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# *Pinus nigra* Arn. ssp. *salzmannii* seedling recruitment is affected by stand basal area, shrub cover and climate interactions

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## Abstract

• **Key message** Shrub cover has visible effects on *Pinus nigra* Arn ssp. *salzmannii* seedling emergence, but only in drier years under moderate basal area (25–30 m<sup>2</sup> ha<sup>-1</sup>). In the wetter year, shrub cover favours seedling survival without basal area influence.

• **Context** Shrubs are known to favour tree seedling recruitment in difficult climate environments, but facilitation may prove optimal in intermediate-level rather than harsh conditions, although such an effect remains to be specified.

• **Aims** The main aim of this work was to evaluate the influence of stand basal area (15–20, 25–30 and 35–40 m<sup>2</sup> ha<sup>-1</sup>) × with/without shrub facilitation on seedling recruitment in Spanish black pine (*P. nigra* Arn. ssp. *salzmannii*), which has been struggling to regenerate natural

forests since the late nineteenth century, sometimes jeopardizing stand persistence.

• **Methods** In spring 2012 and 2013, 72 subplots of 25 × 25 cm were established in Cuenca Mountains, central-eastern Spain (four replicates, each counting three subplots × 2 shrub conditions, both under and outside shrub cover, in three different stand basal areas). All plots were randomly distributed at least 500 m apart within each stand basal area. Each subplot comprised 20 seeds previously collected at the experimental forest site. Seedling emergence and early survival were monitored every month in 2012 and 2013.

• **Results** Seedling recruitment was strongly influenced by year. The year 2012 was drier and warmer than 2013. In 2012, seedling emergence was higher under the 25–30-m<sup>2</sup> ha<sup>-1</sup> stand basal area and favoured by shrub cover, whereas in 2013, neither basal area nor shrub cover had significant effects. No seedling survived the summer in 2012. Higher seedling survival was found outside shrub cover at every basal area interval in 2013 with living seedlings until the end of the year.

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**Contribution of the co-authors** ME Lucas-Borja and D Candel-Pérez conceived the research, obtained the field data, run the data analysis and wrote the manuscript. PA Tíscar, T Onkelinx and P Balandier run the data analysis and wrote the manuscript. FA García Morote obtained the field data and wrote the manuscript.

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• **Conclusion** Climate is the main factor controlling seedling emergence and survival. Shrub facilitation has visible effects on seedling emergence, but only in drier years under moderate light (moderate basal area). Facilitation collapses at extreme stress levels. In wetter years, shrub cover does not promote seedling survival and basal area is not an influential factor.

**Keywords** Tree regeneration · Nurse plants · Climate change · Resource limitation · Light · Water · Plant interactions · Facilitation

## 1 Introduction

Under the Convention for the Conservation of European Wildlife and Natural Habitats (Resolution 4/1996), EU directives classified Spanish black pine forests as ‘habitats of European interest’ requiring specific conservation measures, partly due to the lack of basic understanding on the regeneration biology of this long-lived species (Kerr 2000, 2008). Several authors have cited a number of big problems for successful natural regeneration of Spanish black pine: irregular masting conditions, dry summers over a period of at least 3 years, excessive grazing and uncontrolled ploughing (Del Cerro et al. 2009; Lucas-Borja et al. 2010, 2011, 2012; Tiscar and Linares 2014). Moreover, forecasts suggest these pine forests will be soon one of the forest ecosystems hit hard by climate change, and their current distribution could shrink dramatically (López Serrano et al. 2009). In this context, nurse shrubs and adequate overstorey density may facilitate seedling germination, emergence and survival in harsh environments. However, the issue of how combinations of tree canopy and shrub covers affect initial Spanish black pine seedling recruitment has never been investigated.

Plant–plant interactions range from negative (e.g. competition or allelopathy) to positive relationships (e.g. facilitation). Numerous studies consistently demonstrated that shrubs can act as nurse plants favouring plant seedling establishment through facilitative interactions with beneficiary plants in extreme environments worldwide (e.g. Kitzberger et al. 2000; Gómez-Aparicio et al. 2005; Howard et al. 2012). Shrubs are also recognized as important niches for tree regeneration (Callaway 1995; Rodríguez-García et al. 2011). The positive impact of one plant on another has been shown to occur by offering shading that helps improve microclimate (evapotranspiration lessening), increase nutrient availability and increase soil stability (see Callaway 1995 for a review). However, facilitation, competition and the direction of plant–plant interactions frequently change in response to environmental conditions (Kitzberger et al. 2000; Callaway et al. 2002; Maestre and Cortina 2004; Le Bagousse-Pinguet et al. 2014). Callaway and Walker (1997) pointed out that the importance of facilitation and competition varies inversely across a gradient of abiotic

