

Regeneration of *Betula albosinensis* in Strip Clearcut and Uncut Forests of the Qinling Mountains in China

Yaoxin Guo¹, Gang Li², Youning Hu³, Di Kang³, Dexiang Wang³, Gaihe Yang¹*

1 College of Agronomy, Northwest A&F University, Yangling, Shaanxi, China, 2 College of Life Science, Northwest A&F University, Yangling, Shaanxi, China, 3 College of Forestry, Northwest A&F University, Yangling, Shaanxi, China

Abstract

To contribute to a better understanding of the regeneration strategy of Betula albosinensis forests and the likely reasons behind either the successful recovery or failure after strip clearcutting, we compared the population structures and spatial patterns of B. albosinensis in eight B. albosinensis stands in Qinling Mountains, China. Four cut and four uncut stands were selected, and each sampled using a single large plot (0.25 ha). Results indicated that, on the one hand, B. albosinensis recruitment was scarce (average of 48 stems ha⁻¹) in the uncut stands, relative to the mature population (average of 259 stems ha-1), suggesting a failure of recruitment. On the other hand, the subsequent regeneration approximately 50 years after the strip clearcutting showed that the density of the target species seedlings and saplings has increased significantly, and the current average density of seedlings and saplings was 156 stems ha^{-1} . The clumped spatial pattern of B. albosinensis suggested that their regeneration was highly dependent on canopy disturbance. However, recruitment remained poor in the uncut stands because most gaps were small in scale. The successful regeneration of sunlight-loving B. albosinensis after strip clearcutting was attributed to the exposed land and availability of more sunlight. Bamboo density did not influence B. albosinensis recruitment in the uncut stands. However, stand regeneration was impeded after strip clearcutting; thus, removing bamboo is essential in improving the competitive status of B. albosinensis at the later stage of forest regeneration after clearcutting. The moderate severity of disturbance resulting from strip clearcutting reversed the degeneration trend of primary B. albosinensis stands. This outcome can help strike a balance between forest conservation and the demand for wood products by releasing space and exposing the forested land for recruitment. Life history traits and spatiotemporal disturbance magnitude are important factors to consider in implementing effective B. albosinensis regeneration strategies.

Citation: Guo Y, Li G, Hu Y, Kang D, Wang D, et al. (2013) Regeneration of *Betula albosinensis* in Strip Clearcut and Uncut Forests of the Qinling Mountains in China. PLoS ONE 8(3): e59375. doi:10.1371/journal.pone.0059375

Editor: Ben Bond-Lamberty, DOE Pacific Northwest National Laboratory, United States of America

Received September 23, 2012; Accepted February 13, 2013; Published March 22, 2013

Copyright: © 2013 Guo et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This research was funded by the Key Program of the National Basic Sci-Tech Special Fund of China (2007FY110800). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

1

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: ygh@nwsuaf.edu.cn

Introduction

Betula albosinensis, a deciduous hardwood, is a tree species endemic to China, distributed in the mid-high mountains of warm temperate regions. As one of the most important species in the Qinling Mountains, B. albosinensis thrives over a wide elevation range of 1950 m to 2750 m [1,2]. The community (i.e., species composition, spatial structure, and gap characteristics) [3–5] and seed germination characteristics [6,7] of B. albosinensis forests have been studied. No direct study, however, has reported on the regeneration of B. albosinensis trees in forests where they dominate after natural disturbance or artificial management treatments. This type of ecological knowledge is essential in implementing conservation strategies and ensuring the sustainable utilization of forests [8,9].

Tree regeneration is influenced by many factors, such as the life history attributes and disturbance of species and the competitive interactions among them. Disturbances play an important role in the regeneration dynamics of many mature hardwood forests [8–12]. For pioneer tree species, natural disturbance is regarded as an important measure of population persistence [13]. In China, the natural regeneration of *B. albosinensis* forests in the Qinling

Mountains is poor and may soon be replaced by other stable species [5,14–15]. In response to the increasing demand for forest products through regeneration, strip clearcutting was conducted in several *B. albosinensis* forests in the Qinling Mountains as a sustainable alternative for forest regeneration [7,16–17]. Nevertheless, further studies from ecological and silvicultural perspectives on the regeneration dynamics of *B. albosinensis* after strip clearcutting are required.

