

Restoration of burned and post-fire logged *Austrocedrus chilensis* stands in Patagonia: effects of competition and environmental conditions on seedling survival and growth

M. Florencia Urretavizcaya^{A,B,D} and Guillermo E. Defossé^{A,C}

^AConsejo Nacional de Investigaciones Científicas y Tecnológicas (CONICET),
Centro de Investigación y Extensión Forestal Andino Patagónico (CIEFAP),
CC 14 Esquel (9200) Chubut, Argentina.

^BMinisterio de la Producción, Provincia de Chubut, Argentina.

^CUniversidad Nacional de la Patagonia San Juan Bosco, Facultad de Ingeniería, Esquel,
Chubut, Argentina.

^DCorresponding author. Email: mfurretavizcaya@ciefap.org.ar

Abstract. In Andean Patagonia, Argentina, severe wildfires produce high mortality in *Austrocedrus chilensis* forests. Owing to its high timber quality, *A. chilensis* trees are generally salvage logged right after wildfires. Post-logged areas result in open, denuded stands with low herbaceous or shrub cover, which precludes natural *A. chilensis* regeneration. In severely burned *A. chilensis* stands, we determined how different site conditions (salvage logging, SL, and without logging, WL), and the combination of different methods of control of above- and belowground competition and micro-environmental factors (incident radiation, soil temperature and soil moisture) affected survival, growth and hydric status of planted *A* seedlings. Two growing seasons after plantation, seedling survival was lower than 10% at SL sites, whereas it was near 90% at WL sites. Four seasons after establishment, and regardless of competition treatments, no seedlings survived at SL sites, whereas over 75% survived at WL sites. Radiation attenuation by canopy of WL sites benefitted *A. chilensis* seedling survival, and no additional control of early successional herbaceous species should be necessary to attain high seedling survival 4 years after restoration. Selective logging, by leaving ~50% of burned snags, and active restoration practices, may help reconcile economic needs of society and ecological requirements of *A. chilensis* for recovering former structure and functions.

Additional keywords: Andean cypress, facilitation, seedling performance, temperate forests.

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Introduction

Wildfires have shaped the structure and functioning of most terrestrial ecosystems (Wright and Bailey 1982; Crisp *et al.* 2011). Every ecosystem, however, has its own fire history, specifically defined as its *fire regime*. The fire regime is composed of different elements, identified as (1) *fire type and intensity*, (2) *size* of typical significant fires and (3) *frequency or return intervals* for specific land units (Heinselman 1981). In forest ecosystems, long fire return intervals may produce high severity fires, whereas short fire return intervals may lead to low severity fires (Pyne 1984; Agee 1993; Beschta *et al.* 2004). High severity wildfires usually produce major changes in community and population structure, modifying resources availability and altering the physical environment (Agee 1993). These changes favour the establishment of early successional species, also creating suitable conditions for invasion by new species. High severity fires may also have long lasting effects on low fire-tolerant forests, especially in those whose most prevalent species reproduce strictly by seed, which are killed during,

or because of, the fire disturbance. Moreover, if remaining burned snags are removed from stands after high severity fires, additional changes are created in the post-fire denuded environment, increasing solar radiation, and modifying soil temperature and wind speed (McIver and Starr 2000; Beschta *et al.* 2004). In other cases, in which seeds of forest species are slightly or not damaged by fire, the rapid growth of herbaceous species right after the fire disturbance may constrain their germination and establishment, by out-competing for water and nutrients (Rose and Rosner 2005). The combination of these factors generally delay the natural process of forest recovery and, in extreme situations, preclude burned forests from recovering to pre-disturbance conditions. Under these situations, the fire-disturbed stand needs active restoration practices to return to its normal successional path, by sowing or planting former species. However, in order to guarantee the establishment of the target species, restoration practices sometimes require additional microsite manipulation to avoid or diminish any deleterious biotic or micro-environmental effects on planted

seedlings (Whisenant 1999). These effects are mainly caused by competition for soil water or nutrients by early successional herbaceous vegetation, excessive soil heating by higher incoming radiation reaching denuded soils, or by the lack of nurse plants (generally shrubs or half-shrubs) that protect seedlings from excessive temperature fluctuations (Padilla and Pugnaire 2006; Urretavizcaya *et al.* 2017). Forest-productivity research has identified interactions between competing vegetation and logging debris left after forest harvesting. In general, retention of logging debris in temperate zones inhibits development of herbaceous and sometimes woody species. Retention of logging debris has also been associated with mulching effects. These effects imply conservation of soil water near the soil surface by reducing evaporation and topsoil temperatures (Harrington *et al.* 2013).

In restoration projects, controlling competing species may be sometimes necessary for diminishing their deleterious effects on planted seedlings (Jobidon *et al.* 1998; Löf *et al.* 1998; Dinger and Rose 2009; Maguire *et al.* 2009). This control may involve using herbicides, manual removal of competing vegetation or the application of mulch (Löf *et al.* 1998; Harper *et al.* 2005; Jylhä and Hytönen 2006). Removing competing vegetation by chemical or mechanical methods reduces soil cover and modifies some topsoil physical conditions, increasing incoming radiation and soil-surface temperature, and also influencing soil-water content (Defossé and Robberecht 1996; Harrington *et al.* 2013). For some forest species at seedling stages, high topsoil temperatures at stem-collar level may negatively affect their survival (Kolb and Robberrecht 1996). However, the use of mulch around seedlings may increase survival by reducing temperature variations, conserving soil moisture and diminishing plant evapotranspiration (Flint and Childs 1987). Although mulching is widely used in horticulture, it has scarcely been used in restoration projects aimed at recovering forest species under field conditions (Blanco-García and Lindig-Cisneros 2005).

The forests of *Austrocedrus chilensis* (D. Don) Pic. Ser et Bizarri (commonly known as cypress), are part of the Andean–Patagonian temperate forests that run along the Cordillera de los Andes in southern Chile and Argentina (Cabrera 1976). In Argentina, *A. chilensis* is located in a narrow belt that goes from 37°08' to 43°44'S in the eastern slope of the Andes (Pastorino *et al.* 2006). *Austrocedrus chilensis* forms either pure or mixed stands. Pure stands occupy ~95 500 ha, whereas those in which *A. chilensis* grows intermixed with *Nothofagus dombeyi* Mirb. Oerst. (locally named as coihue) cover ~32 800 ha. National fire statistics show that during the last 20 years, an average of 3000 ha of pure or mixed *A. chilensis* stands have been annually affected by wildfires (Secretaría de Ambiente y Desarrollo Sustentable de la Nación 2013; Mohr Bell 2015).