stress, with facilitation dominating under highly stressful conditions. Sthultz et al. (2007) showed that *Pinus edulis* Engelm. and *Fallugia paradoxa* D. Don (a common shrub in the semiarid south-western USA) interactions shifted from facilitation at high-stress to competition at low-stress sites, which is consistent with the stress gradient hypothesis (SGH) posed by Bertness and Callaway (1994). Conversely, Maestre et al. (2005) found that neither positive nor negative plant–plant interactions increased with increasing abiotic stress in arid and semiarid environments. Furthermore, different studies found that facilitation often collapses at the extreme end of stress and physical disturbance gradients, due to a decrease or disappearance of the effect of the nurse plant species (e.g. Michalet et al. 2006, 2014).

The study of shrub and tree canopy cover interactions may help us to understand the population dynamics involved as a step towards guaranteeing forest persistence. According to different studies, the presence and abundance of tree canopy cover affect natural regeneration success during the early phase of the regeneration cycle (Emborg 1998; Page et al. 2001). Changing shelterwood density has a strong influence on understorey light and soil moisture, which may affect seedling performance and thus ultimately regeneration outcome (Agestam et al. 2003; Balandier et al. 2006a). Recent studies in Mediterranean ecosystems investigated influences of canopy cover on the natural regeneration mechanisms of its main species, including *Pinus sylvestris* L. (Barbeito et al. 2009), *Pinus nigra* Arn. (Del Cerro et al. 2009; Tiscar 2007; Lucas-Borja et al. 2011), *Pinus pinea* L. (Adili et al. 2009), *Pinus halepensis* Mill. (Prévosto et al. 2012), and *Pinus pinaster* Ait. (Rodríguez-García et al. 2011). Lucas-Borja et al. (2011) recommend soil preparation combined with moderate or dense basal area intervals (from 25 to 40 m<sup>2</sup> ha<sup>-1</sup>) during the uniform opening of the tree canopy for Spanish black pine (*P. nigra* Arn. ssp. *salzmannii*) regeneration purposes, but the authors did not address the ‘nurse shrub’ effect on seedling performance.

Shrub and tree covers may create favourable microsites in which environmental conditions are ameliorated to improve emergence, survival and early recruitment of conifer seedlings (Germino et al. 2002; Legras et al. 2010; Maher et al. 2005). However Rodríguez-García et al. (2011) argued that shrubs may also have a negative effect on initial seedling growth by competing for resources, but indirect facilitation can occur if the benefit of suppressing competitive species is higher than the direct negative effects (Pagès et al. 2003; Kunstler et al. 2006). Also, abiotic stress conditions may modulate this trend. Most ‘nurse shrub’ studies only considered the effect of shrubs on natural regeneration success in open areas, ignoring the effect of tree canopy cover on the relationship. Consequently the main objective of this study was to analyse the effects of shrub cover in combination with tree canopy cover on Spanish black pine seedling

emergence and survival during the first growing season of two different years. We expected to find that seedling–nurse shrub interactions would vary depending on tree canopy cover, i.e. stronger and positive interactions at more stressful sites (e.g. low tree density) but negative interactions under closed tree canopy conditions resulting from a higher competition for water.