Understory bamboos in temperate and tropical subalpine forests are particularly effective in reducing seedling recruitment and tree regeneration when they reach a high degree of dominance [18–21]. In the Qinling Mountains, Fargesia qinlingensis is a common understory bamboo in B. albosinensis forests and dominates the understory in most sites. Therefore, understanding the role of F. qinlingensis in B. albosinensis forests may be critical in determining the regeneration dynamics of the latter. In this study, we analyzed the population structures and spatial patterns of B. albosinensis populations in strip clearcut and uncut B. albosinensis forests under different bamboo covers. The objectives included the following: (1) identify the regeneration patterns of B. Albosinensis, (2) to examine whether B. albosinensis regeneration after strip clearcutting was adequate to grow a new forest, and (3) to determine the influence

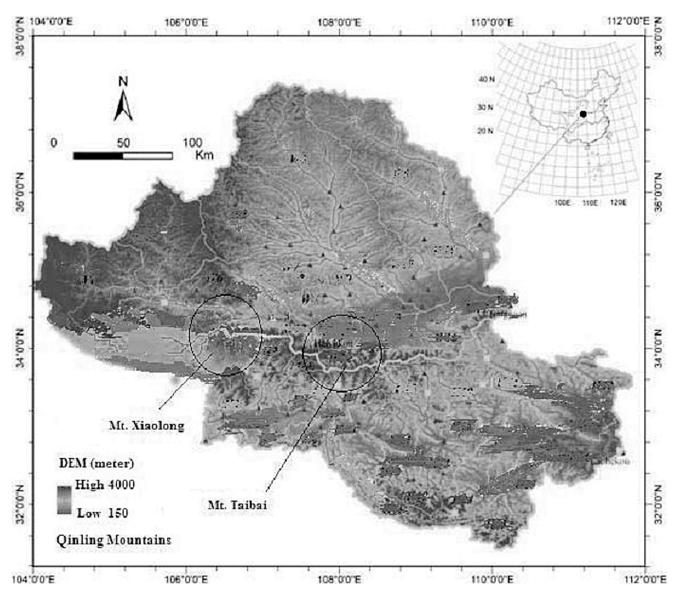


Figure 1. Location of the study areas in the Qinling Mountains of China. doi:10.1371/journal.pone.0059375.g001

of understory bamboo on the regeneration and community structure of *B. albosinensis*.

Methods

Study Area

This study was conducted at the Mt. Taibai National Nature Reserve (33°49 to 34°10′N, 107°19′ to 107°58′E, Shaanxi Province) and Mt. Xiaolong National Nature Reserve (33°35′ to 34°06′N, 106°13′ to 106°34′E, Gansu Province), located in the middle and western areas of the Qinling Mountains in China, respectively (Fig. 1). The Qinling Mountains run east—west and act as an important watershed divider between two great rivers of China, the Yangtze River and the Yellow River, which constitute a transitional zone between northern subtropical zone and warm-temperate zone. Mt. Taibai is the highest mountain in the Qinling Mountains, which spans an altitudinal gradient of 530 to 3767 m. Mean annual rainfall is 750 to 1100 mm, primarily falling in June through August, which are also the warmest months with mean

monthly temperature of 13.9 and 12.1°C, and December and January are the coldest months with monthly temperature of -5.7 and -4.4°C [15,22]. Elevation in the Mt. Xiaolong ranges from 704–3300 m. Annual precipitation ranges from 460–850 mm, most of which falls between July and September. Annual mean temperature ranges from 7 to 13°C [2]. The Nature Reserves were established for multiple-uses, including research, animal protection and forest production. Research activities were conducted under the scientific use permits issued respectively by Forestry Department of Shaanxi Province and Forestry Department of Gansu Province. Our field study did not involve any endangered or protected species in the Nature Reserves.

The numbers in parentheses are the estimated seedlings and saplings established in canopy gaps.

B. albosinensis forest is an important type of forest vegetation of Qinling Mountains, distributed from 1950 to 2750 m in Mt. Taibai and from 2000 to 2600 m in Mt. Xiaolong. B. ablosinensis forests in Mt. Xiaolong were strip clearcut in 1960 s and 1970 s to promote regeneration [2]. In Mt. Taibai, however, human

Table 1. Characteristics of stands and *B. ablosinensis* population in strip clearcut and uncut *B. ablosinensis* forest of Qinling Mountains.

