. The distribution of diameter classes can provide valuable insights into the competitive dynamics and overall growth potential of a forest stand [36], according to one of the basic rules of stand structure [37,38]. Based on the life cycle characteristics and the R^2 between the smoothing curve and survival curve [10], we categorized a diameter class of 6 cm to compile static life table, draw survival curves, and analyze dynamic changes. Based on the life history features of *T. cuspidata*, those with DBH < 2.5 cm were classified as seedlings, those with 2.5 cm ≤ DBH < 4 cm as saplings, and those with DBH ≥ 4 cm as mature trees. The number of plants in each diameter class was counted in 6 cm diameter classes (upper limit exclusion method) to determine the diameter class structure (Table 2).

Table 2. The diameter classification table of *Taxus cuspidata* (upper limit exclusion method).

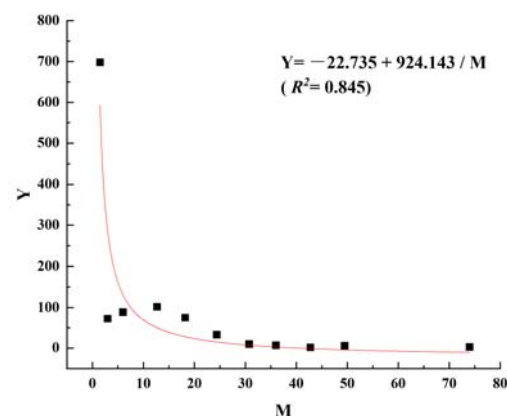
Breast Diameter (cm)	0–2.5	2.5–4	4–10	10–16	16–22	22–28	28–34	34–40	40–46	46–52	52+
Diameter class	1	2	3	4	5	6	7	8	9	10	11

2.3. Smoothing Curve

The wild population in the study is distributed in natural forest. Additionally, the survey method used to collect data may have introduced systematic errors, meaning that the results may be skewed or biased. Furthermore, the data may not fully satisfy the three assumptions necessary for constructing a life table. A life table is a tool used to analyze mortality and survival rates in different age groups of a population. The three assumptions are that the population is closed, that mortality rates are constant over time, and that each individual has an equal chance of dying. If these assumptions are not fully met, the data may show a higher number of individuals in a certain age group than expected and may even indicate a negative mortality rate, which is not possible in reality [38]. Therefore, we used the even-slip technique [39,40] to deal with the relationship between diameter class and median in diameter class to compile a static life table (Figure 2):

$$Y = -22.735 + 924.143/M \quad (R^2 = 0.845)$$

In the formula: M is the median in diameter class; Y is the number of plants.

**Figure 2.** Smoothing curve (note: M is the median in diameter class; Y is the number of plants).

2.4. Static Life Table Compilation

We used diameter class structure to represent time sequence for the woody plants to construct static life table. Static life table illustrates the patterns of population change in birth and death rates and explains the survival strategies of population. The static life table, which is also known as the specific time life table, refers to the sampling within a specific time in the dynamic aging history of multiple generations overlapping in a specific population [38], which actually reflects the population profile at a specific moment. The statistical methods of related indicators in the table are as follows [41,42]:

$$l_x = a_x / a_0 \times 1000$$

$$d_x = l_x - l_{x+1}$$

$$q_x = d_x / l_x$$

$$L_x = (l_x + l_{x+1}) / 2$$

$$T_x = \sum L_x$$

$$e_x = \frac{T_x}{l_x}$$

$$K_x = \ln l_x - \ln l_{x+1}$$

where a_0 is the actual number of survivors, and a_x is the actual number of surviving individuals in diameter class x after smoothing treatment. l_x is the number of survivors standardized to diameter class x , and the number of survivors of the largest survivors in each diameter class is standardized to 1000. Then, in turn, we obtained the standardized survival numbers of other diameter groups in. d_x is the standardized number of dead individuals during the two adjacent diameter groups; q_x is the mortality rate; L_x refers to the average number of surviving individuals (from x to $x + 1$); T_x is the total number of individuals; e_x represents the life expectancy of an individual entering diameter class x ; and K_x represents the vanishing rate.

Based on the life table, we also used four population survival analysis functions, including survival rate function $S(t)$, cumulative mortality function $F(t)$, death density function $f(t)$, and hazard rate function $\lambda(t)$ [43]:

$$S(t) = p_1 p_2 \dots p_x$$

$$F(t) = 1 - S(t)$$

$$f(t) = \frac{(S_{x-1} - S_x)}{h_x} = \frac{S_{x-1} q_x}{h_x}$$

$$\lambda(t) = \frac{2q_x}{h_x(1 + p_x)}$$

where p_x is the survival rate at diameter class x , and h_x is the length of the interval.

2.5. Survival Curve Drawing

Survival curves describe the death rate at a specific age with the number of surviving individuals and reflects the survival status of individuals in the population at various diameters. According to Deevey's theory [44], there are generally three basic types of survival curves. Type I is a convex curve with a high survival rate for young individuals and a high mortality rate for elderly individuals; Type II is a diagonal curve which has the same mortality rate throughout its life span. Type III is a concave curve which means the mortality rate in early years is high. In this paper, we adopted Hett and Loucks [45] exponential equation $N_x = N_0 e^{-bx}$ (N_x is the value of $\ln l_x$, x is the diameter class, and N_0 and b are constants) to describe the Deevey II curve, the power function $N_x = N_0 x^{-b}$ (N_x is the value of $\ln l_x$, x is diameter class, N_0 and b are constants) to describe the Deevey III curve. Then, the corresponding model was established after SPSS 19.0 fitting for comparative analysis.