## 2 Material and methods

### 2.1 Study area

This study was led over 2 years (2012 and 2013) in *Serranía de Cuenca*, a mountainous area located in the region of Castilla-La-Mancha (central-eastern Spain). Spanish black pine (*P. nigra* Arn. ssp. *salzmannii*) is naturally distributed in this area between 1000 and 1500 m a.s.l. where it dominates forest stand composition (Del Cerro et al. 2009). Spanish black pine forests in the Cuenca Mountains have traditionally been managed using the shelterwood method, with a shelter phase of 20–25 years and a rotation period of 100–125 years (Tiscar et al. 2011). The regeneration method used, in both mixed and pure even-aged Spanish black pine stands, involves a uniform opening of the canopy without soil preparation. A dense, continuous and homogeneous mature Spanish black pine forest aged about 100–120 years old was selected in the study area at the core of the distribution range for this species (Los Palancares y Agregados, 1200 m a.s.l., 40° 01' 50" N; 1° 59' 10" W). The herbaceous vegetation is mainly dominated by *Eryngium campestre* L., *Geranium sylvaticum* L., *Centaurea paniculata* L. and *Plantago media* L., whereas shrub understory is composed of *Juniperus oxycedrus* L., *Genista scorpius* L., *Amelanchier ovalis* Medik., *Acer campestre* L., *Corylus avellana* L., *Crataegus monogyna* Jacq., *Viburnum lantana* L., *Rubus* sp., *Rosa* sp., and *Prunus spinosa* L. According to Allué (1990), the climate is classified as Mediterranean humid, with a mean annual temperature of 11.9 °C (mean lowest temperatures of the coldest month at –0.5 °C and mean highest temperatures of the hottest month at 30.5 °C) and mean annual precipitation of 595 mm (99 mm in summer). Calcareous soils dominate the study area, and the soil found in the experimental site is classified as *leptosol* according to the Soil Atlas of Europe (2005). During the last century, the forest stand was not affected by natural disturbances such as windstorms, pests and diseases, or forest fires.

### 2.2 Experimental design

A sowing experiment was conducted over 2 years (2012 and 2013) in 'Los Palancares y Agregados' forest. Based on previous knowledge on stand-structural characteristics (Lucas-

Borja et al. 2011; Candel-Pérez et al. 2012), we selected three stand basal area intervals, the basal area being used as a proxy for tree canopy cover, i.e. BA1 ~15–20 m<sup>2</sup> ha<sup>-1</sup>, low cover; BA2 ~25–30 m<sup>2</sup> ha<sup>-1</sup>, moderate cover; and BA3 ~35–40 m<sup>2</sup> ha<sup>-1</sup>, dense cover (Table 1). Four plots were randomly distributed within each stand basal area interval at least 500 m apart and were thus considered independent and to form the replicates of the experiment. Each plot was occupied by the same shrub species, namely *J. oxycedrus* and *G. scorpius*. In spring 2012 and 2013, six subplots were established in each plot, three under the shrub cover, i.e. inside the space delimited by the vertical projection of the shrub cover, and three outside the shrub cover, i.e. at a minimum of 5 m away from the space delimited by the vertical projection of the shrub cover, giving a total of 72 subplots (3 levels of stand basal area × 4 plots × 3 subplots × 2 shrub conditions). Each subplot was composed of 25 × 25 cm quadrats with a 1-cm<sup>2</sup> mesh size and protected with a wire cage to exclude seed predation from birds and rodents. Twenty seeds collected in local Spanish black pine stands were sown 1 cm deep per sowing point in April 2012 and again in April 2013 (the same subplot was sown in both years). In-lab germination tests to evaluate the potential germination rates of the seeds used in 2012 and 2013 found no differences, with germination rates reaching about 85 % (see Lucas-Borja et al. 2011 for details on the controlled seed germination procedure).

### 2.3 Measurements

Daily air temperature and precipitation were recorded during the entire length of the study at a meteorological station (model METEODATA 1256C) located in the same forest near the plots. Annual water budget was estimated by the standardized precipitation-evapotranspiration index (SPEI) (Vicente-Serrano et al. 2010) based on the sum of monthly differences between precipitation (P) and potential evapotranspiration (PET). Soil temperature and moisture were measured at 10-cm depth every 20 min from March to September in both 2012 and 2013 using six (3 stand basal area intervals × 2 shrub conditions) soil moisture and temperature sensors (HOBO<sup>®</sup> U12). Seedling emergence (early appearance of the cotyledons) and seedling survival were monitored regularly in both 2012 and 2013. Emergence and survival were surveyed every 2 weeks during late spring and monthly over the summer season.

### 2.4 Statistical analysis

The effects of stand basal area, shrub condition and year of sowing on seedling emergence and survival rates were analysed using linear mixed-effects models. Stand basal area, shrub condition and year of sowing were considered fixed effects. Experimental design was considered as nested, with