Stands	Uncut	Uncut	Uncut	Uncut	V Strip clearcut	Strip clearcut	VII Strip clearcut	VIII Strip clearcut
Slop (°)	30	28	17	15	32	25	28	26
Aspect	NW	N	NW	N	N	NE	N	W
Bamboo cover (%)	45	15	8	5	5	10	20	50
Density (DBH \geq 5.0 cm,ha ⁻¹)	244	249	267	276	529	693	590	465
Basal area (m² ha -1)	18.1	17.6	22.8	19.3	24.8	24.7	27.6	23.8
Seedlings ($<$ 1.5 m tall, ha $^{-1}$)	16 (11)	8 (6)	16(12)	8 (7)	72	88	58	46
Saplings($<$ 5.0 cm DBH and \ge 1.5 m tall, ha $^{-1}$)	36 (31)	32 (26)	48(39)	28 (24)	106	86	92	76
Observed no. of species	13	16	20	18	15	18	16	11
Shannon diversity (including <i>F. qinlingensis</i>)	0.79	1.45	1.63	1.76	1.58	1.68	1.38	0.49

doi:10.1371/iournal.pone.0059375.t001

activities are rare due to the relatively difficult accessibility, and some of the only remaining intact forests occur in the region. Thus, the study chooses the stands in Mt. Taibai and Mt. Xiaolong to examine regeneration of *B. ablosinensis* comparatively.

Field Sampling

After reconnaissance, we selected four cut stands (V–VIII) in the strip clearcut area where the primary trees and undergrowth were felled from the base in the 1960 s, except some large individuals as mother trees. Four stands (I–IV) without cut disturbance were also sampled. Stands were selected if they met the following criteria: (i) *B. albosinensis* dominated the stands and represented the typical forest structure at each site; (ii) the trees in the stands were 100 years, thus classifying them as mature stands; and (iii) sampled stands included observable variations in bamboo cover.

In the current study, each stand was sampled with a large plot $(50 \text{ m} \times 50 \text{ m})$. Within each plot, all trees with diameter at breast height (or DBH, i.e., 1.3 m above ground level) longer than 5.0 cm were measured. Trees in the stands with DBH less than 5.0 cm and other woody plants with height taller than 1 m were included in five subplots measuring $5 \text{ m} \times 5 \text{ m}$, with the trees distributed in the middle and at the four corners of each large plot. Bamboo cover in the stands was estimated by tallying the bamboo coverage in the subplots upon the culms.

Data Analyses

Woody plant diversity was computed for each stand using Shannon's formula [23]. Only woody plants that reached 1.0 m in height were considered in the diversity estimation. Species richness in a plot was computed as the number of woody species with stems taller than 1.0 m. Size-structure diagram for *B. albosinensis* was prepared to depict the frequency of different-sized individuals and to interpret the trends in population dynamics.

The spatial pattern of the *B. albosinensis* population was identified using Morisita's index [24]:

$$I_{\delta} = q \sum n_i(n_i - 1) / N(N - 1)$$

Where q is the number of quadrats, n_i is the number of individuals in the ith quadrat, and N is the total number of individuals in all

quadrats. I_{δ} equals 1.0 when the population is randomly distributed, $I_{\delta} > 1$ if a population is clumped, and $I_{\delta} < 1$ if a population is regularly distributed. The intensity of pattern was interpreted from the magnitude of the index value. The greater the index value the greater the intensity of clumping. The scale of pattern (m²) was identified by computing Morisita'index values for quadrats of varying size. Block sizes were computed using quadrats of 5×5 , 5×10 , 10×10 , 10×15 , 15×15 , 15×20 , 20×20 , 20×25 , 25×25 , and 25×30 units. All of the statistical analyses were conducted using SPSS 11.5 software. Figures were plotted by Origin 7.5 software.

Results

Stand Characteristics

The basal areas and densities of B. albosinensis in different stands varied (Table 1). Basal areas ranged from $18.1 \, \mathrm{m}^2 \, \mathrm{ha}^{-1}$ to $22.8 \, \mathrm{m}^2 \, \mathrm{ha}^{-1}$ and from $23.8 \, \mathrm{m}^2 \, \mathrm{ha}^{-1}$ to $27.6 \, \mathrm{m}^2 \, \mathrm{ha}^{-1}$ in the uncut and cut stands, respectively. Densities ranged from $244 \, \mathrm{stems} \, \mathrm{ha}^{-1}$ to $276 \, \mathrm{stems} \, \mathrm{ha}^{-1}$ in the uncut stands, whereas B. albosinensis were more abundant (range $465 \, \mathrm{stems} \, \mathrm{ha}^{-1}$ to $693 \, \mathrm{stems} \, \mathrm{ha}^{-1}$) and had an approximate twofold increase in the cut stands. The bamboo cover ranged from 5% to 50% among the stands. In addition to F. qinlingensis, other woody species were also found in B. albosinensis forests (Table 2). The other woody plants in the uncut stands were more abundant than those in the cut stands because of strip clearcutting.