2.6. Dynamic Analysis of Population Structure

Chen Xiaode [46] population and community structure dynamic quantitative analysis method is used to quantitatively describe the population structure: three dynamic indexes of V_x , V_{pi} , and V'_{pi} . When the population diameter class structure dynamic change index V_n , V_{pi} , V'_{pi} is a positive value, a negative value, and a zero value, it corresponds to the dynamic relationship of growth, decline, and structural stability, respectively. If, in the future, the interference of external k and S_n will cause dilution effect, the maximum probability value can be calculated based on the k and S_x conditional probability method

as V'_{pi} formula. V'_{pi} can also be used as an index to measure the sensitivity of population structure changes to random interference [47]. These indexes indicate population growth when V_{pi} and V'_{pi} are greater than 0; they indicate population decline when they are less than 0; and they indicate that the population is stable when they are equal to 0.

The specific calculation formula of each dynamic index is as follows:

$$V_x = \frac{(S_x - S_{x+1})}{\max(S_x, S_{x+1})} \times 100\%$$

$$V_{pi} = \frac{\sum_{x=1}^{k-1} (S_x V_x)}{\sum_{x=1}^{k-1} S_x}$$

Dynamic index of population changes when considering external disturbance V'_{pi} :

$$V'_{pi} = \frac{\sum_{x=1}^{k-1} (S_x V_x)}{K \times \min(S_1, S_2, \dots, S_k) \sum_{x=1}^{k-1} S_x}$$

In the formula, the dynamic index of population quantity change in the natural state, V_x , represents the individual dynamics from x to $x + 1$ diameter class; V_{pi} is the dynamic index of population quantity change, S_x and S_{x+1} represent the the number of individuals in x and $x + 1$ diameter class, respectively, k is the largest diameter class, and $\max/\min(\dots)$ uses the maximum/minimum value of the number sequence in parentheses.

2.7. Time Series Forecasting

In this study, based on the number of survivals in each diameter class, we use a moving average method of time series analysis [36,39] to predict the subsequent population future development trend in 2, 5, and 10 diameter class; the model is as follows:

$$M_n = \frac{1}{n} \sum_{k=x-n+1}^x D_x$$

where x represents the diameter class; n represents predicted diameter classes; M_n is the moving average of the n -th cycle, which is the number of individuals in the x -diameter class group after n diameter class; and D_x represents a_0 in the x -diameter level.

3. Results

3.1. Population Size Structure of *Taxus cuspidata*

According to the analysis of the diameter class structure in this survey (Figure 3), the diameter class structure of *T. cuspidata* is relatively complete and the population diameter class structure is an inverted L-shape. The number of seedlings is the greatest among all diameter classes, which accounts for 63.57%. During the transition from seedling to saplings, there is a great decrease in the number of *T. cuspidata* (89.54%). The survival rate of seedlings is low, and the survival rate of saplings is considerable. The number of adults between diameter class 3 and diameter class 6 is stable, accounting for 27.23%, among which diameter class 4 is the largest, accounting for 9.29%. The proportion of individuals in the large-diameter class gradually decreases, and there are a certain number of large-diameter individuals in the population. Large-diameter plants, from diameter class 7 to 11, account for 2.55%. There is a phenomenon of missing individuals in the large-diameter class, and the number of individuals in the large-diameter class is sparse. On the whole, the diameter class structure shows a trend of the number of individuals gradually decreasing as the diameter class increases (Figure 3).

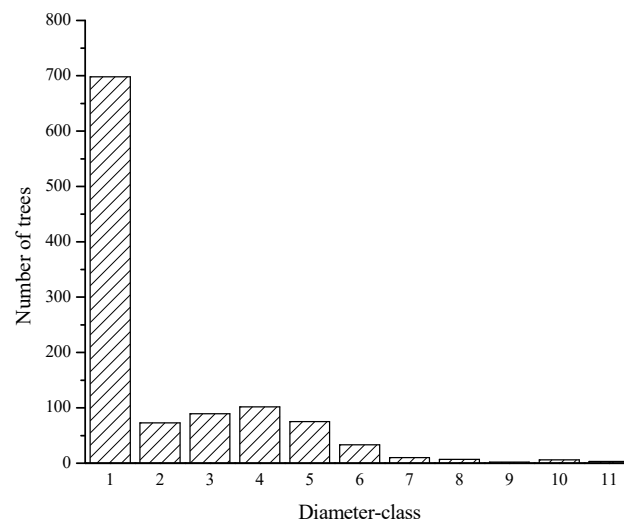


Figure 3. Number of *Taxus cuspidata* in diameter classes.

3.2. The Static Life Table of *Taxus cuspidata*

From the static life table (Table 3), we found that the number of surviving individuals l_x after standardized smoothing treatment gradually slowed down and levelled off as the diameter class increased. The smoothly processed data may neglect certain ecological phenomena in population development [48]. When we compared a_x with l_x , the trend in the data showed consistency. And there are two small peaks in the death rate and vanishing rate (Figure 4). The maximum values of death rate q_x and vanishing rate K_x appear when diameter class 3 transitions to diameter class 4. A greater mortality rate (61.89%) occurred in diameter class 3. The second largest mortality rate in q_x and K_x all appeared when diameter class 7 transitioned to diameter class 8. With the increase in diameter class, a death peak (59.89%) appeared when entering diameter class 7. Compared with the mortality and vanishing rate of other diameter classes, this was more obvious. In addition, the individual life expectancy e_x is less than 10 in diameter class 1, 2, and 3, which indicates that the population fluctuates greatly at low diameter.

