

Carbon storage in biomass, litter, and soil of different plantations in a semiarid temperate region of northwest China

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Abstract

• **Context** A large area of abandoned land in the semiarid temperate region of China has been converted into plantations over the past decades. However, little information is available about the ecosystem C storage in different plantations.

• **Aim and methods** Our objective was to estimate the C storage in biomass, litter, and soil of four different plantations (monospecific stands of *Larix gmelinii*, *Pinus tabulaeformis*, *Picea crassifolia*, and *Populus simonii*). Tree component biomass was estimated using allometric equations. The biomasses of understory vegetation and litter were determined by harvesting all the components. C fractions of plant, litter, and soil were measured.

• **Results** The ecosystem C storage were as follows: *Picea crassifolia* (469 t C/ha) > *Larix gmelinii* (375 t C/ha), *Populus simonii* (330 t C/ha) > *Pinus tabulaeformis* (281 t C/ha) ($P < 0.05$), 59.5–91.1 % of which was in the soil. The highest tree and understory C storage were found in the plantation of *Pinus tabulaeformis* (247 t/ha) and *Larix gmelinii* (1.2 t/ha) respectively. The difference in tree C fraction was significant among tree components ($P < 0.05$), following the order: leaf > branch > trunk > root. The highest soil C (SC) was stored in *Picea crassifolia* plantation (411 t C/ha), while *Populus*

simonii plantation had a higher SC sequestration rate than others.

• **Conclusion** C storage and distribution varied among different plantation ecosystems. Coniferous forests had a higher live biomass and litter C storage. Broadleaf forests had considerable SC sequestration potential after 40 years establishment.

Keywords Biomass carbon · Litter · Semiarid region · Soil carbon · Tree species · Understory vegetation

1 Introduction

The forest ecosystems have large potential to act as a temporary and long-term carbon (C) pool (Dixon et al. 1994). Approximately 80 % aboveground and 40 % underground terrestrial C are stored in forests (Cao and Woodward 1998). However, in the temperate semiarid region of northwest China, many natural forests are degraded to abandoned lands because of climate change and human disturbance. The Chinese government has imposed the ‘Grain-for-Green’ project (i.e., conversion of cropland to forest and grassland) since 1999 (Zhang et al. 2010). Large areas of abandoned land have been converted into forest lands (Fang et al. 2001). Establishment of forest plantation contributes to soil and water conservation, and biodiversity protection (Chazdon 2008). Also, afforestation is regarded as an effective measure to prevent the global warming by sequestering C both in biomass and in soil. Trees and understory vegetation assimilate carbon dioxide (CO₂) from the atmosphere and store C in plant biomass. Accordingly, the storage of soil carbon (SC) increases with the large amount of C input from litterfall and rhizodeposition (Vesterdal et al. 2012).

The magnitude and progress of the changes in C storage following afforestation are highly various because of the influence of different factors, such as climatic condition, soil

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property, tree species, and present management (Jobbágy and Jackson 2000; Paul et al. 2002). Among them, tree species strongly affect the C accumulation of plantation ecosystems in several ways (Díaz-Pinés et al. 2011). Firstly, tree species influences the C storage of live biomass as a result of different tree traits. One of the most determinant tree traits is the growth rate. Fast-growing species, such as poplar, pine, and eucalypt are widely planted for C sequestration (Kaul et al. 2010). However, the fast biomass increment brings in a lower C fraction, suggesting that planting fast-growing tree species sacrifices some C gain by decreasing their C fraction (Zhang et al. 2009). The species-specific C fraction imposes its effect on biomass C storage by varying with growth rates. Moreover, C sequestration in trees is also influenced by the variation of wood density. At identical volume, trees with higher wood density (most deciduous species) accumulate more C than those with light wood density (most coniferous species) (Jandl et al. 2007).

Secondly, interspecific differences in the production and decomposition rate of litters explain the variations in litter C storage (Finzi et al. 1998). Deciduous tree species have a higher annual production of litters than the evergreen coniferous species. However, previous research showed that pine and spruce had more litter C than beech and oak (Jandl et al. 2007). It is mainly attributed to the slow decay rate of needle litters determined by the chemical composition, such as soluble carbohydrates and lignin concentrations (Paul et al. 2002). The decay of litters not only depends on its own property, but also on stand microclimatic conditions. Pérez-Cruzado et al. (2012) demonstrated that light transmission was higher in *Eucalyptus* than in *Pinus* stands determined by the arrangement of the leaves. The higher light transmission leads to a favorable soil temperature and moisture level for decomposers, and then promotes the decay of litters (Martius et al. 2004).