After wildfires, the main factors that influence recovery of affected forests are intensity and severity of each particular fire, size or area covered, and environmental conditions prevailing during and after the fire event (Connell and Slatyer 1977; Veblen *et al.* 1999). For *A. chilensis* forests, the lack of a soil-seed bank after severe wildfires is the initial limiting factor for natural stand recovery (Urretavizcaya and Defossé 2004). Likewise, severe wildfires produce very high *A. chilensis* tree mortality because its very thin bark is rapidly affected by the fire and, unlike other Patagonian species, it does not re-sprout

after fires (Gallo *et al.* 2004). These conditions mean that active-restoration practices, such as seedling planting or seed sowing, are often necessary for restoring *A. chilensis* stands affected by severe fires (Urretavizcaya *et al.* 2012).

Most burned *A. chilensis* forests are logged right after fire events because of their high timber quality and market demands (Perdomo *et al.* 2007). Once burned-log extraction is finished, logged areas result in open, denuded stands with very low herbaceous or shrub cover, similar to what happens in other postfire-logged conifer forests elsewhere (McIver and Starr 2000; Beschta *et al.* 2004). However, for *A. chilensis* forests, there are no studies that analyse these conditions. The new macro- and micro-environmental conditions created in these post-burning logged areas may preclude survival and establishment of planted *A. chilensis* seedlings, sometimes making any restoration plan unsuccessful. Similar to what was stated in Castro *et al.* (2011) and Harrington *et al.* (2013, 2018), we hypothesise that, for *A. chilensis* forests, the presence of burned logs act as structural nurses that, without competing with established vegetation, help reduce incoming radiation and maintain topsoil moisture, crucial for seedling survival during the summer drought. In the case of salvage logging, mulch attenuates the desiccant effects of high levels of direct solar radiation and help reduce evaporative loss by diminishing high topsoil temperatures that are usual in this region during the summer. We also postulate that mulch, as well as a reduction in nearby vegetation competition, has the potential to benefit *A. chilensis* seedling performance by increasing resource availability and modifying microclimate conditions.

The objective of this study was to determine how different site conditions (salvage logging, SL, and without logging, WL), microsite interventions (different methods of control of above- and belowground competition), and micro-environmental factors (incident radiation, soil temperature and moisture) affect survival, growth and hydric status of *A. chilensis* seedlings planted in severely burned stands. The outcomes of this study not only cast light on the effects of these factors on *A. chilensis* seedling establishment, but also help to design appropriate protocols for successfully restoring fire-damaged *A. chilensis* forests. Additionally, this study provides some guidelines that should be followed for sustainable managing post-fire *A. chilensis* stands.

Materials and methods

Study site

The study area was located near the town of Corcovado (43°32'S, 71°33'W), in Chubut province, Patagonia, Argentina (Fig. 1). Its climate is “mediterranean-temperate”, with cold and rainy winters and dry and mild-to-hot summers and early autumn. Mean annual precipitation is 800 mm, 75% of which falls during the winter months. Average annual temperature is 9.3°C. Soils of the region derive from volcanic ashes (Etchevehere 1972) and correspond to Andisols in the study area. These are rich in nutrients (except phosphorous), and have high soil-water retention (Colmet Dâage *et al.* 1993). This area represents the southernmost location of pure stands of *A. chilensis* in the Patagonian region of Argentina (Dezzotti and Sancholuz 1991). At the end of February 2008 (summer in the

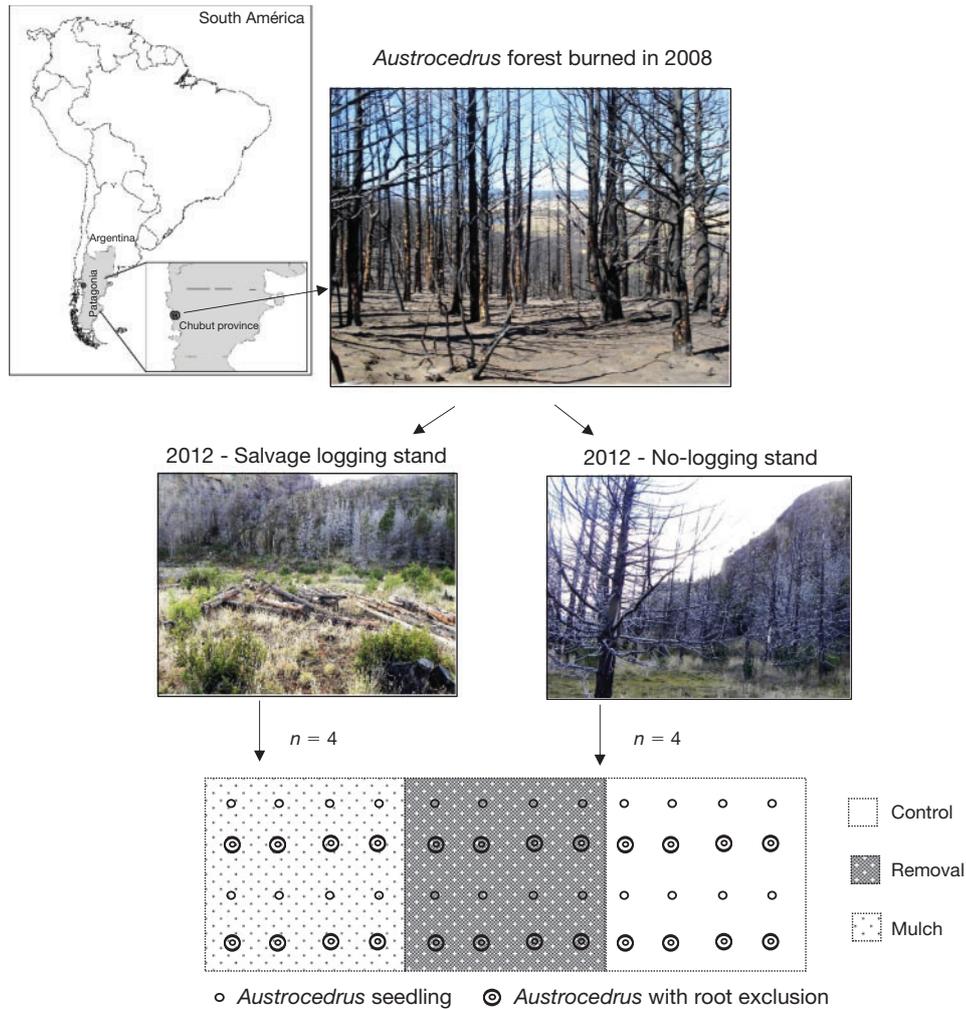


Fig. 1. Geographic location of the study site in Patagonia, Argentina (top left). Images of the study site right after the burn and after setting the logged treatments (top right and centre). Block diagram of *Austrocedrus* seedling plantation comprising aerial (Control, Mulch and Removal) and root competition (root exclusion and root control) treatments in burned stands in Patagonia (bottom).