Table 3. The static life table of *Taxus cuspidata* population.

x	a_0	a_x	M	l_x	$\ln l_x$	d_x	q_x	L_x	T_x	e_x	K_x
1	698	593	1.5	1000	6.9078	519	0.5189	741	741	0.74	0.7316
2	73	285	3	481	6.1761	260	0.5398	351	1092	2.27	0.7762
3	89	131	6	221	5.4000	137	0.6189	153	1245	5.62	0.9647
4	102	50	12.7	84	4.4352	37	0.4395	66	1311	15.53	0.5790
5	75	28	18.2	47	3.8563	22	0.4601	36	1347	28.48	0.6164
6	33	15	24.4	26	3.2399	13	0.5166	19	1366	53.50	0.7269
7	10	7	30.75	12	2.5130	7	0.5989	9	1375	111.38	0.9135
8	7	3	36	5	1.5995	2	0.3187	4	1379	278.50	0.3838
9	2	2	42.75	3	1.2157	2	0.5000	3	1381	409.53	0.6931
10	6	1	49.5	2	0.5226	0	0.0000	2	1383	820.07	0.0000
11	3	1	74	2	0.5226	2	1.0000	1	1384	820.57	0.5226

Note: a_0 is the actual number of survivors in diameter class x ; a_x is the number of survivors at x after smoothing treatment; M is the median value at x ; l_x is the standardized number of survivors at x ; d_x is the standardized number of dead individuals during two adjacent diameter classes; q_x is the mortality rate at each diameter class; L_x refers to the average number of surviving individuals (from x to $x + 1$ class); T_x is the total number of individuals or total life span (x class to over x class); e_x is the life expectancy of an individual at x ; and K_x indicates the vanishing rate.

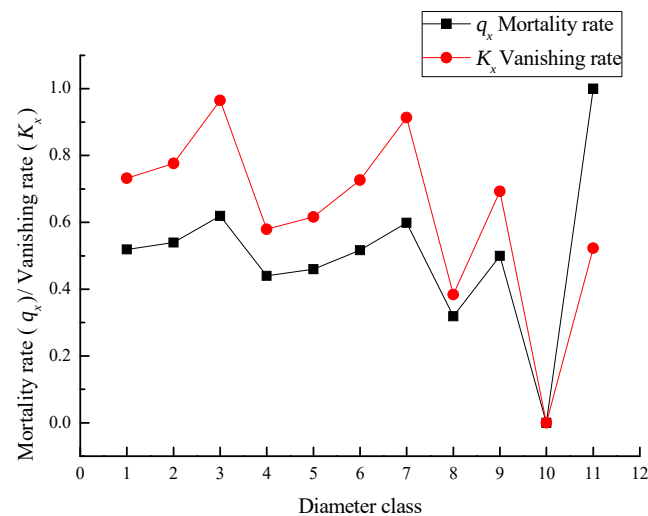


Figure 4. Mortality rate and vanishing rate curves of *Taxus cuspidata* population (note: q_x is the mortality rate; K_x is the vanishing rate).

3.3. Survival Curve of *Taxus cuspidata*

Population survival curves were plotted based on the logarithmic values of population survival numbers ($\ln l_x$) and their corresponding age classes from the static life table and simulated using the exponential and power function models (Figure 5) constructed by Deevey for type II and III survival curves, respectively [45,46]. The fitting results are as follows:

$$\text{Deevey II : } N_x = 12.271e^{-0.270x} \quad (R^2 = 0.930)$$

$$\text{Deevey III : } N_x = 13.136x^{-1.062} \quad (R^2 = 0.725)$$

where N is the $\ln l_x$, which is the logarithm of population survival; x is the diameter class.

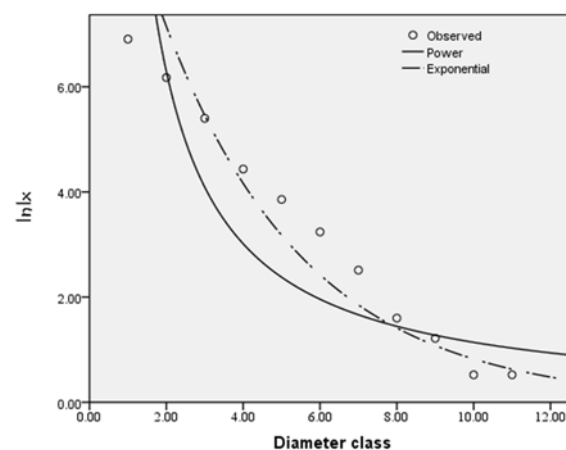


Figure 5. Survival curve of *Taxus cuspidata* population (note: $\ln l_x$ is the logarithm of population survival).

According to formula fitting, the survival curve of the *T. cuspidata* population has a higher correlation with the Deevey II curve. The diameter class is used as the x -axis and $\ln l_x$ as the y -axis to plot the survival curve of *T. cuspidata* (Figure 5). It can be seen that the survival curve is a significant Deevey II type which is diagonal, and the mortality rate is stable in all stages of population development.

3.4. Population Survival Analysis of *Taxus cuspidata*

In order to explain the changing trend of *T. cuspidata* population dynamics, we drew a graph of the survival function based on the static life table (Figure 6). It can be seen that mortality has the same changing trend with vanishing rate (Figure 4), and there are certain fluctuations in diameter class 3 and 7 in the development of the population. The survival rate gradually decreases, the cumulative death rate gradually increases, and both of the changing trends were sharp in the early stage and flatter in the later stage. And the death density curve is basically smooth, less than 3.0%, which shows that the death density of the young and middle-aged is higher than that of the old-aged. The hazard rate curve fluctuates greatly, and it also shows that the risk rate in the low-diameter class is higher than in the large-diameter class. The four survival function curves show that the population dynamics demonstrate a rapid decline in the early stage and are relatively stable in the later stage.