Understory Vegetation and Stand Structure

Except the evergreen bamboo (F. qinlingensis), deciduous species characterize understory vegetation in terms of number of species (Table 2). Differences in B. albosinensis seedling and sapling densities that were observed in the uncut stands seemed were unrelated to bamboo cover (Table 1). In the cut stands, however, B. albosinensis density was lower when bamboo coverage in the stands reached 50%. Moreover, seedling and sapling densities were negatively correlated (r = -0.98, p < 0.01, Spearman rank correlation coefficient) with bamboo coverage (Table 1). Reduction in other woody plant abundance was also evident when bamboo coverage increased to 50%.

Table 2. Number of woody plants (>1.0 m tall) ha⁻¹ in the *B.ablosinensis* stands in Qinling Mountains, China.

Speices	Uncut	Uncut	Uncut	Uncut	V	VI	VII Strip clearcut	Strip clearcut
					Strip clearcut	Strip clearcut		
Pinus armandii	48	84	132	104	32	16	28	56
Betula utilis		16	48	96	24	56	112	48
Tilia latevirens	48	80	46	14	48	32	8	4
Acer robustum	56	64	36	48	6	16	32	
Abies fargesii			32	82	8	14	84	96
Ribes glaciale	68	105	76	38	16	44	80	
Populus davidiana		48	38	16	40	16	32	16
Pterocarya stenoptera						56	84	16
Rose omiensis	96	102	136	12	56	82		30
Lonicera fangutica		64	102	44	84	46	34	
Sorbus discolor	64	72	84	40				
Meliosma cuneifolia		16	32		16	32	64	
Rhododendron purdomli		32	64	96				
Salix matsudana					32	32	64	
Salix caprea	32	48	28	24				
Cerasus tomentosa					48	16	32	24
Populus purdomii	24	16	32	8	8	16		
Quercus liaotungensis	28	32	8					
Cornus controversa	24		16				44	16
Litsea tsinlingensis					8	16	32	
Sorbus koehneana	8	32		6	16		32	
Acer coesium ssp.giraldii			8	40				
Corylus mandshurica	35	24			8	4		
Acer miaotaiense					6		6	34
Populus purdomii		8	32	4				
Acer shenlauensis				36	12	6	4	
llex yunnanensis		14	58					
Pertya sinensis			6	28				
Total	531	857	1014	736	468	500	772	340
Fargesia ginlingensis	1040	790	560	320	470	580	820	1250

Blanks represent no individuals found in the plot.

doi:10.1371/journal.pone.0059375.t002

Bamboo density reduced woody plant diversity (Table 1). There was also a negative correlation (r = -0.76, p = 0.032, Spearman rank correlation coefficient) between species richness and bamboo cover. When *F. qinlingensis* densities were included in the Shannon computation, a negative correlation (r = -0.87, p = 0.005, Spearman rank correlation coefficient) was also found between species diversity and bamboo cover (Table 1).

Size Structure

The population of *B. albosinensis* showed a bell-shaped diameter distribution pattern in the uncut stands (Figure 2). *B. albosinensis* trees mainly thrived in the middle and larger diameter classes, whereas young *B. albosinensis* with DBH less than 15 cm were scarce and accounted for just 11.9% of the total individuals, suggesting poor recruitment over the past decades. *B. albosinensis* seedlings and saplings were sparser, and only several were found in each uncut stand. The diameter class distribution for *B. albosinensis* in strip clearcut stands had a reverse J shape (Figure 2), with young

stems (<20 cm) accounting for 52.8%. Recent recruitment was also abundant, suggesting successful regeneration over the past 50 years after strip clearcutting.