**Table 1** Forest stand structure found at each stand basal area (mean  $\pm$  standard error around the mean, SEM)

Number of plots	Basal area (m <sup>2</sup> ha <sup>-1</sup> )	DBH (cm)	Mean tree height (m)	Number of trees (N ha <sup>-1</sup> )	Shrub cover (%)
4	Dense 35–40	26.0 (2.1)	15.1 (1.1)	646.4 (19.1)	24.1 (3.8)
4	Medium 25–30	30.0 (3.3)	15.0 (0.9)	322.1 (21.1)	25.9 (4.3)
4	Low 15–20	32.0 (3.1)	14.9 (0.9)	183.0 (18.1)	27.3 (4.7)

the plots as the replicates nesting the subplots and the shrub condition. The plots and replicates were considered random terms in the model. Analysis of survival data was limited to the 2013 dataset, because no seedlings survived the summer of 2012. Seedling emergence and survival differences for each basal area interval and sowing position in 2012 and 2013 were evaluated according to the Tukey HSD post hoc test. All statistical analyses were performed using Statgraphics Centurion XV<sup>®</sup>.

### 3 Results

#### 3.1 Precipitation, air and soil temperatures and soil moisture

Mean monthly temperature presented similar values over the 2-year period studied (Fig. 1), with the exception of a warmer summer during the first year (2012). The study area receives an average annual precipitation of 784.6 mm (for the 1948–2005 period). The wettest month is December, with a mean monthly precipitation of 85.7 mm, and the driest month is July, with a mean monthly precipitation of 21.9 mm. The year 2012 was a drier year than 2013 (420 vs. 850 mm in 2012 and 2013, respectively). Precipitation showed a dramatic negative deviation from the mean values throughout most of 2012 (Fig. 2), whereas the opposite situation was recorded in 2013 where precipitation was higher-than-average in January and March and summer (from July to October). The SPEI showed a stark water deficit in the winter months of 2012 (negative values), whereas months with high water availability were observed in the year 2013 (Fig. 3).

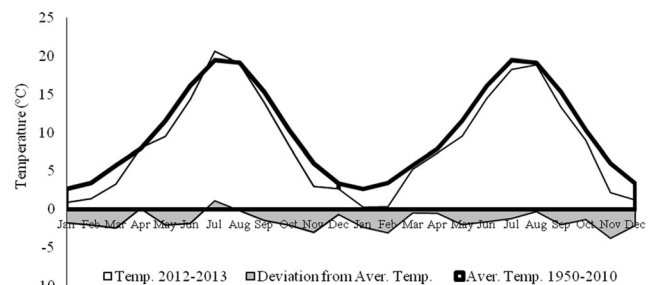
In terms of microclimatic conditions generated by tree canopy cover and shrub cover, in general, soil temperatures were lower (°C) outside shrub cover, both in 2012 and 2013 (Table 2). They also decreased from low to dense basal area. Soil moisture (%) increased with increasing basal area, in particular in 2012. Finally, soil moisture values were lower outside the shrub cover. The minimum value was recorded in 2012 for subplots with low basal area and outside the shrub cover (6.0 %; Table 2).

#### 3.2 Seedling emergence

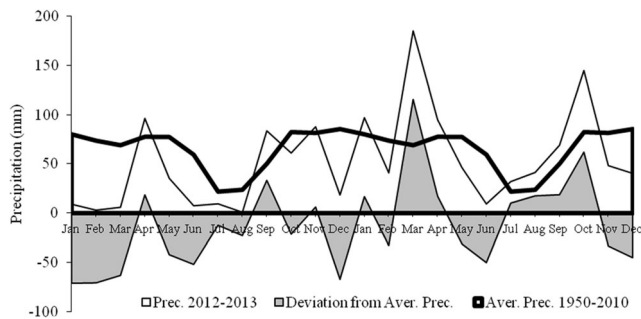
The results of the mixed model analysis for the effects of stand basal area, shrub condition and year of sowing (2012 and 2013) showed that only year of sowing significantly influenced seedling emergence rate (chi-square ratio = 44.77,  $P < 0.001$ ), associated with a significant stand basal area  $\times$  year interaction (chi-square ratio = 4.16,  $P = 0.032$ ) and a barely significant shrub cover  $\times$  year interaction (chi-square ratio = 3.65,  $P = 0.046$ ). In 2012, seedling emergence showed the best results under the moderate basal area interval (BA2 ~25–30 m<sup>2</sup> ha<sup>-1</sup>) and was favoured by shrub cover (emergence rate of 0.4, Fig. 4), whereas seedling emergence rate was very low at both the lowest and highest basal area intervals, only exceeding 10 % in plots protected by a combination of high basal area and shrub cover. In 2013, there was no significant stand basal area effect on seedling emergence nor a shrub cover effect (Fig. 4).