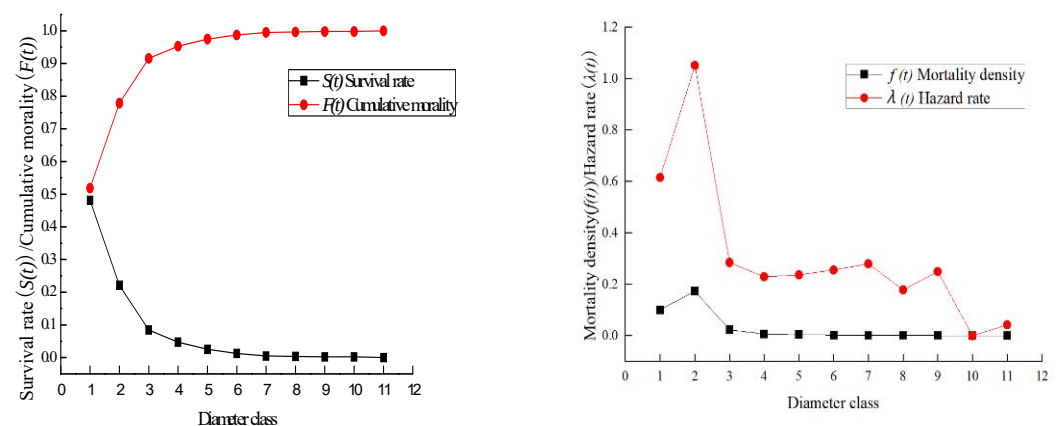


Figure 6. Survival functional rate of *Taxus cuspidata* population. ($S(t)$ is the utilization survival function, $F(t)$ is the cumulative mortality function, $f(t)$ is the mortality density function, and $\lambda(t)$ is the hazard rate function).

3.5. Dynamic Analysis of Population Structure of *Taxus cuspidata*

According to the quantitative analysis results of the individual number in each diameter class (Table 4), there is a large fluctuation phenomenon in the contribution potential between diameter classes in population development. V_2 , V_3 , and V_9 are all less than zero, indicating that the *T. cuspidata* population is in a trend of population decline when the diameter class is 2, 3, and 9. There is fluctuation in diameter class 9, which may be due to human interference. The population structure dynamic indexes in the other diameter stages are all greater than zero, indicating that the population of *T. cuspidata* is growing at most diameter stages. The dynamic index of the population structure $V_{pi} = 0.6395$ when ignoring external environmental interference. When considering external environmental interference, the index is $V'_{pi} = 0.0291$, which is close to 0, suggesting that the population structure is less likely to be growth-oriented and tends to be stable. In terms of the overall dynamic index, V_{pi} is positive, indicating that the population structure is a growth-type. V'_{pi} is much less than the V_{pi} value, indicating that the population is highly sensitive to external environmental disturbances. Or it indicates that the population is highly influenced by external disturbances, but the population is highly resistant to disturbances. In summary, the population is a growth-stable population with a declining trend at young ages and a relatively stable state overall.

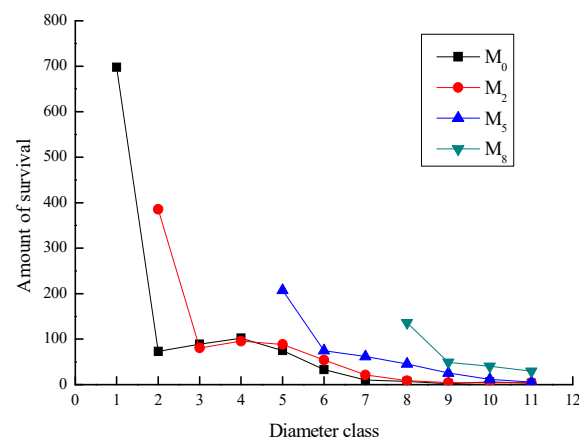
Table 4. The dynamic index (V_x) of the individual number in the *Taxus cuspidata* population in two adjacent diameter classes and the dynamic index of the population structure (V_{pi}).

Item	V_1	V_2	V_3	V_4	V_5	V_6	V_7	V_8	V_9	V_{10}	V_{11}	V_{pi}	V'_{pi}
Dynamic index (%)	0.9	−0.18	−0.13	0.26	0.56	0.7	0.3	0.71	−0.67	0.5	–	0.6395	0.0291

Note: V_x represents the dynamic index of the individual number in population from x to $x + 1$; V_{pi} represents the dynamic index of the entire population structure; V'_{pi} represents the dynamic index of the population's diameter structure when external disturbances are taken into account.

3.6. Dynamic Prediction of *Taxus cuspidata* Population

The stability of *T. cuspidata* was analyzed and the possible changes of population were simulated based on the population pattern of plants at each diameter level (Figure 7). After two diameter classes of time, the individual number in diameter class 2 and diameter classes 5–7 was increased; after five diameter classes of time, the number of individuals in diameter class 5 increased significantly and the number of individuals in diameter classes 6–10 increased slightly; after eight diameter classes of time, the number of individuals in diameter class 9 increased significantly and the number of individuals in diameter classes 9–11 increased slightly. The overall variation is that the number of individuals in large-diameter classes increases, and the population stabilizes, with an increase in diameter class.

**Figure 7.** The time series prediction of *Taxus cuspidata* population (note: M_0 is the current number distribution curve of *Taxus cuspidata* in various diameter classes; M_2 is the predicted 2-diameter classes curve; M_5 is the predicted 5-diameter classes curve; M_8 is the predicted 8-diameter classes curve).