southern hemisphere), a wildfire categorised as extreme behaviour and high severity ravaged 80 ha within this area. The categorisation of this fire was based on regional observations and by applying the classification proposed by Mutch and Swetnam (1995). Under this classification, the studied fire falls in the category 5: very high severity; most sub-canopy trees killed and >50% of canopy trees killed (Fig. 1).

Experimental design and treatments

In an area representative of the burned forest, we selected two adjacent and similarly burned *A. chilensis* stands (Fig. 1). One of them (~4 ha) was salvage cut in 2010 (i.e. all remaining *A. chilensis* snags removed), whereas the other (2 ha) conserved all the standing burned trees. Both stands presented similar aspect and 0 to 5% slope. In August 2012 (winter in the southern hemisphere), a split-split plot experiment was installed. Site condition or logging was the main-effect plot and had two levels

(SL and undisturbed or WL). Aboveground competition was the second-effect plot and had three levels (aboveground vegetation manually removed (RE), mulch application (MU) and no aboveground removal, or control (CO)). Root competition was the third-effect plot and had two levels (root exclusion (EX), and root control or without exclusion (RC)).

The experiment was installed in eight plantation blocks of 8 × 24 m (4 in SL and 4 in WL conditions) (Fig. 1). In every block, 48 *A. chilensis* seedlings were planted at a distance of 2 × 2 m each. At the time of planting, seedlings were 3 years old and were produced at Centro de Investigación y Extensión Forestal Andino Patagónico nursery (Esquel, Chubut, Argentina) from seeds of the same genetic zone of the study site (Pastorino 2015). Each seedling was manually planted in similarly shovel-made holes. Before planting, a sample composed of 20 randomly chosen seedlings were tested to determine their morphological attributes. These attributes (mean ± standard error) were: seedling height (21 ± 0.5 cm); collar diameter (4.2 ± 0.1 mm), aerial



Fig. 2. Images of the study site in Patagonia showing the unlogged stand, in which aerial competition treatments were established. Mulch application is shown on the left, manual removal in the centre, and undisturbed control in the right photograph.

dry weight (3.1 ± 0.2 g), root dry weight (2.5 ± 0.2 g), aerial to root dry weight ratio (1.3) and sturdiness index (50) (Hasse 2007). These values are within the range of customary morphological attributes used for commercial *A. chilensis* seedlings. At planting, plastic root-exclusion tubes 25 cm in diameter and 25 cm long (Cook and Ratcliff 1984; Defossé 1995) were installed in half of the planted seedlings to isolate their root system from those of neighbouring herbaceous species. Each block was divided into three plots (sample units) containing 16 seedlings each. The three aerial competition treatments (RE, MU and CO) were randomly assigned to one of these plots (Fig. 1). In total, 456 *A. chilensis* seedlings were used in the experiment (228 per stand and 48 per block). Right after plantation, all seedlings were identified and their specific height and collar diameter measured for determining further survival and growth. Each seedling was protected by a metallic mesh for avoiding foraging by the exotic European hare (*Lepus europaeus* Pallas).

The mulch came from chipped *A. chilensis* branches and slash residues remaining after the wildfire. Mulch cover was 3–5-cm high in a 40-cm radius around treated seedlings. Aerial vegetation was manually removed with a hoe and a rake from each removal treatment in a 40-cm radius around each seedling, trying not to disturb the topsoil around it (Fig. 2).

Characterisation of the vegetation of the study site

At 3 and 6 months after setting plantation (November 2012, spring in the southern hemisphere, and February 2013, summer in the southern hemisphere), early successional species grown in both SL and WL stands were cut, taken to the laboratory, identified and their dry weight determined. The sampling procedure involved collection of all biomass located within eight 0.5-m² plots randomly distributed within Control plots of SL ($n = 4$) and WL blocks ($n = 4$).

Environmental variables measured

Photosynthetically active radiation (PAR), and soil temperature (ST) and moisture (SM) were measured three times (mid-spring, 11 November 2012; mid-summer, 6 February 2013; and late summer, 22 March 2013) during the first growing season after the start of the experiment. These measurements were performed to determine if there were significant differences among sites (SL and WL) and aerial competition (RE, MU and CO) treatments. For measuring PAR, a Cavadevices Ceptometer

(Cavadevices.com, Buenos Aires, Argentina) with Photosynthetic Photon Flux method was used. Topsoil temperature was measured with an infrared laser thermometer Cole-Parmer Model 39650-02 (Instrument Company, Chicago, IL, USA) and at 10-cm soil depth with a Digi-Sense DualLogR thermometer and thermocouples Model 91100-50 (Cole-Parmer Instrument Company Vernon Hills, IL, USA). Soil samples from 0 to 10 cm (2 per plot in each block, 48 per date) and 20 to 30 cm (1 per plot in each block, 24 per date) were taken and their water content determined gravimetrically. At the beginning of the second growing season (October 2013), automatic data loggers Decagon Em5b (Decagon Devices, Pullman, USA) with moisture sensors ECH2O (Decagon Devices, Pullman, USA) at 10-cm soil depth were installed in randomly selected SL and WL blocks and each aerial competition treatment. These measurements allowed continuous recording of soil moisture data to the end of the growing season (March 2014).

