



**HAL**  
open science

## Tree diversity reduces pine infestation by mistletoe

Inge van Halder, Bastien Castagneyrol, Cristóbal Ordóñez, Felipe Bravo,  
Miren del Río, Lucile Perrot, Herve Jactel

### ► To cite this version:

Inge van Halder, Bastien Castagneyrol, Cristóbal Ordóñez, Felipe Bravo, Miren del Río, et al.. Tree diversity reduces pine infestation by mistletoe. *Forest Ecology and Management*, 2019, 449, pp.1-8. 10.1016/j.foreco.2019.117470 . hal-02618879

**HAL Id: hal-02618879**

**<https://hal.inrae.fr/hal-02618879>**

Submitted on 26 Oct 2021

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

# 1 **Tree diversity reduces pine infestation by mistletoe**

2

3 Inge van Halder\*<sup>1</sup>, Bastien Castagneyrol <sup>1</sup>, Cristóbal Ordóñez <sup>2,4</sup>, Felipe Bravo <sup>2,4</sup>, Miren del Río <sup>2,3</sup>,

4 Lucile Perrot<sup>1</sup>, Hervé Jactel<sup>1</sup>

5

6 1 BIOGECO, INRA, Univ. Bordeaux, 33610, Cestas, France

7 2 Sustainable Forest Management Research Institute, University of Valladolid & INIA, Av. Madrid 44,

8 Palencia 34004, Spain

9 3 Department of Silviculture and Forest Management, INIA, Forest Research Centre, Ctra. A Coruña,

10 km 7.5, Madrid 28040, Spain

11 4 Departamento de Producción Vegetal y Recursos Forestales, E.T.S de Ingenierías Agrarias,

12 Universidad de Valladolid, Palencia, Spain

13

14 \* **Corresponding author:** [inge.van-halder@inra.fr](mailto:inge.van-halder@inra.fr)

15

## 16 **Abstract**

17 The pattern that a given tree species suffers less damage when growing with heterospecific

18 neighbors than amongst conspecific plants, i.e. associational resistance, is common for insect

19 herbivores and many fungal pathogens. However, associational resistance to parasitic plants has

20 never been tested in a replicated study. Using paired forest plots, we investigated whether tree

21 diversity triggered associational resistance to a tree parasite, the European mistletoe *Viscum album*

22 ssp. *austriacum*, by comparing pure stands of Scots pine (*Pinus sylvestris*) with mixtures of Scots pine

23 and Maritime pine (*Pinus pinaster*) in northern Spain. Maritime pine, with 1.2% of trees being

24 infested, was considered a non-host species in the study area. The infestation level of Scots pines

25 was significantly higher in pure plots (45.1%) than in mixed plots of Scots pines and Maritime pines

26 (25.4 %). Our study is the first to quantify associational resistance to a plant parasite in mixed vs.

27 pure forest stands and suggests that mechanisms proposed to explain associational resistance to  
28 insects and pathogens also apply to plant parasites. Scots pine trees that were taller than the  
29 surrounding trees had a higher infestation probability, in both pure and mixed stands. Scots pine  
30 trees growing in mixtures were slightly lower than Maritime pines, suggesting that associational  
31 resistance was partly driven by reduced relative tree height. However, the effect of plot type (pure  
32 vs. mixed) remained significant after the effect of tree height was accounted for, thus indicating that  
33 other factors also contributed to lower mistletoe infestation in mixed plots. In particular, the  
34 behavior of birds dispersing mistletoe seeds might differ in mixed vs. pure stands.