#### 3.3 Seedling survival

In 2012, all emerged seedlings died in all shrub conditions and all basal area intervals. Consequently, only 2013 data were analysed. The results of the mixed model analysis for the effects of stand basal area and shrub condition showed that only shrub condition significantly influenced seedling survival rate (chi-square ratio = 27.72,  $P < 0.010$ ). Neither basal area nor the interaction between stand basal area and shrub condition was significant (chi-square ratio = 2.44,  $P = 0.095$ ; chi-



**Fig. 1** Range of mean monthly temperature (°C), normal monthly temperature (1950–2010), and deviation of mean monthly temperature from historical average temperature, for the period January 2012 to December 2013



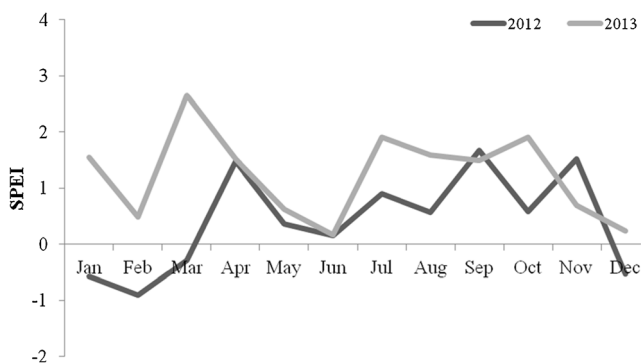
**Fig. 2** Range of monthly precipitation (mm), historical average precipitation (1950–2010) and deviation of monthly precipitation from historical average precipitation, for the period January 2012 to December 2013

square ratio = 0.11,  $P = 0.898$ , respectively). Results showed that seedling survival was higher outside shrub cover at every basal area interval (Fig. 5).

#### 4 Discussion

Seedling emergence rate was strongly influenced by the considered year and the two-way interactions with shrub condition and basal area interval, likely related to the contrasted climatic features of the 2 years studied; 2012 was particularly dry and a little warmer than usual, whereas 2013 was wetter than usual. In 2012, seedling emergence was higher under the medium 25–30 m<sup>2</sup> ha<sup>-1</sup> stand basal area and was favoured by being inside shrub cover, whereas in 2013, there were no significant effects of basal area or shrub condition on emergence.

Light on the forest floor is a direct consequence of canopy structure and stand/shrub density (Catovsky and Bazzaz 2000; Balandier et al. 2006b; Barbier et al. 2008); a high light level may increase soil temperature and climate demand, thus decreasing accordingly soil moisture by increasing soil evaporation, as recorded in this particularly difficult conditions under the lowest tree canopy cover (Table 2). Soil temperature and



**Fig. 3** Standardized Precipitation-Evapotranspiration Index (SPEI) for 2012 and 2013, estimated from a standardized sum of monthly differences between precipitation (P) and potential evapotranspiration (PET)

humidity are likely to be the two main drivers of germination and seedling emergence under field conditions (Lee et al. 2004; Castro et al. 2005; Del Cerro et al. 2009). There is broad evidence (Holmgren et al. 1997; Del Cerro et al. 2009) that soil and air temperatures are lower under tree canopy, wind velocity is reduced and air humidity is higher than in open areas, thus reducing seedling transpiration demand and soil evaporation. Furthermore, field experiments conducted on the forest floor under pine stands indicated that germination is soil–water content-dependent (Lee et al. 2004).

In the Cuenca mountains conditions, the micrometeorological and light conditions found under moderate basal area interval and inside shrub cover conditions would have generated a better environment for Spanish black pine seedling emergence in the drier and warmer year 2012 (Granda et al. 2012). On one hand, higher basal area was unfavourable, probably because the competition for light, or other resources, or through allelopathic processes sometimes mentioned, is too high. Higher light conditions (low tree canopy cover and outside shrub cover) are less favourable as they do not generate adequate conditions for emergence in drier and warmer years. Therefore, it seems that for the conditions of the Cuenca Mountains, the considered shrub species (*J. oxycedrus* and *G. scorpius*), and the considered dryness in 2012, interactions between pine seedlings, shrubs and adult trees moved from competition at high and low stand basal area to facilitation at moderate basal area. The positive effect of shrub cover on seedling emergence may have been driven by increased moisture resulting from the effects of shrub cover on soil temperature and relative air humidity (Holmgren et al. 1997). These results support the Bertness and Callaway (1994) theoretical model predicting a switch from competition to facilitation with increasing harshness of the environment and also the ‘collapse of facilitation’ model (Michalet et al. 2006) for the harshest sites. Further evidence that facilitation collapses under extreme stress levels was the zero survival in 2012 both with and without shrub cover protection, as also found by Kitzberger et al. (2000). However, as our results came from only 2 years and a single forest, they obviously need to be replicated in other places and years to be generalized.