4. Discussion

4.1. Population Structure Analysis

The number of seedlings was significantly higher than the number of individuals in the larger-diameter classes, and a large number of young individuals in the population in diameter classes 1, 2, and 3 ($0 \text{ cm} \leq \text{DBH} < 10 \text{ cm}$) were able to survive under the main forest layer occupied by various dominant trees in the community, indicating that germination and seedling growth were adapted to be completed in a shaded environment. And we found that it was difficult for its seedlings to grow into young and small trees, due to the fact that other tree species occupied the upper layer for a long time. When the population reached a certain threshold, with the increase in individuals' demand for nutrients and living space, the overlap with neighboring individuals and upper trees increased and the competition for living space among individuals intensified. But the environmental loading capacity reached its limit, resulting in self-thinning and combing due to density constraints. In this study, the first peak of population mortality may be attributed to the increase in intraspecific and interspecific competition. The population of *T. cuspidata* showed a rapid decline with increasing age, making it more difficult to enter the succession layer through strong environmental filtering. Most seedlings either grow slowly

and wait to enter the main forest layer or gradually die while waiting. It is difficult for these seedlings to grow into small- and medium-sized trees, which is related to the competition for light and nutrients within and among species. To overcome these obstacles, a suitable balance between light availability, nutrient availability, and competition management is crucial for the successful growth and development of *T. cuspidata*.

The population size fluctuated in diameter class 9 ($40 \text{ cm} \leq \text{DBH} < 46 \text{ cm}$), and it is speculated that the population suffered from strong anthropogenic disturbance and deviated from a stable population structure. During the period from the 1950s to 1990s, there was a significant increase in unlawful logging, particularly targeting large-diameter plants. This had detrimental effects on the stand type and structure of forests. The logging activities resulted in the destruction of mature and old trees that held higher economic value [13]. Luckily, the forests were left with a substantial soil seed bank, which played a crucial role in the subsequent regeneration of the forests. Furthermore, the logging activities had a profound impact on the distribution of habitats, leading to habitat fragmentation. This fragmentation posed a major challenge for the development of the population. Therefore, logging is considered the primary culprit behind habitat fragmentation, which has had significant implications for the population dynamics of the species in question. The study by Diao Y.F. [20] on the structural characteristics of *T. cuspidata* in Muling of Heilongjiang concluded that the middle-aged, mature, and old trees in Murong are relatively stable and abundant, which is also consistent with the findings of our paper, indicating that *T. cuspidata* can form a large number of large-diameter populations and maintain population stability. And the survival curves did not show the complete Deevey's classification because there were only three individuals with $8 \text{ cm} \leq \text{DBH} < 16 \text{ cm}$ in the Muling area. In addition, the results of Liu T. [2] showed that the natural population of *T. cuspidata* in the nature reserve had a large population size and relatively concentrated distribution, but the population structure was unreasonable, with a large number of seedlings and large-diameter individuals in the renewal layer and few individuals in the succession layer. In this paper, the dynamics of the natural populations in Jilin were studied to remedy this deficiency and to describe the structural characteristics of the population in a more comprehensive diameter structure. It can also be further proved that the environment in Jilin is indeed a highly suitable area for *T. cuspidata* [14].

4.2. Population Dynamics Analysis and Conservation Recommendations

The dynamic index showed that there were large fluctuations in the contribution potential between diameter classes in the process of population development, and V_2 , V_3 , and V_9 were all less than 0. Regarding the analysis of the overall dynamic index, V_{pi} was positive, which indicated that the population structure of *T. cuspidata* was growth-oriented. The fact that V'_{pi} was much less than V_{pi} indicated that the population was more sensitive to external environmental disturbances. And when the external environmental disturbances were considered, $V'_{pi} = 0.0291$, the dynamic index (V'_{pi}) was greater than 0, but close to 0, which indicated that the population is more sensitive to external environmental disturbances—even moderately intense human activities. Further, under external disturbances, the *T. cuspidata* population tends to be stable and shows some resistance to disturbance. So, the establishment of in situ reserves is a key measure to maintain *T. cuspidata* habitat integrity, prevent or slow down the decline of the population structure, and protect this rare and endangered species effectively.

The choice of appropriate measures will depend on the causes of the decline [12]. According to the results of the static life table and dynamic indices, the *T. cuspidata* population undergoes a total of two fluctuations in its growth and developmental stages, one during juvenile age and the other in diameter class 9. The first fluctuation is related to the competition for light and nutrients within and among species. Based on our analysis, we conclude that the second fluctuation observed, in diameter class 9, is likely attributable to human disturbance. This conclusion is drawn from two main pieces of evidence. Firstly, the survival curve for this diameter class follows a significant Deevey II pattern, which