Finally, tree species also influences the amount, quality, and distribution of soil C (SC) by regulating C input (litterfall and rhizodeposition) and C output (respiration and leaching) (Vesterdal et al. 2012). For instance, pine forests have remarkably low SC storage, whereas beech forests have the highest SC storage among common European tree species (Jandl et al. 2007). As mentioned above, coniferous forest had lower litter decomposition rate than deciduous forest, slowing down the C input from litter to the soil. Moreover, the rooting depth is relevant for SC, because root growth is the most effective way of introducing C to the soil (Jobbágy and Jackson 2000). Coniferous species with shallow roots tended to accumulate less C in the soil. In addition, Vesterdal et al. (2012) reported that the soil respiration increased in the order beech < lime < spruce, oak, maple < ash. Thus, C outputs by soil respiration of different tree species were various, which partly influenced the SC storage.

Recently, many case studies involved in forest ecosystem C storage and dynamics have been conducted by Chinese

researchers, using forestry survey data (Fang et al. 2001; Guo et al. 2010). However, studies about C storage of plantation ecosystems in the semiarid region are scarce, and the quantification of total plant biomass C, including understory vegetation and litter, is even rarer. The objectives of this study were (1) to find which plantation ecosystem has higher C sequestration capacity, and (2) to explore the distribution of C storage in the plant biomass, litter, and soil. The information concerning C storage at the ecosystem level would provide more accurate assessments of the C sequestration.

2 Materials and methods

2.1 Study area

The study was carried out in Ningxia Hui Autonomous Region (northwest China), which has a continental climate. The mean annual temperature is between 5 °C and 9 °C. The mean monthly maximum temperature is in July, and the mean monthly minimum temperature is in January. The mean annual precipitation is between 180 mm and 800 mm. The majority of precipitation falls during the period July to September. Typical soil types in the region are gray cinnamon soil, black loam, and aeolian soil. The average pH of soil here ranges between 6.5 and 8.5 (Sun et al. 2009). Plantations here are monospecific stands dominated by larch, pine, spruce, poplar and etc. The understory shrub species are *Cotoneaster acutifolius*, *Corylus heterophylla*, *Spiraea mongolica*, *Elaeagnus pungens*, *Lonicera microphylla*, *Sorbaria sorbifolia* and etc. The understory herb species are *Eragrostis pilosa*, *Cymbopogon citratus*, *Fragaria orientalis*, *Carex duriuscula*, *Heteropappus altaicus*, *Artemisia subdigitata* and etc.

2.2 Experimental setup, sampling, and laboratory analysis

Larix gmelinii, *Pinus tabulaeformis*, *Picea crassifolia*, and *Populus simonii* stands were selected as tested sites. Abandoned lands (the initial land use before afforestation) in the main experiment area served as control sites (Table 1). In each plantation and control site, three replicated 50 m × 20 m plots (at least 100 m away from each other) were established. Each plot was separated into ten 10 m × 10 m quadrats. In all the quadrats, the height and diameter at breast height (DBH) for each individual tree were measured to estimate tree biomass. Five samples of tree leaf, branch, trunk, and root were randomly collected for C fraction determination. All understory vegetation (shrub and herb) was harvested from three subquadrats (2 m × 2 m) randomly located in each quadrat. Shrubs were separated into leaves, branches, and roots; herbs were separated into aboveground and belowground parts. All litter was collected from three subquadrats (1 m × 1 m) located within each quadrat. Samples of understory vegetation and

Table 1 Description of the study sites

Site	Property				
Plantation	Tree species	<i>Larix gmelinii</i>	<i>Pinus tabuliformis</i>	<i>Picea crassifolia</i>	<i>Populus simonii</i>
	Altitude (m)	2,300	2,150	2,710	2,080
	Forest type	Deciduous	Evergreen	Evergreen	Deciduous
		Conifer forest	Conifer forest	Conifer forest	Broadleaf forest
	Soil type	Gray cinnamon soil	Gray cinnamon soil	Gray cinnamon soil	Gray cinnamon soil
	Stand age (a)	46	32	48	40
	Density (tree/ha)	1,520±64	2,830±256	1,570±144	1,420±51
	<i>H</i> (m)	12.8±0.3	15.0±1.0	12.4±3.1	7.4±0.8
	<i>D</i> (cm)	10.8±0.2	7.9±0.7	6.9±1.6	5.8±0.6
	Control	Land use	Abandoned land	Abandoned land	Abandoned land
Vegetation type		Grassland	Grassland	Grassland	Grassland
Soil type		Gray cinnamon soil	Gray cinnamon soil	Gray cinnamon soil	Gray cinnamon soil

Data of density, *H* and *D* represent the mean ± standard deviation

H is tree height, *D* is diameter at breast height

litter were dried at 65 °C to constant weight for determination of biomass and C fraction.