Spatial Patterns

Spatial distributions of small (DBH<10 cm) and larger (DBH >10 cm) *B. albosinensis* differed between the cut and the uncut stands (Figure 3). In the uncut stands, small *B. albosinensis* were clumped at almost all scales (25 m² to 750 m²), and the clumping magnitude fluctuated severely among different scales, suggesting a heterogeneously natural disturbance. In comparison, small and larger *B. albosinensis* clumps of highest intensity both occurred at small scales (25 m² to 150 m²), suggesting that recruitment in single-tree gaps was common. In the cut stands, the high clumping of small *B. albosinensis* occurred at small to intermediate scales, whereas taller *B. albosinensis* trees were clumped only at intermediate scales (200 m² to 400 m²). Such results suggested that

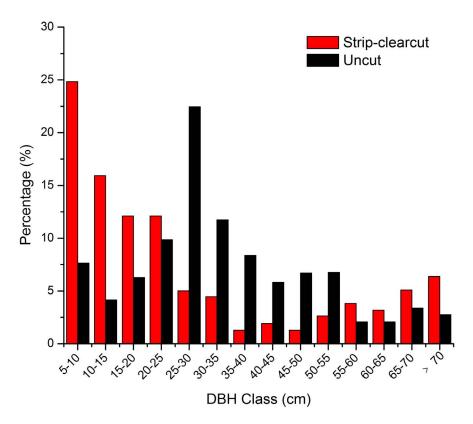


Figure 2. Size class distribution of the B. ablosinensis trees in strip clearcut and uncut B. ablosinensis forests of Qinling Mountains, China.

doi:10.1371/journal.pone.0059375.g002

B. albosinensis recruitment in the past period was initiated by large disturbance.

Discussion

Seedling and sapling pools of B. albosinensis in the uncut stands were scarce, suggesting a degeneration trend. Poor natural regeneration of B. albosinensis forests elsewhere in China has also been reported [25,26]. The absence of B. albosinensis recruitment generally indicates an unfavorable environment for regeneration. Thick forest litter is a main factor influencing regeneration because of restrictive seed germination in deciduous forests. Based on laboratory simulation experiment results, the germination rate of B. albosinensis seeds declined when covered with mulch, especially broad-leaf mulch [6,7]. Similar effects of forest litter on germination have also been observed in Betula alleghaniensis in Canada [27] and Betula maximowicziuna in Japan [28]. General field observations demonstrate that B. albosinensis are prone to germination in places with less litter fall (e.g., under open-canopy or near roads). Even after a successful germination, B. albosinensis seedlings beneath closed canopies easily die as a result of shade sensitivity, although fast-growing seedlings can compensate for less shade-tolerance.

Disturbances drive the regeneration dynamics of most closed-canopy forests by creating opportunities that facilitate the establishment of new individuals through canopy opening [8,11,29]. In the present work, poor *B. albosinensis* regeneration was observed in the uncut stands, and these mainly occurred in the canopy gaps produced by disturbances. This finding suggests that *B. albosinensis* are dependent on gaps for regeneration. The intensely clumped distributions of small (<10 cm DBH) and

larger (>10 cm DBH) *B. albosinensis* at small scales (25 m² to 150 m²) are consistent with the canopy gap sizes (20 m² to 100 m²) surveyed in the forests of the Qinling Mountains [4]. Larger *B. albosinensis* were found clumped with lower intensity, indicating a characteristic of thinned population [11]. *B. albosinensis* possess life history traits associated with the gap characteristics, and are less shade-tolerant. In addition, *B. albosinensis* produce more frequent seed crops, have smaller seeds that disperse farther, and have seedlings that grow faster. These life history traits promote the rapid colonization and early dominance of *B. albosinensis* in the gaps. Therefore, gap disturbance seems critical for the maintenance of *B. albosinensis* populations.

However, B. albosinensis recruitment in canopy gaps in this report was very limited in number, which may be related to the disturbance scale. Betula in other subalpine forests [30–32] exhibit great dependency on relatively large gaps for regeneration. B. albosinensis also appear to require large gaps for persistence. Natural disturbances that have occurred in B. albosinensis forests, including standing death and snapping of canopy trees caused by heavy snow, diseases, and climber twining, are frequently smallscale and rarely large-scale [4,14]. As a result of these disturbances, light and ground layers are only slightly changed over a small area. Most gaps are filled by either vegetation growth of the surrounding adults or replacement of shade-tolerant species that are already present as suppressed individuals. These frequent small-scale disturbances may not fulfill the demand of B. albosinensis regeneration for an environment with much light and exposed soil. Therefore, without large-scale disturbances to clear space and expose the covered land, it may be difficult for B. albosinensis stands to persist for generations to come.