is characterized by a diagonal curve, indicating stable mortality rates across all stages of population development. This suggests that the mortality rate for trees in diameter class 9 remains consistent over time, indicating external factors influencing the population. Secondly, the existence of a large number of large diameter trees is consistent with the hypothesis of unlawful logging. Typically, a healthy forest population would display a more balanced distribution of tree diameters, with a gradual decrease in numbers as diameters increase, just like our time series prediction. However, an abrupt increase in the number of large diameter trees suggests that selective logging or illegal activities targeting specific tree sizes have occurred. So, it is important to take into account the specific local environmental conditions and to strengthen the protection of mature trees from felling and logging. The temporal prediction curves indicate that the proportion of large-diameter individuals increases after the population has passed through the seedling and juvenile stages. The population in Jilin is a stable population with a low growth rate, and the conservation of this population should take into account the local environmental conditions. Different conservation measures should be adopted for the population with different diameter structures under anthropogenic disturbance. We need to reduce anthropogenic disturbance and enhance the population of saplings and seedlings at the same time. As the large-diameter populations are rare and few in the wild, we recommend allowing them to grow naturally through enclosure protection. But this may lead to a further decline in population after the mature mother tree has reached its fruiting peak. And random simulation studies have shown that even large existing populations in nutrient-rich areas are unlikely to survive in the long term unless favorable conditions for establishing young plants are restored [49]. According to Su [50], both natural *T. cuspidata* and transplants of *T. cuspidata* are characterized by high genetic diversity, no apparent genetic structure, and abundant gene flow, suggesting that this population is recently endangered. In addition, integration analyses conducted by Aguilar et al. [51] and Vranckx et al. [52] showed that the longer widespread species, recently endangered allopatric species, and forest tree species persist in fragmented habitats—which is more detrimental to the recovery of their genetic diversity and the maintenance of their allopatric breeding systems—and they are highly susceptible to the loss of genetic diversity in future generations. The adverse effects of severe habitat fragmentation may be partially offset by frequent gene exchange [53]. These measures can be crucial in ensuring the long-term conservation of species in fragmented habitats. Let us examine each measure in detail: 1. Conservation and expansion of the mother tree population: by conserving and expanding the population of mother trees, we ensure that there are enough individuals producing viable seeds or other reproductive structures. This allows for gene exchange to occur through sexual reproduction, increasing genetic diversity within the population. 2. Preservation of germplasm resources: germplasm refers to the genetic material of plants, including seeds, pollen, and tissue cultures. By preserving a diverse range of germplasm resources, we maintain a bank of genetic diversity that can be used for future gene exchange. This involves collecting and storing seeds, pollen, or tissue samples from different individuals or populations. 3. Establishment of pollen banks: pollen is the male reproductive structure of plants, and it plays a crucial role in gene exchange. By establishing pollen banks, we can collect and store pollen from a variety of individuals or populations. This ensures that specific genetic traits can be introduced or maintained through artificial pollination, even if individuals or populations are not geographically close to each other. Moreover, measures such as artificially assisted pollination, lifting seed dormancy, and timely replenishment of the seed bank can also be carried out among fragmented habitats to maintain genetic connectivity among populations and population stability during long-term conservation of the species [50,54]. Therefore, this study suggests that while conserving wild populations, in situ conservation of remaining populations, relocation of germplasm resources, and reduction of human activities will be beneficial to the long-term survival of these endangered species and that natural habitat restoration and relocation of conservation efforts should be carried out in order to increase the size of the population and genetic diversity. Furthermore, it is necessary to take further effective

conservation measures for *T. cuspidata* and other dominant species in the community to protect the seed stock and ecological balance of the community and ensure the development and survival of the seed bank of other dominant species in the absence of large-diameter *T. cuspidata* [55]. Conservation actions for this threatened plant should focus on protecting seedlings so that they grow into mature trees and on the existing population; this strategy is the key element to increasing the population size and to ensuring the persistence of the population [56].

5. Conclusions

The static life table suggested that the population of *T. cuspidata* was of a stable type overall, and the population survival curve tended to be Deevey II in Jilin. The diameter-class structure of *T. cuspidata* was stable overall, and the dynamic indices showed that the mortality rate of young individuals was high and the population was influenced by external disturbances and showed some resistance to human disturbance. Time sequence prediction analysis showed that natural renewal could be maintained and the population size would remain at a certain amount in the future. The result can prove that the Jilin region is a highly suitable area for the growth of *T. cuspidata* in Northeast China. In situ conservation of remaining wild populations, relocation of germplasm resources, and reduction of human activities are recommended as actions that will be beneficial to the long-term survival of *T. cuspidata*.

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References

1. Zang, R.G.; Cheng, K.W.; Li, J.Q.; Zhang, W.Y.; Chen, X.F.; Tao, J.P. *Natural Forest Biodiversity Conservation and Restoration*; Science and Technology Press: Beijing, China, 2005; pp. 464–480. (In Chinese)
2. Liu, T. *Population Ecology of Natural Japanese Yew*; Northeast Forestry University: Harbin, China, 2007; pp. 1–2. (In Chinese with English abstract)
3. Chen, J.; Long, T.; Yang, L.; Wang, Y.; Xu, C.; Li, J.W. Habitat suitability assessment of *Taxus cuspidata*. *J. Beijing For. Univ.* **2019**, *41*, 51–59. (In Chinese with English abstract)
4. Menges, E.S. Population viability analyses in plants: Challenges and opportunities. *Trends Ecol. Evol.* **2000**, *15*, 51–56. [[CrossRef](#)]
5. Dang, H.; Zhang, Y.; Zhang, K.; Jiang, M.; Zhang, Q. Age structure and regeneration of subalpine fir (*Abies fargesii*) forests across an altitudinal range in the Qinling Mountains, China. *For. Ecol. Manag.* **2010**, *259*, 547–554. [[CrossRef](#)]
6. Dolanc, C.R.; Thorne, J.H.; Safford, H.D. Widespread shifts in the demographic structure of subalpine forests in the Sierra Nevada, California, 1934 to 2007. *Glob. Ecol. Biogeogr.* **2013**, *22*, 264–276. [[CrossRef](#)]
7. Diao, Y.F.; Jin, G.Z.; Tian, S.Y.; Liu, Y.K.; Liu, Y.L.; Han, L.D.; Li, Y.H. Species composition and community structure of a *Taxus cuspidata* forest in Muling nature reserve of Heilongjiang province, China. *Sci. Silvae Sin.* **2016**, *52*, 26–36. (In Chinese with English abstract)
8. Easterling, M.R.; Ellner, S.P.; Dixon, P. Size specific sensitivity: A new structured population model. *Ecology* **2000**, *81*, 694–708. [[CrossRef](#)]
9. Ramula, S.; Kerr, N.Z.; Crone, E.E. Using statistics to design and estimate vital rates in matrix population models for a perennial herb. *Popul. Ecol.* **2019**, *62*, 53–63. [[CrossRef](#)]
10. Werner, P.A.; Peacock, S.J. Savanna canopy trees under fire: Long-term persistence and transient dynamics from a stage-based matrix population model. *Ecosphere* **2019**, *10*, e02706. [[CrossRef](#)]
11. Gao, W.Q.; Ni, Y.Y.; Xue, Z.M.; Wand, X.F.; Kang, F.F.; Hu, J.; Gao, Z.H.; Jiang, Z.P.; Liu, J.F. Population structure and regeneration dynamics of *Quercus variabilis* along latitudinal and longitudinal gradients. *Ecosphere* **2017**, *8*, 1–15. (In Chinese with English abstract) [[CrossRef](#)]
12. Jin, H.; Zhao, Y.; Yin, H.; Qin, L.W.; Liu, L.J.; Wang, C.; Jia, X.; Li, B.Y. Quantitative characteristics and dynamic analysis of the endangered species *Rhododendron chrysanthum* population in Changbai Mountain. *Chin. J. Ecol.* **2017**, *36*, 3123–3130. (In Chinese with English abstract)