Seedling survival, growth and hydric conditions measurements

We recorded seedling survival and height growth at the end of the growing season in 2013, 2014 and 2016. In April 2014 and 2016, seedling-collar diameter was also measured. Survival, height and collar-diameter measurements were performed in all seedlings. In mid-summer (February 2013), the hydric status of seedlings was determined both at predawn and midday at both SL and WL sites, and also in aerial (RE, MU and CO) and belowground (EX and RC) competition treatments. For these measurements, we used a Scholander pressure chamber (BioControl Model 4, Buenos Aires, Argentina). The measuring technique involved cutting of a small portion of a top *A. chilensis* seedling branch, using 24 seedlings at predawn and 24 at midday. Two blocks were randomly selected for each WL and SL site treatment and 4 seedlings were sampled per each aerial (RE, MU and CO) treatment, 2 in EX and 2 in the RC treatments.

Analyses

Vegetation biomass was analysed by analysis of variance (ANOVA) of one factor: site (WL and SL), and independently for each measuring date. PAR, ST and SM determined during the first growing season, were analysed by ANOVA with two factors: site (WL and SL) and aerial competition (RE, MU and CO) treatments, and independently for each measuring date.

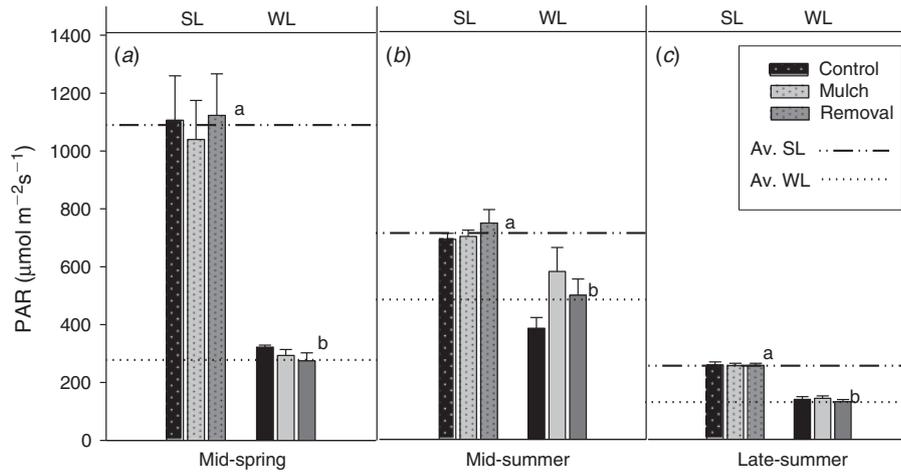


Fig. 3. Photosynthetically active radiation (PAR, mean \pm 1 s.e.) per logging site (SL, salvage logging; WL, without logging) and aerial competition treatments from mid-spring to late summer of the first growing season after *Austrocedrus* seedling planting. Av., average per logging treatment. Different letters above each bar graph indicate statistical significant differences at $P < 0.05$.

Soil moisture, which started to be continuously measured from the beginning of the second growing season, was analysed through mixed lineal models. Soil-moisture values were averaged at monthly intervals for each treatment, and significant differences among treatments determined for each month during the growing season.

Austrocedrus chilensis seedling survival was evaluated as a function of the first (SL and WL), second (RE, MU and CO) and third (EX and RC) effect treatments. The analyses involved repeated measures through lineal mixed models by considering the three effects as fixed, and blocks and time as random factors. These analyses were performed after the first summer, second winter and second summer (April and October 2013 and April 2014 respectively). Survival at the fourth summer (April 2016) was only measured in aerial (RE, MU, CO) and root (EX and RC) competition in WL treatment, because no survival was registered at SL plots and blocks. In these analyses, P -values with Bonferroni correction and a $P \leq 0.05$ were used. When significant differences were found, mean comparison was made by using Fisher’s least significant difference. Height growth after the first growing season (2012–2013) was analysed with the same models used for survival. After that date, the same models were used but only for one site (WL), because most seedlings at SL plots had already died. Height and collar-diameter growth was analysed for the periods 2012–2014 and 2014–2016 and then for the entire study period (2012–2016). Seedling hydric status was analysed by ANOVA with three factors: site (SL and WL), aerial (RE, MU and CO) and root (EX and RC) competition treatments. For all analyses, we used INFOSAT Statistical Package (Facultad de Ciencias Agropecuarias, Universidad Nacional de Córdoba, Córdoba, Argentina, www.infostat.com.ar, accessed 7 February 2019) (Balzarini *et al.* 2008; Di Rienzo *et al.* 2017) and their interface with the R Foundation for Statistical Computing Platform (R version 3.4.0 The R-CoreTeam 2017, <https://www.r-project.org/>, accessed 7 February 2019). Unless stated otherwise, the probability level was set at $P < 0.05$.

Results

Vegetation

At the beginning of the experiment, herbaceous plant cover was similar at all treatment plots and blocks, and the predominant species were *Holcus lanatus*, *Acaena ovalifolia*, *Carduus thoermeri*, *Rumex acetosella* and other grasses. Total herbaceous (grasses and herbs) aboveground biomass was not significantly different at both SL and WL plots. Three and six months after plantation (4.5 and 5 years after the fire), the biomass averaged 272 ± 38 and 414 ± 25 g m⁻² at SL plots and 395 ± 44 and 431 ± 34 g m⁻² at WL plots.

Environmental variables

Photosynthetically active radiation

Photosynthetically active radiation was significantly higher in SL treatments as compared with WL treatments, but was not significantly different among aerial (RE, MU and CO) treatments. Within sites (SL and WL), the higher PAR differences were registered during mid-spring 2012. Salvage logging showed PAR values above $1000 \mu\text{mol m}^{-2} \text{s}^{-1}$, almost four times the PAR registered ($290 \mu\text{mol m}^{-2} \text{s}^{-1}$) in WL treatments. Photosynthetically active radiation differences started to decline towards mid- and late summer (Fig. 3).