Ten representative soil samples were randomly collected for each plot (50 m × 20 m) at depths of 0–10 cm, 10–20 cm, 20–30 cm, 30–50 cm, and 50–100 cm using a soil core (5 cm diameter). Soil samples from the same layer in each plot (plantation plots and control plots) were mixed for a more representative sample for the measurement of soil C (SC) fraction. In addition, soil cores (5-cm-height, 5-cm-diameter) with two replications of each plot were sampled for bulk density measurement as suggested by the Chinese Editorial Committee of Soil Analysis (1996).

The plant, litter, and soil samples were dried and ground to pass through 0.25-mm sieve prior to the laboratory analysis. The C fraction was determined using an Elementar Vario EL cube Analyzer.

2.3 Destructive tree sampling

To estimate the biomass of tree components, we harvested 20 trees of each species in the sites with similar characteristics to the study area. These trees were selected with the aim of covering the range of DBH from 5 cm to 40 cm. Stems were cut at the ground level. Total tree height and DBH were measured and recorded. The trunk was marked into three parts (top, middle, and bottom), cut into 1 m sections, and weighed. At the end of each trunk section, a 5-cm thick disc was cut, weighed, and taken to laboratory for moisture content determination. Branches were also separated into three different size components (diameter >20 cm, 5–20 cm, and <5 cm). All the leaves were collected in the field. The stump and coarse root (diameter ≥ 5 mm) of

sample trees was excavated manually. The fresh weight of each component (trunk, branch, leaf, and root) was measured to the nearest 1.0 g by using an electronic balance. Approximately 500 g of fresh samples of each tree component were randomly collected for moisture content determination. Dry weight was obtained by drying the samples at 65 °C until they reached a constant weight. The total dry biomass of each component was calculated through multiplying the fresh weight by the dry/wet ratio. Biomass for the whole tree was calculated by summing the biomass of trunk, branch, leaf, and root.

2.4 Data analysis

Statistical analysis was performed by the software SPSS, ver. 16.0 (SPSS Inc., Chicago, IL, USA). Allometric equations between tree component biomass and independent variable (squared DBH multiplied by the tree height (D^2H)) were developed using curve fitting. The optimum equations were selected to calculate the tree component biomass in the plantation sites. Plant and litter C storage were calculated from the C fraction multiplied by component biomass. SC storage was calculated from the SC fraction multiplied by the bulk density and the thickness of the soil layer. SC sequestration rate was calculated by subtracting the SC storage of the control site from that of the plantation and then dividing the result by stand age. ANOVA analyses were used to determine the statistically significance differences between species for the biomass, C fraction, and C storage; multiple comparisons were carried out by Duncan's method, with differences in the $P < 0.05$ significance level.

Table 2 Allometric equations for different tree species and component

Tree component	Allometric equations	R ²	SEE	F
<i>Larix gmelinii</i>				
Leaf	B=0.113(D ² H) ^{0.406}	0.841	0.282	184.391***
Branch	B=0.373(D ² H) ^{0.397}	0.920	0.187	184.149***
Trunk	B=0.241(D ² H) ^{0.552}	0.933	0.236	223.137***
Root	B=1.011(D ² H) ^{0.321}	0.813	0.246	69.347***
Whole tree	B=1.282(D ² H) ^{0.465}	0.962	0.147	406.789***
<i>Pinus tabuliformis</i>				
Leaf	B=0.007(D ² H) ^{0.890}	0.896	0.412	164.088***
Branch	B=0.001(D ² H) ^{1.246}	0.924	0.486	231.508***
Trunk	B=0.016(D ² H) ^{1.020}	0.911	0.435	193.603***
Root	B=0.005(D ² H) ^{1.060}	0.963	0.284	489.867***
Whole tree	B=0.031(D ² H) ^{1.039}	0.938	0.364	287.694***
<i>Picea crassifolia</i>				
Leaf	B=0.056(D ² H) ^{0.608}	0.918	0.251	212.772***
Branch	B=1.334(D ² H) ^{0.421}	0.826	0.267	90.369***
Trunk	B=0.073(D ² H) ^{0.805}	0.912	0.346	197.608***
Root	B=0.193(D ² H) ^{0.617}	0.832	0.384	94.062***
Whole tree	B=1.026(D ² H) ^{0.602}	0.899	0.280	168.304***
<i>Populus simonii</i>				
Leaf	B=1.264(D ² H) ^{0.262}	0.734	0.464	37.810***
Branch	B=1.017(D ² H) ^{0.472}	0.776	0.554	24.204***
Trunk	B=0.464(D ² H) ^{0.646}	0.884	0.484	57.222***
Root	B=0.483(D ² H) ^{0.550}	0.809	0.498	30.294***
Whole tree	B=1.973(D ² H) ^{0.561}	0.835	0.354	80.870***

D²H is squared DBH (D, (cm)) multiplied by tree height (H, (m)), B is the biomass of tree component (kg/tree), SEE is the standard error of estimate
*** means the equation is significant ($\alpha=0.001$)