In contrast, neither basal area nor shrub condition had significant effects on seedling emergence under the more favourable climate conditions and higher soil moisture recorded in 2013. Drought alleviation and higher water availability during 2013 allowed Spanish black pine to emerge without shrub and canopy cover protection (Castro et al. 2005).

Seedling survival rate was null in the drier and warmer year (2012). As different authors argued, seedling survival is one of the most critical stages in a plant’s life history, and is often reduced by drought and high temperatures in the Mediterranean basin (Madrigal et al. 2004; Padilla and Pugnare 2007), where water deficit in the summer appears to be the dominant cause of seedling mortality (Valladares 2004).

**Table 2** Soil temperature and moisture (mean  $\pm$  standard error around the mean, SEM) measured at 10 cm depth from March to September in 2012 and 2013 tabulated by stand basal area interval and shrub conditions

Stand basal area interval <sup>a</sup>	2012				2013			
	Soil temperature (°C)		Soil moisture (%)		Soil temperature (°C)		Soil moisture (%)	
	ISC	OSC	ISC	OSC	ISC	OSC	ISC	OSC
Dense	7.1 (0.5)	7.3 (0.4)	12.1 (0.4)	10.9 (0.3)	5.0 (0.5)	6.2 (0.3)	12.3 (0.5)	11.3 (0.4)
Medium	8.2 (0.4)	10.0 (0.3)	11.0 (0.3)	6.2 (0.2)	6.1 (0.5)	8.1 (0.2)	12.2 (0.3)	8.7 (0.3)
Low	8.0 (0.3)	10.5 (0.3)	10.0 (0.2)	6.0 (0.5)	7.9 (0.4)	8.5 (0.4)	12.1 (0.4)	8.5 (0.5)

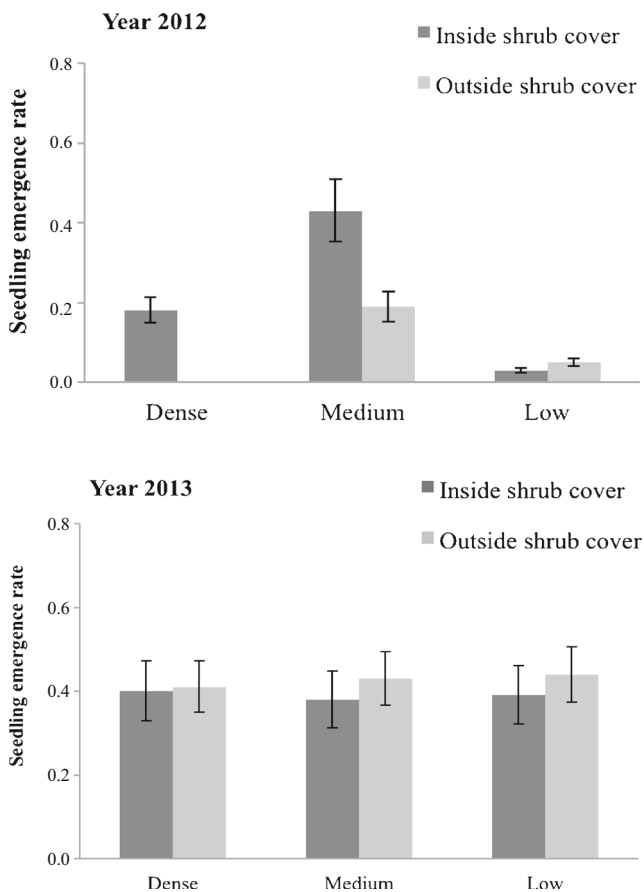
ISC inside shrub cover, OSC outside shrub cover)

<sup>a</sup> Basal area intervals (low (BA1 ~15–20 m<sup>2</sup> ha<sup>-1</sup>), medium (BA2 ~25–30 m<sup>2</sup> ha<sup>-1</sup>) and dense (BA3 ~35–40 m<sup>2</sup> ha<sup>-1</sup>))