Soil temperature

Topsoil temperature in mid-spring and mid-summer of the first growing season was not significantly different between sites (SL and WL), but it was significantly different among competition treatments (Table 1a). In both SL and WL plots, topsoil temperature varied between 30 and 40°C in mid-spring, increasing to 40 to 50°C by mid-summer. The MU treatment significantly reduced topsoil temperature from mid-spring to mid-summer as compared with the other aerial competition treatments. During the summer, SL plots showed slightly higher topsoil temperatures as compared with WL sites, although they were significantly different only at the end of

Table 1. Temperature per logging site and aerial competition treatments: (a) in the topsoil; and (b) at 10-cm soil depth

Different lower-case, superscript letters indicate statistical significant differences among treatments at $P < 0.05$. SL, salvage logging; WL, without logging

a)	SL			WL		
	Control	Mulch	Removal	Control	Mulch	Removal
Mid-spring (Nov-12)	35.7 ^b	30.6 ^a	37.8 ^{bc}	36.5 ^{bc}	33.8 ^{ab}	40.6 ^c
Mid-summer (Feb-13)	44.3 ^{ab}	42.3 ^a	49.7 ^{bc}	42.4 ^a	39.9 ^a	45.4 ^c
Late-summer (Mar-13)	23.2 ^b	23.1 ^b	24.5 ^b	18.0 ^a	18.3 ^a	17.8 ^a
b)	SL			WL		
	Control	Mulch	Removal	Control	Mulch	Removal
Mid-spring (Nov-12)	23.8 ^c	15.8 ^{ab}	19.2 ^b	16.3 ^{ab}	15.0 ^a	17.4 ^{ab}
Mid-summer (Feb-13)	20.6 ^c	19.2 ^{ab}	21.2 ^c	19.3 ^{bc}	18.7 ^a	20.1 ^{bc}
Late-summer (Mar-13)	12.8 ^b	12.1 ^a	12.8 ^b	12.1 ^a	11.6 ^a	12.1 ^a

the summer (18.0 and 23.4°C for WL and SL respectively, Table 1a).

Soil temperature at 10 cm belowground was not significantly different between SL and WL plots, but it was significantly different among aerial competition treatments (Table 1b). The MU treatment also showed a significantly lower 10-cm soil depth temperature as compared with the other aerial competition treatments. Temperature at 10 cm soil depth was, in average, 18, 24 and 8°C lower than topsoil temperature in mid-spring, and mid- and late summer respectively.

Soil moisture

During the first growing season after seedling plantation, soil moisture in the first 10-cm soil depth was not significantly different between SL and WL plots in mid-spring, but it was significantly higher at WL plots as compared with SL sites as the growing season advanced (Table 2a). Soil moisture was also significantly different among aerial competition treatments (CO < RE < MU) in mid-spring and mid-summer. In mid-summer, soil moisture was significantly lower in RE as compared with MU treatments (Table 2a). In late summer (end of the first growing season), soil moisture was significantly lower in all competition treatments at SL sites as compared with those located at WL sites. At 20- to 30-cm depth, soil moisture was not significantly different at SL and WL sites in the mid-spring measurement, although it was significantly lower in CO as compared with the other aerial competition treatments (Table 2b). During mid-summer, interactions between sites (SL and WL) and aerial competition treatments were marginally significant ($P = 0.0522$). As it was expected, soil moisture in the CO treatments at SL sites was significantly lower than that at WL sites, whereas CO in this treatment showed the higher moisture values (Table 2b). At the end of the summer,

Table 2. Soil moisture per logging site and aerial competition treatments: (a) average of the first 10-cm soil depth and (b) from 20- to 30-cm soil depth

Different lower-case, superscript letters indicate statistical significant differences among treatments at $P < 0.05$. SL, salvage Logging; WL, without logging

(a)	SL			WL		
	Control	Mulch	Removal	Control	Mulch	Removal
Mid-spring (Nov-12)	7.3 ^a	13.4 ^c	10.7 ^b	11.9 ^{bc}	17.2 ^d	13.3 ^c
Mid-summer (Feb-13)	4.3 ^a	5.8 ^b	4.6 ^{ab}	9.6 ^d	9.4 ^d	7.5 ^c
Late-summer (Mar-13)	12.8 ^a	12.6 ^a	12.6 ^a	19.2 ^b	18.6 ^b	12.7 ^a
(b)	SL			WL		
	Control	Mulch	Removal	Control	Mulch	Removal
Mid-spring (Nov-12)	13.3 ^a	13.9 ^a	14.8 ^a	15.2 ^a	18.4 ^b	17.9 ^b
Mid-summer (Feb-13)	8.4 ^a	10.4 ^{ab}	9.1 ^{ab}	11.2 ^b	9.0 ^{ab}	11.1 ^b
Late-summer (Mar-13)	11.2 ^a	12.3 ^a	11.2 ^a	13.5 ^a	14.5 ^a	13.3 ^a

no significant differences in moisture values were recorded between sites or among aerial competition treatments.

Dynamics of soil moisture at 10-cm soil depth

Soil moisture in the CO and RE aerial competition treatments at WL site was highest during mid-spring, with 25.6 and 23.5% respectively. Aerial competition treatments at the same date but in SL showed intermediate soil moisture values, ~20%, while MU treatment showed the lowest soil moisture values 4.7 and 14.4% in SL and WL, respectively. Soil-moisture dynamics showed a constant decline from the beginning of the spring towards the summer, with some upward pulses that represent mid-summer precipitation events (Fig. 4). Application of the MU treatments appears to have a deleterious effect in both SL and WL site treatments, although the WL treatment seems to rapidly recover moisture values after two rainfall events. After three consecutive rainfall events in mid-summer (February), all treatment effects (i.e. sites and aerial competition) disappeared. Soil moisture steadily diminished in all treatments towards the end of the summer (Fig. 4).

Austrocedrus chilensis performance

Seedling survival

Seedling survival was significantly affected by sites (SL and WL) during the whole study period, and by aerial competition treatments (RE, MU and CO) during the second winter and the second summer analysed (Fig. 5). However, during the first 2 years after starting the experiment, survival was not significantly different in both root (EX and RC) competition treatments.