3 Results

3.1 Biomass and C storage of tree layer

According to the component allometric equations (Table 2), we estimated the tree biomass of different plantations. Individual tree biomass (kg/tree) followed the order *Pinus tabuliformis*>*Picea crassifolia*>*Populus simonii*>*Larix*

gmelinii (Table 3), and annual biomass increment was 2.7 ± 0.4 kg/tree/year, 1.4 ± 0.1 kg/tree/year, 1.5 ± 0.2 kg/tree/year, and 0.8 ± 0.2 kg/tree/year respectively. There was a common tendency for all the species that trunk is a large proportion of tree biomass. This pattern was particularly evident in *Pinus tabuliformis*, whose trunk represented 55.8 % of the total biomass. Biomass of leaves in all the four plantations was relatively small, accounting for 5.9–12.8 % of the total biomass. *Picea crassifolia* and *Populus simonii* had higher branch proportion, while *Larix gmelinii* had extremely higher root proportion when compared with other tree species.

There were significant interspecific differences in the stand tree biomasses ($F=9.577$, $P=0.005$) and C storage ($F=12.072$, $P=0.002$) (Fig. 1). The stand tree biomass was 56 t/ha in the *Larix gmelinii* plantation, 247 t/ha in the *Pinus tabuliformis* plantation, 106 t/ha in the *Picea crassifolia* plantation, and 146 t/ha in the *Populus simonii* plantation. Accordingly, tree C storage decreased in the order of *Pinus tabuliformis*>*Picea crassifolia*>*Populus simonii*>*Larix gmelinii*. Two-way ANOVA analysis showed that the difference in tree C fraction was not statistically significant among the tree species ($P>0.05$), but significant among the tree components ($P<0.05$) (Table 4). C fraction of different tree components decreased following the order leaf>branch>trunk>root.

3.2 Biomass and C storage of understory layer

Shrub biomass ranged between 1.0 t/ha and 2.4 t/ha, and decreased as following order, *Picea crassifolia*, *Populus simonii*, *Larix gmelinii*>*Pinus tabuliformis* ($F=32.960$, $P<0.001$). Shrub branch generally contributed to higher biomass proportion than leaf or root did in three plantations (*Picea crassifolia*, *Larix gmelinii* and *Pinus tabuliformis*). Whereas shrub roots of *Populus simonii* plantation had the highest biomass proportion (Fig. 2). Herb biomass was quite various among plantations ($F=39.880$, $P<0.001$), ranging between 0.3 t/ha and 1.0 t/ha. The belowground biomass of herb was about twice of the aboveground biomass.

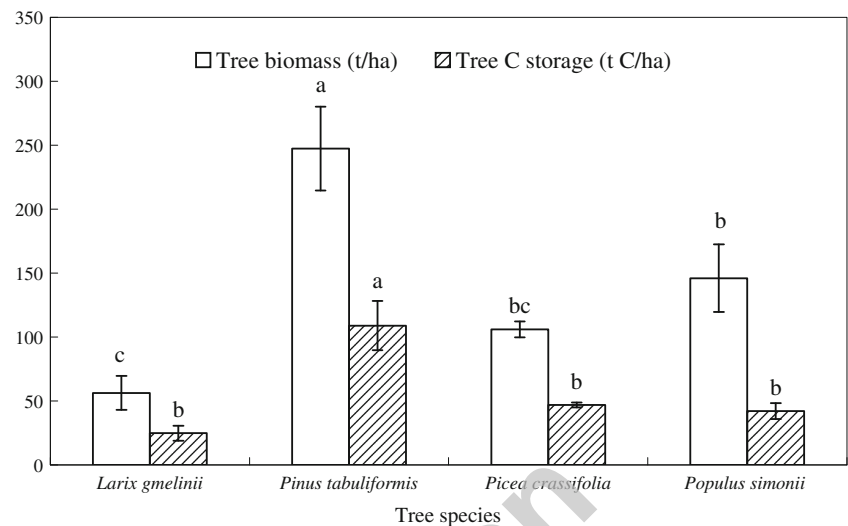
The *Picea crassifolia* plantation had significantly lower shrub C fraction than other plantations (Table 5). For different

Table 3 Biomass of tree components (kg/tree) of different plantations

Tree species	Tree component				Total tree
	Leaf	Branch	Trunk	Root	
<i>Larix gmelinii</i>	2.1±0.2 (5.7)	6.6±0.6 (17.6)	13.5±1.8 (45.4)	10.3±0.8 (27.3)	37.5±4.2c
<i>Pinus tabuliformis</i>	6.1±0.4 (7.1)	15.0±1.7 (17.4)	47.9±3.4 (55.8)	16.8±1.6 (19.6)	85.7±7.9a
<i>Picea crassifolia</i>	4.0±0.1 (5.9)	25.2±0.6 (36.9)	24.2±1.0 (35.6)	14.8±0.5 (21.7)	68.2±2.4b
<i>Populus simonii</i>	5.9±0.3 (10.4)	17.0±1.7 (29.9)	22.7±3.1 (39.8)	13.0±1.5 (22.9)	57.0±6.7b