13. Kroon, H.D.; Groenendaal, J.V.; Ehrlen, J. Elasticities: A review of methods and model limitations. *Ecology* **2000**, *81*, 607–618. [[CrossRef](#)]
14. Wang, D.D.; Zhang, Y.W. Identification and genetic diversity analysis of *Taxus cuspidata* hybrid. *J. Northeast Norm. Univ.* **2019**, *51*, 113–118. (In Chinese with English abstract)
15. Wu, S.X.; Liu, Y.H.; Zhang, L.M.; Shang, F.Q.; Tan, C.Q. Growth stability analysis of ex situ conservation of *Taxus cuspidata* seedling from different sources. *J. Northeast Norm. Univ.* **2018**, *40*, 27–37. (In Chinese with English abstract)
16. Wang, T. *Study on Taxus cuspidata var Seeding Raising Technology*; Northeast Normal University: Harebin, China, 2017; pp. 11–22. (In Chinese with English abstract)
17. Liu, T.; Xu, H.Y.; Jin, H.Y.; Zhang, Y.H.; Zhou, Z.Q. Diurnal changes of chlorophyll fluorescence characteristics of Japanese Yew seedlings under different light conditions. *Bull. Bot. Res.* **2015**, *35*, 848–853. (In Chinese with English abstract)
18. Liu, T.; Jin, H.Y.; Xu, H.Y.; Zhou, Z.Q.; Zhang, Y.H. Diurnal changes of photosynthetic characteristics of *Taxus cuspidata* seedlings under different light conditions. *J. Northeast Norm. Univ.* **2015**, *37*, 67–72. (In Chinese with English abstract)
19. Li, Y.L. *Interspecific Relationship of the Taxus cuspidata*; Northeast Forestry University: Harebin, China, 2008; pp. 17–35. (In Chinese with English abstract)
20. Diao, Y.F.; Liu, Y.K.; Liu, Y.L.; Tian, S.Y. Population structure characteristics of *Taxus cuspidata*. *Sci. Sin. Vitae* **2020**, *50*, 391–397. (In Chinese with English abstract)
21. Caswell, H. *Matrix Population Models: Construction, Analysis and Interpretation*; Sinauer: Sunderland, MA, USA, 1989.
22. Caswell, H. Life cycle models for plants. *Lect. Math. Life Sci.* **1986**, *18*, 171–233.
23. Vandermeer, J. Choosing category size in a stage projection matrix. *Oecologia* **1978**, *32*, 79–84. [[CrossRef](#)]
24. Caswell, H. Prospective and retrospective perturbation analyses: Their roles in conservation biology. *Ecology* **2000**, *81*, 619–627. [[CrossRef](#)]
25. Dong, M.Z.; Wang, L.T.; Lv, M.J.; Meng, D.; Yang, C.Y.; Zhao, C.J.; Fu, Y.J.; Yang, Q. Communities and protection strategy of endangered species wild *Taxus cuspidata*. *Bull. Bot. Res.* **2020**, *40*, 416–423. (In Chinese with English abstract)
26. Zhang, X.S. *Vegetation Map of China and Its Geographic Pattern: Illustration of the Vegetation Map of the People's Republic of China (1:1 000 000)*; Geological Publishing House: Beijing, China, 2007. (In Chinese)
27. Bridie, C.; Fortin, H.M.J. Development of a *Populus Balsamifera* Clone in subarctic Quebec reconstructed from spatial analyses. *J. Ecol.* **1995**, *83*, 309–320. [[CrossRef](#)]
28. Johnson, J.B. Stand structure and vegetation dynamics of a subalpine treed fen in Rocky Mountain National Park, Colorado. *J. Veg. Sci.* **1997**, *8*, 337–342. [[CrossRef](#)]
29. Rebertus, A.J.; Veblen, T.T. Structure and tree fall gap dynamics of old-growth *Nothofagus* forests in Tierra del Fuego, Argentina. *J. Veg. Sci.* **1993**, *4*, 641–654. [[CrossRef](#)]
30. Harper, J.L. *Population Biology of Plants*; Academic Press: London, UK, 1977.
31. Frost, I.; Rydin, H. Spatial pattern and size distribution of the animal-dispersed tree *Quercus robur* in two spruce-dominated forests. *Ecoscience* **2000**, *7*, 38–44. [[CrossRef](#)]
32. Harcombe, P.A.; Marks, P.L. Tree diameter distributions and replacement processes and replacement processes in southeast Texas forests. *For. Sci.* **1978**, *24*, 153–166.
33. Lorimer, C.G. Age structure and disturbance history of a southern Appalachian virgin forest. *Ecology* **1980**, *61*, 1169–1184. [[CrossRef](#)]
34. Gao, H.Z.; Huang, X.; Su, H.; Qiao, P.F.; Jiang, Z.M.; Shen, Y.R.; Cai, J. Structure and dynamic characteristics of *Betula albo-sinensis* populations in two regions in the Qinling Mountains of northwestern China. *J. Beijing For. Univ.* **2022**, *44*, 12–20. (In Chinese with English abstract)
35. Zhang, J.; Pan, Z.L.; Tian, Y.H.; Hou, J.L.; Peng, M.C.; Duan, H.; Li, Y.F.; Li, Y.P. Age structure and dynamics of *Pinus yunnanensis* population in Yunlong Tianchi Nature Reserve. *Acta Ecol. Sin.* **2022**, *42*, 9091–9099. (In Chinese with English abstract)
36. Li, F.R. Study on diameter distribution and models prediction yields for natural Dahurian larch stands. *J. North-East For. Univ.* **1987**, *15*, 8–16. (In Chinese with English abstract)
37. Ding, G.M.; Liu, X.M.; Ni, Z.Y.; Peng, J.Y. A preliminary study of population dynamics of *Picea crassifolia* in Qilian Mountains Gansu Province. *J. Plant Resour. Environ.* **2005**, *14*, 36–41. (In Chinese with English abstract)
38. Begon, M.; Mortimer, M.; Thompson, D.J. *Population Ecology: A Unified Study of Animal and Plants*; Blackwell Science Ltd.: London, UK, 1981; pp. 1–1088.
39. Jiang, H. *Population Ecology of Spruce*; Chinese Forestry Press: Beijing, China, 1992; pp. 1–26. (In Chinese)
40. Liu, J.H.; Hong, W. A study on forecast of population dynamics of *Castanopsis kawakamii*. *Chin J. Appl. Environ. Biol.* **1999**, *5*, 247–253. (In Chinese with English abstract)
41. Wratten, S.D.; Fry, G.I.A. *Field and Laboratory Exercises in Ecology*; Edward Arnold: London, UK, 1996; pp. 98–103.
42. Wang, M. *The Study on Population Dynamics and Spatial Distribution Pattern of Main Plants in Xiaolingshan of Gansu Province*; Northwest Normal University: Lanzhou, Gansu, 2012; pp. 7–10. (In Chinese with English abstract)
43. Yang, F.X.; Wang, S.Y.; Xu, H.G.; Li, B.Q. The theory of survival analysis and its application to life table. *Acta Ecol. Sin.* **1991**, *11*, 153–158. (In Chinese with English abstract)
44. Deevey, E.S. Life tables for natural populations of animals. *Q. Rev. Biol.* **1947**, *22*, 283–314. [[CrossRef](#)] [[PubMed](#)]
45. Hett, J.M.; Loucks, O.L. Age structure models of balsam fir and eastern hemlock. *J. Ecol.* **1976**, *64*, 1029–1044. [[CrossRef](#)]