35

36 **Keywords** : Associational resistance; Biodiversity; Forest management; Mistletoe; *Pinus pinaster*;  
37 *Pinus sylvestris*; *Viscum album*

38

## 39 **1. Introduction**

40 Increasing evidence is showing that tree diversity contributes to forest ecosystem functioning and  
41 the provision of ecosystem services (Brocknerhoff et al., 2017). Mixed-species forests exhibit higher  
42 productivity, plant and animal biodiversity, resistance to disturbances and less insect damage than  
43 tree monocultures ( Castagneyrol et al., 2014; Jactel and Brocknerhoff, 2007; Jactel et al., 2018, 2017).  
44 Associational resistance, i.e. the fact that a given tree suffers less damage when growing with  
45 heterospecific neighbors than amongst conspecific trees (Barbosa et al., 2009) is a common pattern  
46 for herbivore insects (Castagneyrol et al., 2014) and root pathogens (Jactel et al., 2017), while for  
47 foliar pathogens the effect of mixed stands seems more variable (Jactel et al., 2017). The effect of  
48 tree diversity on insect herbivores and pathogens can be attributed to two, often non-independent,  
49 processes: a lower density of host plants in mixtures or a pure associational effect (Hambäck et al.,  
50 2014). Two theories explain the relationship between host density and insect/pathogen abundance  
51 in pure vs. mixed stands. The resource concentration hypothesis (Root, 1973) predicts higher

52 herbivore abundance in pure stands because insects are more likely to find, remain and reproduce  
53 on host trees that are more abundant in such stands. On the contrary, when host density is low in  
54 mixed stands, herbivores may concentrate on the few available hosts, leading to a higher infestation  
55 level per tree (resource dilution hypothesis (Damien et al. 2016; Otway et al., 2005)). Non-host trees  
56 can also trigger associational resistance independently of host density. For example, reduced  
57 apparency of focal tree species, whereby non-host trees in the mixture disrupt visual and chemical  
58 cues emitted by host trees, can explain associational resistance to actively dispersing herbivores and  
59 insect vectored pathogens (Castagneyrol et al., 2013). For airborne pathogens or those dispersing  
60 through root contact the presence of non-host trees can provide a physical barrier to contamination  
61 of neighboring host trees, leading to lower infestation levels in mixed stands (Jactel et al., 2017). The  
62 presence of non-host trees can also promote the presence and abundance of natural enemies  
63 providing biological control of insect herbivores (Jactel and Brockerhoff, 2007) or pathogens (Jactel et  
64 al., 2017). While these resistance effects of tree diversity have frequently been observed for pest  
65 insects and, to a lower extent, for pathogens, they have very rarely been studied for parasitic plants  
66 such as mistletoes. In this study, we examined the prevalence of European mistletoe (*Viscum album*  
67 *ssp. austriacum*) in pure vs. mixed pine forests.

68

69 Mistletoes are hemi-parasitic plants, with about 1300 species from five families within the Santales  
70 (Watson, 2001). The European mistletoe, *Viscum album*, is a perennial, hemi-parasitic plant that only  
71 lives on woody plants (Zuber and Widmer, 2009) and extracts water and minerals from its host. In  
72 Europe four subspecies occur that differ in distribution and host range (Zuber, 2004). *V. album ssp.*  
73 *austriacum* occurs in Spain and Central Europe, mainly on *Pinus* species and rarely on *Larix* and *Picea*  
74 (Zuber, 2004; Zuber and Widmer, 2009). *V. album* is a species of interest because it is a host for  
75 several specialized insect species, a food source for birds and it contains pharmacological substances  
76 (Briggs, 2011; Lázaro-González et al., 2017; Zuber, 2004). However, high levels of *V. album* infestation  
77 have negative effects on tree growth (Noetzli et al., 2003; Rigling et al., 2010; Sangüesa-Barreda et

78 al., 2013) and contribute to tree death especially when associated with drought stress (Dobbertin  
79 and Rigling, 2006; Mutlu et al., 2016; Tsopelas et al., 2004). The relationship with drought stress  
80 indicates that with climate change the damage caused by *V. album* will probably increase in the  
81 future. Moreover, *V. album* is expanding its range. An upward shift in altitude has been observed in  
82 the last century, which seems linked with global warming (Dobbertin et al., 2005).