Data represent the mean ± standard deviation, basing on measured tree D and H in Table 1 and the equations described in Table 2. Significant differences among plantations are indicated with lowercase letters ($\alpha=0.05$). Values in brackets are percentages of component biomass to the total tree biomass

Fig. 1 Tree biomass and C storage of different plantations. Significant differences in tree biomass or tree C storage among plantations are indicated with lowercase letters ($\alpha=0.05$). Vertical lines are standard deviation



shrub components, C fractions of leaf (43.7 ± 1.8 %) and branch (43.0 ± 1.1 %) were higher than that of root (40.8 ± 2.6 %) ($P < 0.05$). In addition, herb C fraction in aboveground biomass was significantly higher than that in belowground biomass ($P < 0.05$). C storage of understory vegetation ranged between 0.5 t C/ha and 1.0 t C/ha among different plantations. The shrub:herb ratio of C storage was much higher in *Populus simonii* plantation (13.4) than that in other plantations (Fig. 3).

3.3 C storage of litter layer

Biomass of litter ranged from 9.7 t/ha to 31.6 t/ha and significantly differed among the four plantations ($F = 11.116$, $P = 0.003$) (Table 6). *Picea crassifolia* plantation had significantly higher litter biomass than other plantations ($P < 0.05$), while *Populus simonii* plantation had the lowest one. Litter C storage ranged between 2.4 t C/ha and 7.9 t C/ha. The annual increment of litter C storage decreased in the following sequence, *Picea crassifolia* > *Larix gmelinii*, *Pinus tabuliformis* > *Populus simonii* ($F = 30.262$, $P = 0.003$).

3.4 SC fraction and storage

The mean values of SC fraction decreased with increasing of soil depth (Fig. 4). This pattern was particularly evident in

Pinus tabuliformis plantation. SC fraction decreased by 75.5 % from 0–10 cm to 50–100 cm in *Pinus tabuliformis* plantation. Throughout the soil profile, SC fraction varied among the plantations, and decreased in the order *Picea crassifolia* > *Larix gmelinii* > *Populus simonii* > *Pinus tabuliformis*. *Picea crassifolia* plantation had the highest SC storage (411 t C/ha), followed by *Larix gmelinii* (342 t C/ha) and *Populus simonii* (283 t C/ha) plantations (Table 5). *Pinus tabuliformis* plantation had extremely lower SC storage than other plantations ($P < 0.05$). SC sequestration rates were higher in *Populus simonii* and *Picea crassifolia* plantations, while lower in *Larix gmelinii* and *Pinus tabuliformis* plantations.

3.5 Forest ecosystem C storage

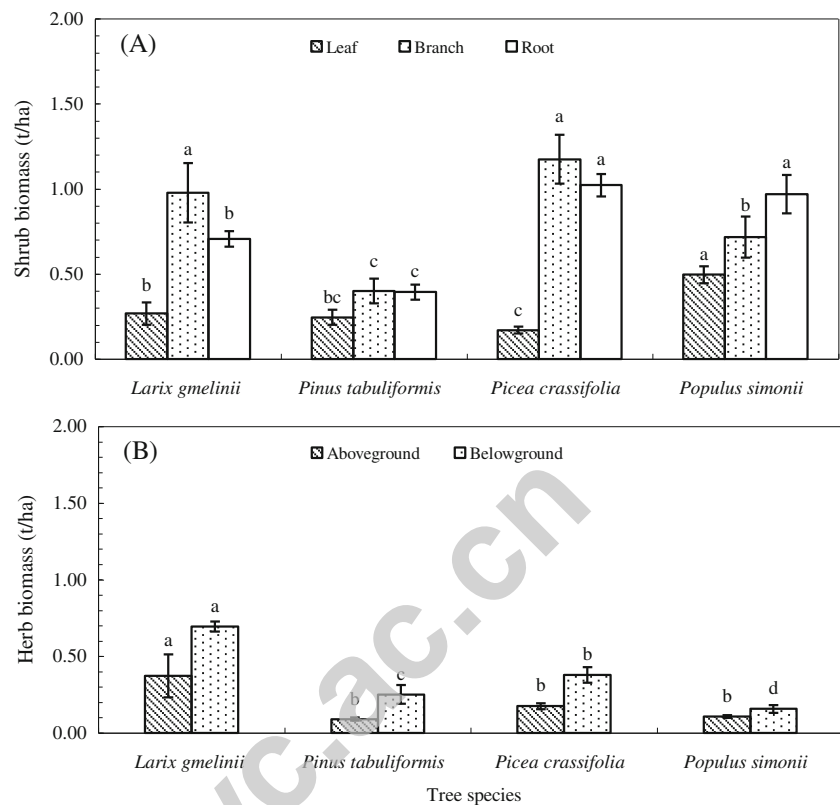
We compared the C storage in live biomass, litter, soil, and the whole ecosystem (Fig. 5). There were significant differences in forest ecosystem C storage among the plantations ($F = 30.993$, $P < 0.001$). *Picea crassifolia* plantation had significantly higher litter C storage (7.9 t C/ha), SC storage (411 t C/ha), and ecosystem C storage (469 t C/ha) than the other plantations ($P < 0.05$). In contrast, *Pinus tabuliformis* had the lowest ecosystem C storage (281 t C/ha) but the highest biomass C storage (110 t C/ha). SC storage proportion of the