46. Chen, X.D. A study on the method of quantitative analysis for plant population and community structural dynamics. *Acta Ecol. Sin.* **1998**, *18*, 104–107. (In Chinese with English abstract)
47. Liu, G.J.; Zhang, X.M.; Zhu, J.T.; Lv, C.Y.; Lu, Y. Structure and dynamics of *Haloxylon ammodendron* population along the southeastern margin of the Zhunger basin. *Acta Bot. Boreal.-Occident. Sin.* **2011**, *31*, 1250–1256. (In Chinese with English abstract)
48. Yang, L.; Zhang, Z.L.; Yun, Y.; Yan, W.P.; Chen, X.; Zhang, L.; Zheng, D.J.; Chen, J.L. The population structure and dynamics of *Dracaena cambodiana*, an endangered tree on Hainan island. *Acta Ecol. Sin.* **2018**, *38*, 201–214. (In Chinese with English abstract)
49. Colling, G.; Matthies, D. Effects of habitat deterioration on population dynamics and extinction risk of an endangered, long-lived perennial herb (*Scorzonera humilis*). *J. Ecol.* **2006**, *94*, 959–972. [[CrossRef](#)]
50. Su, J.Y.; Yan, Y.; Li, C.; Li, D.; Du, F. Informing conservation strategies with genetic diversity in wild plant with extremely small populations: A review on gymnosperms. *Biodivers. Sci.* **2020**, *28*, 376–384. (In Chinese with English abstract)
51. Aguilar, R.; Quesada, M.; Ashworth, L.; Herrera's-Diego, Y.; Lobo, J. Genetic consequences of habitat fragmentation in plant populations: Susceptible signals in plant traits and methodological approaches. *Mol. Ecol.* **2008**, *17*, 5177–5188. [[CrossRef](#)]
52. Vranckx, G.U.Y.; Jacquemyn, H.; Muys, B.; Honnay, O. Meta-analysis of susceptibility of woody plants to loss of genetic diversity through habitat fragmentation. *Conserv. Biol.* **2012**, *26*, 228–237. [[CrossRef](#)] [[PubMed](#)]
53. Kramer, A.T.; Ison, J.L.; Ashley, M.V.; Howe, H.F. The paradox of forest fragmentation genetics. *Conserv. Biol.* **2008**, *22*, 878–885. [[CrossRef](#)] [[PubMed](#)]
54. Ellstrand, N.C.; Elam, D.R. Population genetic consequences of small population size: Implications for plant conservation. *Annu. Rev. Ecol. Syst.* **1993**, *24*, 217–242. [[CrossRef](#)]
55. Jiang, Y.H.; Shen, W.H.; Tan, C.Q.; Xiang, W.H.; Peng, Y.H.; Pan, G.B. The population structure and dynamics analysis and protection strategy of *Vatica guangxiensis* extremely small population. *Ecol. Sci.* **2016**, *35*, 67–72.
56. Cogoni, D.; Sulis, E.; Bacchetta, G.; Fenu, G. The unpredictable fate of the single population of a threatened narrow endemic Mediterranean plant. *Biodivers. Conserv.* **2019**, *28*, 1799–1813. [[CrossRef](#)]

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