83 As for many other mistletoe species, the seeds of *V. album* are dispersed by birds. Seed dispersal, the  
84 first step in the infestation process, seems an essential process in explaining *V. album* infestation and  
85 spatial distribution and bird behavior may lead to a higher seed deposition on certain trees, for  
86 example on tall trees or on trees at stand edges (Durand-Gillmann et al., 2014; Vallauri, 1998). The  
87 most important dispersers of *V. album* seeds are mistle thrush (*Turdus viscivorus*), other *Turdus*  
88 species, waxwing (*Bombycilla garrula*) and blackcap (*Sylvia atricapilla*) (Mellado and Zamora, 2014;  
89 Zuber, 2004). The most effective dispersers of *V. album* in southern Europe are thrushes (Mellado and  
90 Zamora, 2014). Thrushes eat the berries and defecate the seeds. With a transit time of about half an  
91 hour the seeds can be dispersed over distances of more than 20 km by migrating birds (Frochot and  
92 Sallé, 1980). However, most seed dispersal occurs at shorter distances by thrushes foraging in areas  
93 with *V. album* infested trees or by thrushes holding and defending territories of groups of *V. album*  
94 infested trees (Skórka and Wójcik, 2005; Snow and Snow, 1984). Blackcaps, another seed disperser,  
95 disperse the seeds at even closer distances, mainly within the same tree, as they feed on the skin of  
96 the berry and leave the seed on a shoot nearby the *V. album* shrub (Zuber, 2004). The behavior of  
97 birds to spend more time on infested hosts than non-infested hosts thus leads to an aggregation of  
98 mistletoes within hosts (Aukema and Martinez de Rio, 2002).

99

100 The effects of tree diversity on mistletoe infestation, including effects of host density and of pure  
101 associational effects of the accompanying tree species, can be multiple, since tree diversity can  
102 influence both the behavior of seed dispersing birds and mistletoe-host interactions (see Figure 1).  
103 Birds are active seed dispersers and it is likely that processes generating associational effects for

104 insect herbivores also act upon birds, such as disruption of host finding cues. Mistletoe infestation is  
105 often higher on taller trees as observed for *V. album* (Durand-Gillmann et al., 2014; Kolodziejek and  
106 Kolodziejek, 2013) and for several other mistletoe species (Aukema and Martínez del Rio, 2002;  
107 Donohue, 1995; Roxburgh and Nicolson, 2008; Shaw et al., 2005; Smith and Reid, 2000; Teodoro et  
108 al., 2010). This pattern was proposed to result mainly from bird preferences for more apparent trees  
109 rather than to differences in host tree suitability (Aukema and Martínez del Rio, 2002; Roxburgh and  
110 Nicolson, 2008). In mixed stands, where infested trees can be partly hidden by non-infested  
111 neighbours, birds foraging for mistletoe fruits may have greater difficulty to find their resource. Birds  
112 can also react to local mistletoe abundance. The behavior of birds to spend more time in groups of  
113 trees with high mistletoe abundance gives that in those areas both infested and uninfested hosts  
114 have a higher exposure to seed dispersers than in areas with a low infestation level (Aukema, 2003).  
115 This mechanism may lead to a direct effect of the non-host density in mixtures as birds will  
116 encounter less mistletoe hosts in these stands and shorten probably their foraging time.  
117 Tree diversity can also affect host-mistletoe interactions. Host plants have developed structural and  
118 biochemical defenses to mistletoe infestation (Aukema, 2003) and the expression of tree defensive  
119 traits have been shown to be influenced by the identity of neighboring trees (Castagneyrol et al.,  
120 2018; Rosado-Sánchez et al., 2018). Trees may also differ in quality for mistletoes. For example, in  
121 areas where water is limiting, mistletoes are more likely to establish on host trees with better access  
122 to water (Watson, 2009). This process probably differs between pure and mixed stands, with drought  
123 responses of tree species varying according to the composition of mixtures (Forrester and Bauhus,  
124 2016; Grossiord, 2018).

125

126 The objective of our study was to evaluate the effect of mixed vs. pure stands on the infestation level  
127 by *V. album* and to identify tree and stand characteristics linked to associational resistance or  
128 susceptibility. We studied the presence of *V. album* ssp. *austriacum* in pure Scots pine (*Pinus*  
129 *sylvestris*) forests and mixed forests of Scots pine and Maritime pine (*P. pinaster*) in northern Spain.