Table 4 Tree component C fraction (%) of different plantations

Component	<i>Larix gmelinii</i>	<i>Pinus tabuliformis</i>	<i>Picea crassifolia</i>	<i>Populus simonii</i>	Mean
Leaf	45.3±0.8	47.9±0.7	46.0±0.7	44.0±1.6	46.8±1.3a
Branch	47.6±1.2	46.3±2.4	44.9±2.9	44.5±2.4	45.6±2.0ab
Trunk	46.9±1.3	44.2±1.2	45.3±2.0	45.6±0.4	45.1±1.7b
Root	47.4±0.2	44.0±0.5	44.3±1.0	43.8±0.6	44.4±1.3b
Mean	45.8±1.6a	45.8±2.4a	45.5±1.5a	44.9±1.6a	

Data represent the mean ± standard deviation. Significant differences among components or tree species are indicated with lowercase letters ($\alpha=0.05$)

Fig. 2 Component biomass of understory vegetation [shrub (A) and herb (B)] of different plantations. Significant differences in shrub and herb component biomass among plantations are indicated with lowercase letters ($\alpha=0.05$). Vertical lines are standard deviation



ecosystem C storage ranged from 59.5 to 91.1 %, while litter C storage proportion ranged from 0.7 to 1.5 %.

4 Discussion

There were significant differences in tree biomass among the four plantations. The tree biomass was in the order of *Pinus tabuliformis* > *Picea crassifolia* > *Populus simonii* > *Larix gmelinii* (Fig. 1). The interspecific differences in tree biomass were mainly caused by inherent variations in growth rates

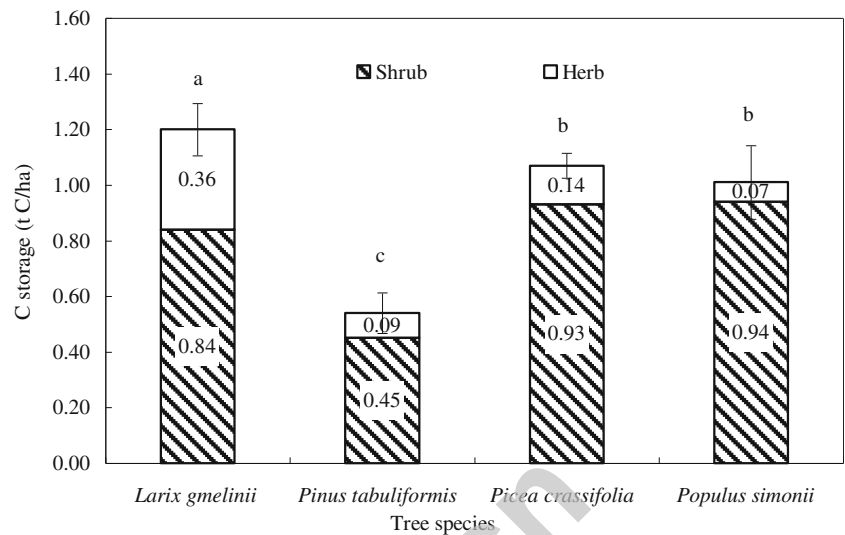
(Houghton 2005). In this study, *Pinus tabuliformis* had the highest tree biomass (247 t/ha) and growth rate (2.7 ± 0.4 kg/tree/year). As reported by Zheng and Shangguan (2007), *Pinus tabuliformis* kept a stable photosynthetic rate over a range of climates. Also, its strong drought tolerance was exhibited in the leaf morphological acclimation, such as leaf mass per area, chlorophyll content and etc. Therefore, *Pinus tabuliformis* was well-adapted to the semiarid Loess Plateau. In addition, the tree biomass of *Pinus tabuliformis* plantation was higher than the mean biomass (73–184 t/ha) observed by previous research (Ma 1989). This may be due to the high