Survival slowly diminished at SL sites as compared with WL sites after the first summer, and did not present significant differences related to aerial (RE, MU and CO) or root (EX and RC) competition treatments (Fig. 5). After the second

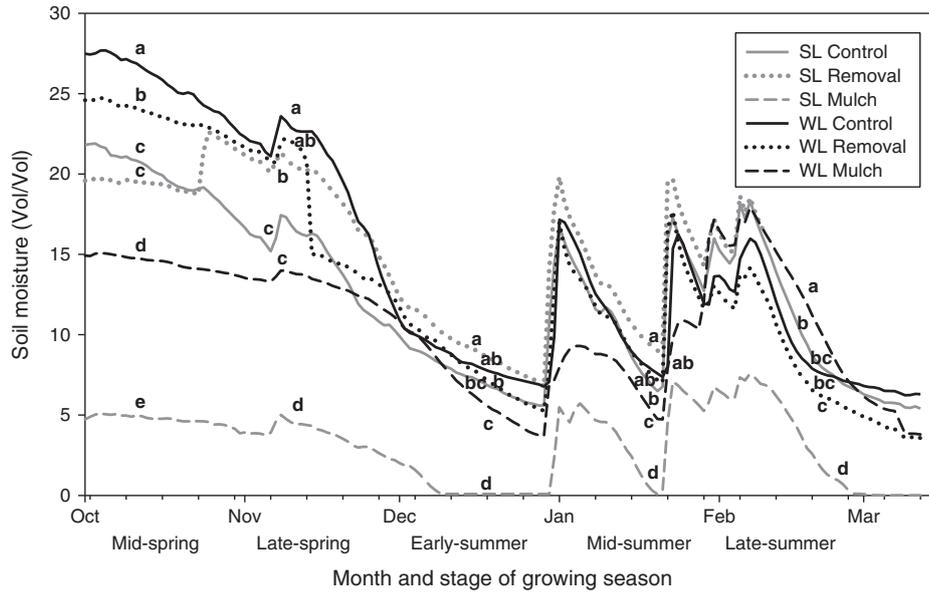


Fig. 4. Soil moisture dynamics at 10-cm soil depth per logging and ground aerial competition treatments during 2013–2014 growing season. Different letters on each trend line for the month and stage of growing season indicate statistical significant differences at $P < 0.05$. SL, salvage logging; WL, without logging.

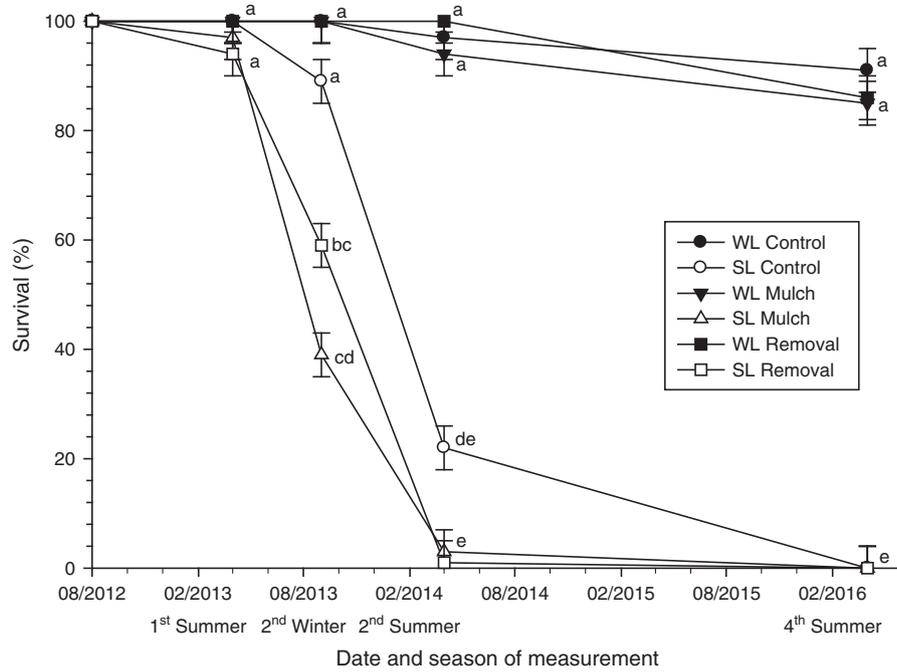


Fig. 5. *Austrocedrus* seedling survival (mean percentage \pm 1 s.e.) during the different seasons considered, and as affected by the combination of logging and aerial competition treatments. Different letters on each trend line for the date and season of measurement indicate statistical significant differences at $P < 0.05$. SL, salvage logging; WL, without logging.

winter, however; seedling survival steadily diminished at SL sites for all aerial competition treatments. At the beginning of the second summer, most seedlings at SL treatment had died, and only 20% of seedlings in the aerial competition control (CO)

remained alive. No seedlings survived at SL sites by the fourth summer after plantation, whereas for all competition treatments of WL, survival remained above 85%, with the only exception of the combination of EX and MU treatments, which was 75%

Table 3. Survival (%) of *Austrocedrus* seedlings during the first, second and fourth summer after plantation in without logging sites (WL, $n = 4$) and as affected by aerial vegetation ($n = 16$ seedlings for each sample unit), and root exclusion treatments ($n = 8$ for each sample unit)

Different lower-case, superscript letters indicate statistical significant differences among treatments at $P < 0.05$. EX, exclusion of roots; RC, root competition

Aerial vegetation treatment	Root exclusion	1st summer and 2nd winter	2nd summer	4th summer
Removal	Yes (EX)	100	100	85 ^{bc}
	No (RC)	100	100	88 ^b
Mulch	Yes (EX)	100	91	75 ^c
	No (RC)	100	97	94 ^{ab}
Control	Yes (EX)	100	97	97 ^a
	No (RC)	100	97	85 ^{bc}

Table 4. Mean height growth (cm) of *Austrocedrus* seedlings during the first summer after plantation per logging site, aerial vegetation and root exclusion treatments

Different lower-case, superscript letters indicate statistical significant differences among treatments at $P < 0.05$. SL, salvage logging; WL, without logging; EX, exclusion of roots; RC, root competition

Site	Aerial vegetation treatment	Root exclusion	Height growth (cm)
SL	Control	No (RC)	0.25 ^f
		Yes (EX)	0.62 ^{ef}
	Mulch	No (RC)	0.80 ^{cdef}
		Yes (EX)	1.08 ^{bcddef}
	Removal	No (RC)	0.72 ^{def}
		Yes (EX)	1.11 ^{bcddef}
WL	Control	No (RC)	1.25 ^{bcdde}
		Yes (EX)	2.03 ^{ab}
	Mulch	No (RC)	1.86 ^{abcde}
		Yes (EX)	2.28 ^{ab}
	Removal	No (RC)	1.92 ^{abcd}
		Yes (EX)	2.56 ^a

(Table 3). At the end of the fourth growing season, significant interaction was found between aerial (RE, MU, and CO) and root competition (EX and RC) treatments. The highest survival was registered in the combination between CO and EX treatments, and in MU and RC treatments. Unexpectedly, the MU and EX treatments showed the lowest survival. The other treatments presented intermediate survival values (Table 3).