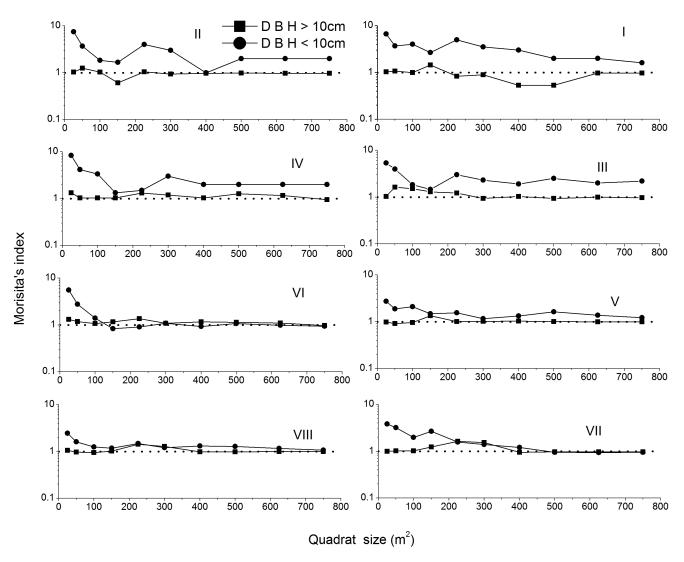


Figure 3. Values of Morisita's index (I_{∂}) for different-sized *B. ablosinensis* in 8 stands in Qinling Mountains, China. Random distribution ($I_{\delta=}$ 1.0) is shown by the dot line. doi:10.1371/journal.pone.0059375.g003

Bamboo, as a common understory plant in subalpine forests, restricts tree regeneration and species diversity [20-21,29,33]. Where bamboo fully occupies a forest understory, the frequency and number of tree seedlings and shrubs are lower because of intense competition with bamboo for space and resources. In our study, a significant negative correlation was found between understory woody plant diversity (richness and Shannon's formula) and F. qinlingensis cover, suggesting that dense bamboo reduced plant diversity. This finding is consistent with published reports on other subalpine forests where bamboos dominate the forest understory [18,34,35-37]. However, no significant correlation was found between B. albosinensis recruitments and F. qinlingensis cover despite the wide range (5% to 45%) in the uncut stands. Previous studies in mixed hardwood-conifer forests have suggested that understory bamboos impede Betula regeneration, although it did so with less intensity than conifer [36,37].

It is likely that there are other restricting factors (e.g., forest litter, closed canopy, and dense shrubs) that weaken the impeding effect of bamboo alone on *B. albosinensis* regeneration. In addition, the ability of *B. albosinensis* to disperse into gaps as well as go through a fast-growing juvenile stage may more or less help them

break the shade of bamboo layers. Therefore, bamboo density has no significant effect on the distribution and establishment of *B. albosinensis* seedlings and saplings. However, understory bamboo may contribute greatly to the persistence of *B. albosinensis* population by restricting coexisting conifer species [32,37]. Furthermore, bamboo flowering usually creates a favorable environment for *B. albosinensis* regeneration as a large forest disturbance.

Contrary to failed natural regeneration, a large number of recruitments were found in stands with subsequent regeneration approximately 50 years after strip clearcutting. *B. albosinensis* dominated the regenerating stands as a pioneer species. After strip clearcutting, the sudden exposure of previously forested lands and more sunlight reaching the forest floor seemed responsible for the successful regeneration of sunlight-loving *B. albosinensis*. The establishment of *B. albosinensis* in large numbers after strip clearcutting suggests that disturbance magnitude is important for understanding the regeneration strategy of *B. albosinensis*. However, we found that high bamboo coverage in the cut stands reduced *B. albosinensis* seedling and sapling recruitment. With vegetation restoration after the cut, *B. albosinensis* regeneration became prone

to bamboo restriction when they attained a high degree of dominance. As such, other artificial silvicultural methods, such as bamboo removal, may be necessary in improving the competitive status of *B. albosinensis* at the later stage of forest regeneration after clearcutting.