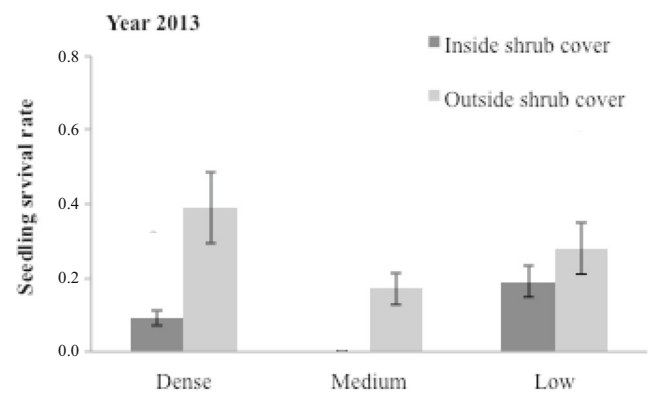
Drought pressure is the most important factor controlling regeneration density of *P. nigra* ssp. *pallasiana* (Fyllas et al. 2008), especially in the first growing season (Del Cerro et al. 2009). Note that the more favourable climatic conditions in 2013 allowed seedlings to survive until the end of the experiment. We observed that survival was better achieved outside shrub cover under the more favourable conditions of 2013 without basal area influence. This fact suggests a light effect

(seedlings receive higher light outside shrub cover), as already reported by Adili et al. (2013) and also a shrub competition effect with decreasing harshness (Bertness and Callaway 1994) without adult trees influence.

In fact, seedling establishment and early growth may be prevented by successive filters, including variability in seed production, light and understory vegetation. Water availability is thus the primary limiting factor, but when it becomes non-limiting in case of favourable climate conditions (2013), then light becomes the next biggest limiting factor. Indeed, solar radiation appears to be an important environmental factor defining the regeneration niche of pine species: higher solar radiation implies higher seedling survival and higher initial seedling growth (Rodríguez-García et al. 2011). Simard et al. (1998) stated that increased light favours a longer taproot and longer shoots in seedlings, enabling them to penetrate the soil and survive better. When seedling roots extend deeper into soil, they uptake sufficient water to increase the chances of seedling survival. Here, the absence of soil water deficit during 2013, as demonstrated in the calculated SPEI, may have helped the Spanish black pine seedlings to survive until the arrival of autumn rains. Finally, basal area was a not significant factor in relation to seedling survival in 2013,



**Fig. 4** Seedling emergence rate at different sowing positions (inside or outside shrub cover) and different basal area intervals (low (BA1 ~15–20 m<sup>2</sup> ha<sup>-1</sup>), medium (BA2 ~25–30 m<sup>2</sup> ha<sup>-1</sup>) and dense (BA3 ~35–40 m<sup>2</sup> ha<sup>-1</sup>)) in 2012 and 2013. Error bars are the Tukey HSD at  $P < 0.05$



**Fig. 5** Seedling survival rate at different sowing positions (inside or outside shrub cover) and different basal area intervals (low (BA1 ~15–20 m<sup>2</sup> ha<sup>-1</sup>), medium (BA2 ~25–30 m<sup>2</sup> ha<sup>-1</sup>) and dense (BA3 ~35–40 m<sup>2</sup> ha<sup>-1</sup>)) in 2013. Error bars are the Tukey HSD at  $P < 0.05$

suggesting that the strength of tree–seedling interactions is significantly lower in wet years (Gómez-Aparicio et al. 2004).

## 5 Conclusion

As shown by Lucas-Borja et al. (2012), the natural regeneration of Spanish black pine has to go through conflicting situations between conditions suitable for seedling emergence (medium basal area interval and inside shrub cover) and conditions suitable for seedling survival (outside shrub cover without basal area influence), this effect being modulated by climate of the given year. Shrub facilitation was only confirmed for seedling emergence in the drier year under moderate light (medium basal area), whereas shrub cover does not promote seedling survival and basal area is not an influential factor in the wetter year. Indeed the findings reported here point to bleak prospects for natural Spanish black pine regeneration success in the driest years like 2012, with no survival at the end of the growing season, whatever the basal area or shrub cover conditions. In addition we have to highlight that the climate conditions of 2012 could become more frequent with the ongoing climate changes. Therefore, forest managers should indeed take advantage of the wettest years to promote black pine regeneration by clearing shrubs after seedling emergence to increase light availability and avoid shrub competition.

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## Compliance with ethical standards

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