Seedling growth

During the first summer after plantation, seedling height growth was significantly different in all treatments: site (SL and WL), aerial (RE, MU and CO) and root competition (EX and RC), although it was not significant with regard to any treatment interactions. Seedlings grew more at WL sites ($P < 0.0001$) and in RE and MU ($P = 0.0099$) and EX ($P = 0.0022$) treatments. Growth was minimal, with absolute values that varied from 0.25 to 1.11 cm at SL sites, although they varied from 1.25 to 2.56 cm at WL sites. The combination of WL with RE and EX treatments showed significantly higher growth as compared

with all SL treatments and to WL-CO-RC treatment combination (Table 4).

From the first summer up to the end of the experiment, seedling growth was only measured at WL sites because all seedlings at SL sites had already died. Within WL treatment, seedling height growth was similar in all aerial and root competition treatments, and averaged 5.6 ± 1.4 cm during the period 2012–2014 (Fig. 6a). From 2014 to the end of the experiment in 2016, seedlings grew at an average of 19.3 ± 1.5 cm in the CO treatment, significantly higher than the RE and MU treatments, which grew 14.1 ± 0.5 and 16.3 ± 1.5 cm respectively (Fig. 6a). No significant differences in height growth were found between EX and RC root competition treatments during this period. By considering the whole study period (2012–2016), all seedlings in this WL site treatment grew 21.0 ± 1.2 cm on average, presenting no significant differences among all aerial treatments (RE, MU and CO) or between root competition (EX and RC) treatments (Fig. 6a). Seedling collar diameter was not significantly different for all aerial and root competition treatments, and grew on average 1.9 ± 0.3 mm for the 2014–2016 period and 3.4 ± 0.4 mm for the 2014–2016 period. For the complete 2012–2016 period, collar diameter grew on average 5.1 ± 0.5 mm (Fig. 6a).

Hydric status of *A. chilensis* seedlings

In mid-summer of the first growing season, predawn and midday hydric status of cypress seedling branches was associated with site (SL and WL) but not to aerial or root competition treatments. At predawn, seedlings grown at SL sites showed significantly ($P < 0.001$) lower (more negative) water potential average values (-1.6 MPa) than at WL sites (-1.2 MPa) (Fig. 7a). At midday, when the were higher evapotranspiration rates, average values were -2.2 MPa for SL plots and -1.6 MPa for WL plots (Fig. 7b).

Discussion

This study showed that, in *A. chilensis* post-burned stands, salvage logging significantly affected survival and growth of restored *A. chilensis* seedlings. Results showed that two growing seasons after plantation, seedling survival was $< 10\%$ at SL sites, whereas it was near 90% at WL sites. Four seasons after establishment, and regardless of sub-treatments, no seedlings survived at SL sites, whereas over 75% survived at WL plots.

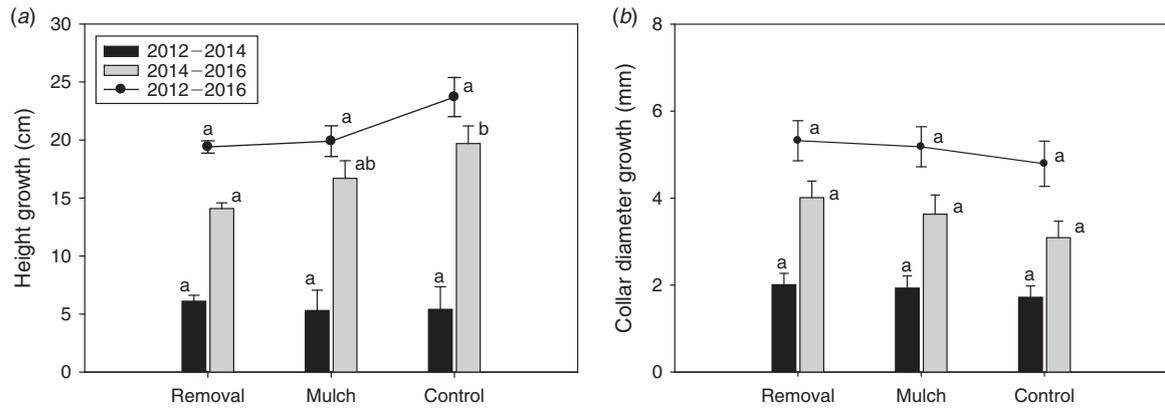


Fig. 6. (a) *Austrocedrus* height (cm) and (b) collar diameter (mm) growth (mean \pm 1 s.e) in without logging sites and as affected by aerial competition treatments during 2012–2014, 2014–2016, and the complete 2012–2016 study period. Different letters indicate statistical significant differences at $P < 0.05$.

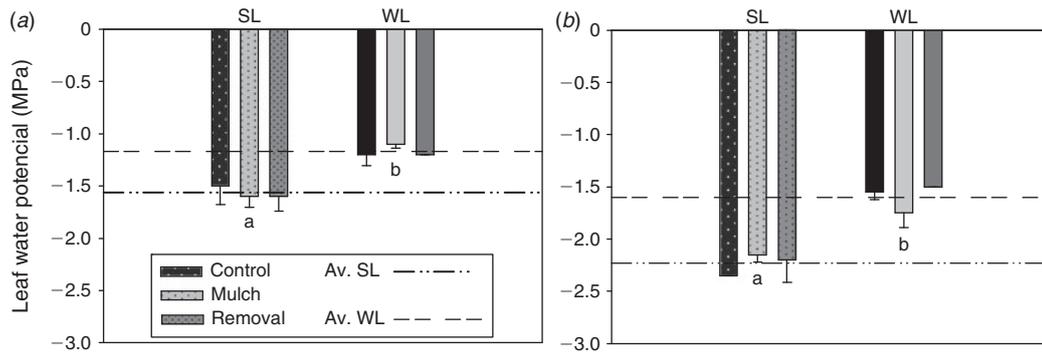


Fig. 7. Leaf water potential (mean \pm 1 s.e.) at predawn (a) and midday (b) in salvage logging (SL) and without logging (WL) sites and as affected by aerial competition treatments. Different letters below each bar graph indicate statistical significant differences at $P < 0.05$.