The above results confirm that strip clearcutting can prevent primary *B. albosinensis* stands from degenerating. The moderate severity of disturbance caused by strip clearcutting may be the best approach to achieve the dual objectives of forest conservation and timber production. Palynological evidence suggests that *Betula* forests have existed as zonal forests in the geological period and in modern times in the Qinling Mountains [38]. Naturally, *B. albosinensis* population may be capable of maintaining their stability as a whole despite the particularly poor regeneration. Thus, the aggregation of different-spatiotemporal cohorts driven by disturbances may be the pattern and strategy of natural *B. albosinensis*

References

- Zhang YQ (1989) Shaanxi Forest. Xi'an: Xi'an Science and Technology Press. (In Chinese.).
- Li ZY, Zhang YC, Li J (2002) The forest of Xiaolong Mountain. Tianshui: Xiaolongshan Forest Experiment Bureau. (In Chinese.).
- Fu ZJ, Guo JL (1994) Preliminary studies of Betula albosinensis forest in Taibai Mountain. Chinese Journal of Plant Ecology 18(3): 261–270. (In Chinese with English abstract.).
- Su JW, Yue M, Wang YJ (2006) Gap characteristics of Betula albosinensis forest in Taibai Mountains. Chinese Journal of Applied and Environmental Biology 12(2): 195–199. (In Chinese with English abstract.).
- Guo YX, Kang B, Li G, Wang DX, Yang GH, et al. (2011) Species composition and point pattern analysis of standing trees in secondary *Betula albosinensis* forest in Xiaolongshan of west Qinling Mountains. Chinese Journal of Applied Ecology 22(10): 2574–2580. (In Chinese with English abstract.).
- Ren JY, Lin Y, Yue M (2008) Seed germination characteristics of Betula albosinensis at Mountain Taibai, China. Chinese Journal of Plant Ecology 32(4): 883–890. (In Chinese with English abstract.).
- Wu Y, Liu Q, He H, Liu B, Yin H (2004) Effects of light and temperature on seed germination of *Picea asperata* and *Betula albosinensis*. Chinese Journal of Applied Ecology 15(12): 2229–2232. (In Chinese with English abstract.).
- Cullen LE, Stewart GH, Duncan RP, Palmer JG (2001) Disturbance and climate warming influences on New Zealand Nothofagus tree-line population dynamics. Journal of Ecology 89: 1061–1071.
- Antos JA, Parish R (2002) Structure and dynamics of a nearly steady-state subalpine forest in south-central British Columbia, Canada. Oecologia 130: 126-125
- Yahner RH (2000) Eastern Deciduous Forest: Ecology and Wildlife Conservation. Minneapolis, MN: University of Minnesota Press.
- Veblen TT, Donoso C, Schlegel FM, Escobar B (1981) Forest dynamics in south-central Chile. Journal of Biogeography 8: 211–247.
- Pebyelas J, Ogaya R, Boada M, Jump AS (2007) Migration, invasion and decline: changes in recruitment and forest structure in a warming-linked shift of European beech forest in Catalonia (NE Spain). Ecography 30, 829–837.
- Takahashi A, Koyama H, Takahashi N (2008) Habitat expansion of Robinia pseudoacacia L. and role of seed banks in the Akagawa River basin. Journal of the Japanese Forest Society 90: 1–5.
- Lin Y, Ren JY, Yue M (2008) Population structure and spatial analysis of Betula albosinensis at Taibai Moutain, northwestern China. Chinese Journal of Plant Ecology 32(6): 1335–1345. (In Chinese with English abstract.).
- Li JJ, Zhu ZC, Min ZL (1989) Comprehensive Suvey of the Taibai Mountain Preserve. Xi'an: Shaanxi Normal University Press. (In Chinese.).
- Basnet K (1992) Effect of topography on the pattern of trees in Tabonuco (Dacryodes excelsa) dominated rain forest of Puerto Rico. Biographica 24: 31–42.
- Allison D, Art W, Cunningham E, Teed R (2003) Forty-two years of succession following strip clearcut in a northern hardwoods forest in northwestern Massachusetts. Forest ecology and management 182: 285–301.
- Taylor AH, Shiwei J, Lianjun Z, Chunping L, Changjin M, et al. (2006) Regeneration patterns and tree species coexistence in old-growth Abies-Picea forests in southwestern China. Forest Ecology and Management 223: 303–317.
- Gonzalez ME, Veblen TT, Donoso C, Valeria L (2002) Tree generation response in a lowland Nothofagus-dominated forest after bamboo die-back in south-central Chile. Plant Ecology 161: 59–73.

population stability. However, persistent and periodic artificial disturbance by strip clearcutting is necessary from a forest production perspective.