Table 5 C fraction (%) of understory vegetation in different plantations

	<i>Larix gmelinii</i>	<i>Pinus tabuliformis</i>	<i>Picea crassifolia</i>	<i>Populus simonii</i>	Mean
Shrub					
Leaf	43.3±1.3	45.0±1.0	41.6±2.2	45.4±1.0	43.7±1.8a
Branch	43.7±0.2	43.4±0.8	41.9±0.8	44.0±0.5	43.0±1.1a
Root	41.7±1.4	41.8±0.3	36.3±1.9	41.1±0.2	40.8±2.6b
Mean	42.9±1.3a	43.4±1.5a	39.9±3.1b	43.5±2.0a	
Herb					
Aboveground	41.4±1.0	37.4±3.4	35.4±0.8	34.7±1.4	37.9±3.2a
Belowground	39.8±4.4	31.6±1.1	31.0±3.0	33.9±0.8	33.0±4.4a
Mean	40.6±6.9a	34.5±9.0b	33.2±8.1b	34.3±5.9b	

Data represent the mean ± standard deviation. Significant differences among components or tree species are indicated with lowercase letters ($\alpha=0.05$)

Fig. 3 Understory C storage of different plantations. Significant differences among plantations are indicated with lowercase letters ($\alpha=0.05$). Vertical lines are standard deviation



stand density ($2,830 \pm 256$ tree/ha), which could maximize the tree biomass (Jandl et al. 2007).

C storage of understory layer decreased in an inverse trend to that of tree layer (Fig. 2). There was a significant negative correlation between understory C storage and tree C storage ($r = -0.836, P = 0.001$). The result can be partly attributed to the increasing of tree cover canopy with the increment of tree biomass. An increased canopy closure resulted in a dramatic change in understory microclimatic condition and resources availability (Abdallah and Chaieb 2012). For example, plantations characterized by a higher canopy closure prevented larger shares of sunlight from reaching the understory layer (von Arx et al. 2012). The photosynthesis of understory vegetation was suppressed, and the biomass and C accumulation was decreased. Moreover, a lower air temperature caused by a high canopy closure, particularly during the growing season, elevated metabolic rates of understory vegetation (Holst et al. 2004). High metabolic rates brought the relative lower biomass accumulation.

In this study, *Populus simonii* plantation had significantly lower litter C storage than *Larix gmelinii* and *Picea crassifolia* plantations (Table 6). To explain this result, the deciduous leaf

litter exhibited a greater decomposition rate than other tree leaf litter (Schulp et al. 2008). Its greater decomposition rate was mainly caused by the high concentration of soluble carbohydrates and low concentration of lignin (Hobbie et al. 2000). In addition, *Populus simonii* plantation was located in a lower altitude where there was a higher temperature to accelerate the decomposition (Table 1). *Picea crassifolia* plantation had the highest litter C storage among plantations. This was attributed to the species-specific influence on the C-mineralization patterns in litter layer. It was reported that nitrification of litter layer beneath the spruce stand was very low (Trum et al. 2011). Thus, the tree species had played an important role in C accumulation in litter layers.

Table 6 Litter biomass, C fraction and C storage and annual increment in litter C storage of different plantations

Tree species	Biomass (t/ha)	C fraction (%)	C storage (t C/ha)	Annual C storage increment (t C/ha/year)
<i>Larix gmelinii</i>	27.6±5.1a	21.2±4.0a	5.7±0.5b	0.12±0.01b
<i>Pinus tabuliformis</i>	12.0±1.5b	30.2±2.6a	3.6±0.6c	0.11±0.02b
<i>Picea crassifolia</i>	31.6±5.7a	25.9±6.4a	7.9±1.0a	0.16±0.02a
<i>Populus simonii</i>	9.7±2.1bc	25.8±5.1a	2.4±0.6c	0.06±0.01c

Data represent the mean ± standard deviation. Significant differences among plantations are indicated with lowercase letters ($\alpha=0.05$)