These findings are useful to help alleviate the lack of reforestation typically observed at burned and logged sites.

Tree seedlings, in general, experience transplantation shock during the first growing season after planting, while they try to acclimate to their new site conditions (Hobbs 1984; Ouzts *et al.* 2015). During their acclimation period, *A. chilensis* seedlings showed a little shoot growth, while post-fire herbaceous vegetation established vigorously. Competition by neighbouring species is one of the main interactions affecting vegetation growth in early post-fire successional stages (Jobidon *et al.* 1998; Löf *et al.* 1998; Dinger and Rose 2009; Maguire *et al.* 2009). However, competition might affect growth in a great variety of ways, and it may differ among communities (Reader *et al.* 1994). In a mesic area similar to our study site, *A. chilensis* seedling survival and growth were positively associated with post-fire cover during the 2 years after plantation (Urretavizcaya *et al.* 2017). In our study, however, this nursing effect appeared two seasons after acclimation. Neighbouring plants seemed to exert a nursing rather than a competing effect, as *A. chilensis* seedlings grew vigorously from the third season onwards. These results agree with the study of Holmgren and Scheffer (2010), who reported results for a similar mesic site, and with other

authors (Veblen *et al.* 1995; Gobbi 1999) dealing with other pure and fire-undisturbed *A. chilensis* stands. These studies showed that facilitation overcomes other interaction effects, and occurs in small to intermediate canopy gaps (Veblen *et al.* 1995) associated with high herbaceous and shrub cover (Gobbi 1999). Our study also showed that the use of root-exclusion tubes neither benefited nor harmed seedling performance.

At the beginning of the first summer after seedling plantation, PAR at SL sites was almost 4 times higher than at WL plots, significantly increasing topsoil temperature at the end of that period. These factors created a more stressful microenvironment at SL sites, perhaps causing its high seedling mortality as compared with WL plots by the end of the second summer. By contrast, at WL sites, standing snags reduced PAR reaching the soil surface, attenuating topsoil heating and improving overall seedling growth micro-environmental conditions. This situation is similar to what Castro *et al.* (2011) reported for plants grown in other Mediterranean environments. Although PAR and topsoil temperature at the end of the summer were significantly higher at SL sites as compared with WL plots, soil moisture was similar in both sites. Why, then, when having similar soil moisture levels, was higher mortality produced at SL sites as

compared with WL plots? The answer could be found by considering the way *A. chilensis* developed physiological mechanisms to avoid hydric stress. Gyenge *et al.* (2005) demonstrated that *A. chilensis* avoids water stress by strong stomatal control. In our study, *A. chilensis* seedlings were able to tolerate soil-water deficit in both SL and WL treatments, but were unable to cope with high radiation levels at SL sites as compared with WL plots. This inference is supported by other studies with the same species, which showed that, even with adequate soil moisture, high levels of incident radiation increase the risk of photo inhibition and high cell damage because of overheating, causing high *A. chilensis* seedling mortality (Gyenge *et al.* 2007; Caselli *et al.* 2019). Additionally at WL sites, shade protection exerted by standing snags not only reduced PAR and diurnal topsoil temperatures, but also attenuated low night temperatures especially during the winter, thus creating a better microclimate for planted seedlings. This inference is supported by Castro *et al.* (2011), whose study showed that stands with snags maintain higher topsoil temperatures in winter as compared with nearby bare-tree areas.

After the first dry summer season, soil water returned to field capacity in all treatments except in those covered by mulch. This MU treatment may have obstructed normal water infiltration into the soil profile, being more noticeable at SL sites than in WL plots, because of its higher levels of radiation and the desiccant effect of winds. This may help explain the poor performance of seedlings grown in this MU treatment as compared with other treatments. Mulch is a mechanical barrier that may protect the seedling collar from high PAR and high topsoil temperatures, but also intercepting precipitation and dew condensation. This may be the reason why, in other study dealing with *A. chilensis* seedlings (Lallement *et al.* 2010), no differences in survival and growth were detected by using 2-cm shredded wood as mulch, as compared with unprotected control.

Conclusions

Severe fires kill *A. chilensis* trees and seeds, producing drastic modifications in stand structure. Post-fire macro- and micro-environmental conditions, particularly high PAR levels, may limit seedling survival and growth if salvage logging is applied to remaining *A. chilensis* snags, making any restoration efforts futile. In burned sites without logging, however, PAR attenuation by a canopy of remaining snags benefitted *A. chilensis* seedling survival, and no additional control of early successional herbaceous species should be necessary to attain high seedling survival. Also at SL sites, diminution of aerial or belowground competition did not ameliorate the deleterious effects of PAR. The addition of mulch as it was applied and, contrary to what was expected, did not improve seedling survival or growth. Our results agree with other studies (i.e. McIver and Starr 2000; Castro *et al.* 2011), which showed that post-fire standing snags provide significant shade, diminishing soil-surface heating and modifying surface micro-environmental conditions as compared with nearby salvage logging areas. However, standing post-fire snags are often logged for economic reasons (Castro *et al.* 2011), posing socio-ecological conflicts should the stand need to be actively restored, because seedling survival under this situation is almost nil (Keyser *et al.* 2009).

Taking into account these results, and to achieve high seedling establishment rates, limited logging is suggested in burned forest, such as the one presented in this case. One generally accepted recommendation is to leave ~50% of the snags of each diameter class, and all snags whose diameter at breast height is >50 cm or were >150 years old (Beschta *et al.* 2004). This recommendation may also be compatible with objectives related to the many other ecological roles of snags in the recovering landscape, including the provision of habitat for a variety of species and playing a key role in biological and physical processes (Thomas 1979; Donato *et al.* 2013). In burned *A. chilensis* stands, if salvage logging is the only alternative, a restoration plan should consider waiting for seedling plantation till the lower strata of plant cover and some shrubs has recovered. These shrubs may act as nurse plants, facilitating seedling survival and assuring further establishment (Kitzberger *et al.* 2000; Urretavizcaya *et al.* 2012). Aerial devices could also be used to attenuate direct radiation during early stages of seedling establishment (Urretavizcaya and Defossé 2013). Finally, post-fire management of burned forests should consider a variety of options to help reconcile and better balance the competing society's economical needs with the benefits of sustainable management through ecological restoration.

Conflicts of interest

The authors declare that they have no conflicts of interest.

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