Acknowledgments

Great thanks to the Mt. Taibai National Nature Reserve and the Mt. Xiaolong National Nature Reserve for their helps in field work and logistical support. We are also grateful to Junjie Yang and Lingtong Kong for the assistance in field work.

Author Contributions

Provided direction and suggestions for the manuscript: GL DW GY. Conceived and designed the experiments: YG GL. Performed the experiments: YG DK YH. Analyzed the data: YG. Wrote the paper: YG.

- Narukawa Y, Yamamoto S (2002) Effects of coniferous and broad-leaved trees in a Japanese temperate mixed forest. Journal of Vegetation Science 2,413–418.
- Dang HS, Zhang YJ, Zhang KR, Jiang MX, Zhang QF (2010) Age structure and regeneration of subalpine fir (Abies fargesii) forests across an altitudinal range in the Qinling Mountains, China. Forest Ecology and Management 259: 547– 554.
- Ren Y, Liu MS, Tian LH, Tian XH, Li ZJ (2006) Biodiversity, conversation and management of Taibaishan Nature Reserve. Beijing: China forestry Publishing House. (In Chinese.).
- Shannon CE, Weaver W (1949) The Mathematical Theory of Communication. Urbana, IL: University of Illinois Press.
- Morisita M (1959) Measuring of the dispersion of individuals and analysis of the distributional patterns. Memoirs of the Faculty of Science Kyushu University, Series E 2: 215–235.
- Liu SJ, Su ZX (2004) T-square study on spatial pattern and regeneration of Betula albosinensis population on west slope of Jiuding Mountain. Chinese Journal Applied Ecology 15(1): 1–4. (In Chinese with English abstract.).
- Miao N, Liu SR, Shi ZM, Yu H, Liu XL (2009) Spatial pattern of dominant tree species in subpine Betula-Abies forest in west Sichuan of China. Chinese Journal of Applied Ecology 20(6), 1263–1270. (In Chinese with English abstract.).
- Peterson CJ, Facelli JM (1992) Contrasting germination and seedling growth of Betula allenghaniensis Britton and Rhus typhina L. subjected to various amounts and type of plant litter. American Journal of Botany 79: 1209–1216.
- Osumi K, Sakurai S (1997) Seedling emergence of Betula maximowicziana following human disturbance and the role of buried viable seeds. Forest Ecology and Management 93: 235–243.
- Taylor AH, Qin Z, Liu J (1996) Structure and dynamics of subalpine forests in the Wang Lang Natural Reserve, Sichuan, China. Vegetation 124: 25–38.
- Koyama T (1984) Regeneration and coexistence of two Abies species dominating subalpine forests in central Japan. Oecologia 62: 156–161.
- Nakamura T (1985) Forest succession in the subalpine region of Mt, Fuji, Japan. Vegetation 64: 15–27.
- Taylor AH, Qin Z (1988) Tree replacement patterns in subalpine Abies-Betula forests, Wolong Natural Reserve, China. Vegetation 78: 141–149.
- Takahashi K, Kohyama T (1999) Size-stucture dynamics of two conifers in relation to understorey dwarf bamboo: a simulation study. Journal of Vegetation Science 10: 833–842.
- Veblen TT, Veblen AT, Schlegel FM (1979) Understorey patterns in mixed evergreen-deciduous Nothofagus forests in Chile. Journal of Ecology 67: 809– 823.
- Peet RK (1981) Forest vegetation of Colorado Front Range: composition and dynamics. Vegetation 45: 3–75.
- Taylor AH, Qin Z (1988) Regeneration patterns in old-growth Abies-Betula forests in the Wolong National Reserve, Sichuan, China. Journal of Ecology 76: 1204–1218.
- Wang W, Franklin SB, Ren Y, Ouellette JR (2006) Growth of bamboo Fargesia qinlingensis and regeneration of trees in a mixed hardwood-conifer forest in the Qinling Mountains, China. Forest Ecology and Management 234: 107–115.
- Zhu ZC (1991) Stability of the Betula forest in the Taibai mountain of Qinling mountain range. Journal of Wuhan Botanical Research 9: 169–175. (In Chinese with English abstract.).