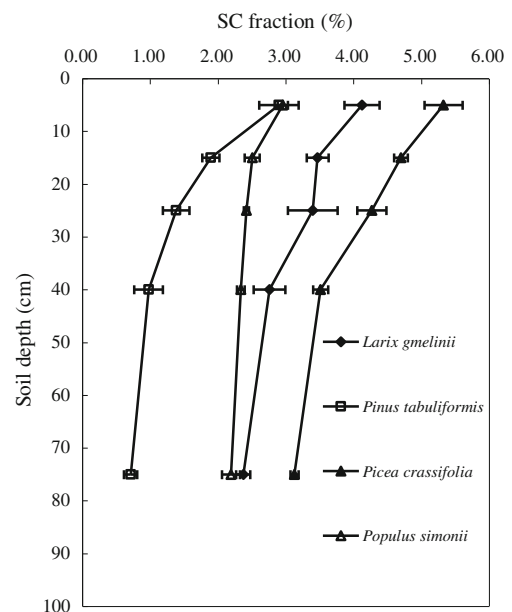
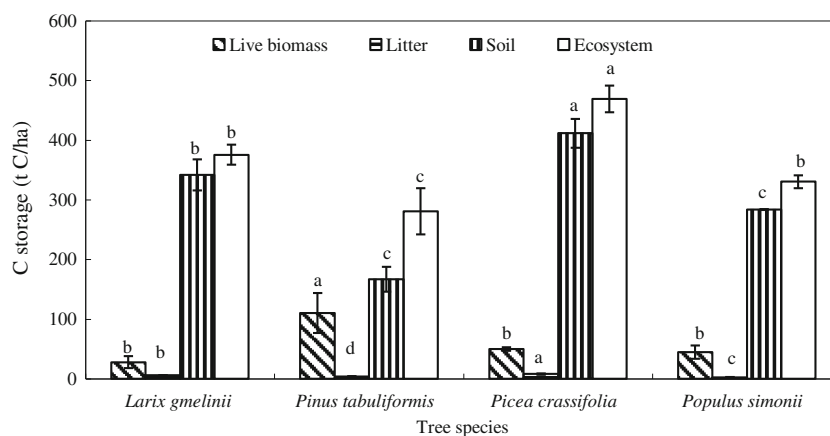


Fig. 4 SC fraction (0–100 cm) of different plantations. Horizontal lines are standard deviation

Fig. 5 Ecosystem C storage of different plantations. Significant differences in live biomass, litter, soil and ecosystem C storage among plantations are indicated with lowercase letters ($\alpha=0.05$). Vertical lines are standard deviation



SC sequestration rate in the *Populus simonii* plantation was higher than that in the three coniferous plantations (Table 7). Differences in SC sequestration among tree species may be attributable to the various C input or output patterns (Jandl et al. 2007). For this study, the most determinant factor is the rate of transfer of C from litter and root to soil. Litterfall from the *Populus simonii* plantation decayed faster than that from other plantations due to its chemical composition and micro-climatic condition. Moreover, deciduous trees presented a deeper rooting pattern than the coniferous trees, which resulted in a higher rate of root production and turnover (Johnson 1992). The greater C input from both litter and root decomposition sequestered a higher SC in the *Populus simonii* plantation. In contrast, *Larix gmelinii* and *Pinus tabuliformis* plantation had negative SC sequestration rate (Table 7). The explanation was that *Larix gmelinii* and *Pinus tabuliformis* planting had disturbed soil properties and stimulated the mineralization of SC. These losses were not offset by the low C input from conifer litter within a brief period (Pérez-Cruzado et al. 2012). The result was in agreement with previous findings (Farley et al. 2004; Laganière et al. 2010). The researches suggested that SC was decreased in coniferous plantations at the decade scale. SC accumulation occurred until the soil reached a new equilibrium between C input and C output.

In the study, SC storage was demonstrated as the largest C pool in the ecosystem throughout the four plantations (Fig. 4). The differences in ecosystem C storage among plantations

were mainly determined by the magnitude of SC pool. There was a significant correlation between C storage in soil and that in ecosystem ($r=0.971$, $P<0.001$). This was consistent with the report of Dixon et al. (1994) regarding the soil pool forming the major part of forest C storage. With the growth of forest trees, there will be more organic matter input to the soil. The ecosystem C storage will increase with the increment of SC storage (Li et al. 2011). Biomass C storage was the second largest C pool in the ecosystem. Winjum and Schroeder (1997) reported that mean biomass C storage was 64 t C/ha in temperate planted forests. With the exception of *Pinus tabuliformis*, the biomass C storage of the other three plantations tested in our study was lower than the mean value (64 t C/ha). This was probably attributed to the negative effect of semiarid climate on tree growth and plant biomass (Silvester and Orchard 1999). However, Li et al. (2003) estimated the mean biomass C storage was 3.4 t C/ha in grassland, and 5.7 t C/ha in agricultural land in China. The biomass C storage of the four plantations in our study was much higher than the above mentioned ecosystems. Consequently, afforestation carried out in this region pronouncedly increased ecosystem C storage.

Table 7 SC storage and sequestration rates of different plantations

Tree species	Plantation (t C/ha)	Abandoned land (t C/ha)	SC sequestration rate (t C/ha/year)
<i>Larix gmelinii</i>	342±26b	395±87	-1.16
<i>Pinus tabuliformis</i>	167±21d	341±96	-9.86
<i>Picea crassifolia</i>	411±24a	263±36	3.08
<i>Populus simonii</i>	283±1c	67±13	8.32

Data represent the mean ± standard deviation. Significant differences among plantations are indicated with lowercase letters ($\alpha=0.05$)

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Contribution of the co-authors Yang Gao: measuring the samples, running data analysis and writing the paper. Jimin Cheng: supervising the work. Zhengrui Ma, Yu Zhao and Jishuai Su: collecting the samples.

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