130 Preliminary observations in the study area indicated that Scots pine was much more sensitive to *V.*  
131 *album ssp. austriacum* than Maritime pine that could be considered a non-host. As such,  
132 associational effects in mixed stands of Scots pine and Maritime pine would result from a  
133 combination of both host density effects, whereby Scots pine density is lower in mixtures as  
134 compared to monocultures, and pure associational effects whereby, for a given Scots pine density,  
135 the presence of Maritime pines might reduce the probability of infestation on neighboring Scots  
136 pines.

137 In particular, our study aimed to answer the following questions:

- 138 i) does *V. album* infestation level of Scots pines differ between pure and mixed stands?
- 139 ii) does the presence of *V. album* depend on relative tree height (i.e. how much a given tree is  
140 higher than its neighbors) ?
- 141 iii) what are the relative effects of host and non-host density on *V. album* infestation?

142

143

## 144 2. Material and methods

145

### 146 2.1 Study area and plots

147 The study was conducted in northern Spain, in an area of approximately 50,000 ha covered with  
148 Mediterranean forests of Scots pine and Maritime pine. The area covers the transition zone between  
149 the natural Scots pine (higher elevation) and Maritime pine (lower elevation) forests in the Northern  
150 Iberian mountain range, belonging to the provenance regions “Montaña Soriano Burgalesa” and  
151 “Montaña de Soria Burgos” respectively (Martín et al 1998). Mean annual temperature of the area is  
152 9.0 °C, mean annual precipitation ranges from 715 to 888 mm and elevation ranges from 1090 to  
153 1277 m a.s.l.. To study the effect of species mixture on forest productivity and structure in this area,  
154 Riofrio et al. (2017) selected in 2014-2015 36 circular plots with a radius of 15 m. Plots were selected  
155 as representative parts of forest composition and structure in the surrounding area. Plots were

156 grouped into 12 triplets of mixed plots and the corresponding pure plots of Scots pine and Maritime  
157 pine (i.e. 36 plots in total). All triplets were situated in an area of 40 km length by 20 km width, with  
158 coordinates of plots between 41°46'15.2"N - 41°53'46.6"N and 2°55'39.9"W - 3°20'43.4"W. Distance  
159 between plots within the same triplet was always shorter than 1 km. Tree age of plots ranged  
160 between 38 and 139 years. For Scots pines, the median difference in age between the pure and  
161 mixed plot of a triplet was 7.5 years, with a minimum of 2 years for the triplet with the youngest  
162 plots and a maximum of 38 years for the triplet with the oldest plots. For Maritime pine the median  
163 value was 8 years, and varied between 2 years to 34 years for the oldest triplet. Additional  
164 information about stand characteristics are included in Riofrío et al (2017, Supplementary Material).  
165 For each tree, the diameter at breast height and the height were measured (see for more details on  
166 forest management, plot selection, and measurements Riofrío et al. (2017).

167 Preliminary observations on Maritime pine trees in pure Maritime pine stands and in mixed stands  
168 revealed that *V. album* was nearly absent on Maritime pine in this area, in sharp contrast with the  
169 high prevalence on Scots pine. We therefore considered Maritime pine as non-host and did not  
170 survey pure Maritime pine plots. The study was thus based on *V. album* infestation in 12 pairs of  
171 pure stands of *P. sylvestris* and mixtures of *P. sylvestris* and *P. pinaster*. Mixed plots had varying  
172 proportions of tree species, with *P. sylvestris* representing 37-77% of the total number of trees and  
173 32-71% of total basal area. In plots classified as pure plots, *P. sylvestris* accounted for at least 91% of  
174 the total number of trees and 85% of total basal area (**Table 1**). Other plot characteristics are  
175 indicated in **Table 1** and **Appendix A (Fig. A.1 and A.2; Table A.1)**. Since the average total number of  
176 trees was the same in mixed and pure plots and Maritime pine thus partly replaced Scots pine in  
177 mixed plots, the number of trees of the two species was negatively correlated for the 24 plots ( $r = -$   
178 0.54) as was their basal area ( $r = -0.74$ , Appendix **Fig. A.1**).

179 In March 2017, two observers, positioned at different sides of the tree, assessed together the  
180 presence/absence of *V. album* on each tree inspecting the complete tree crown and stem with  
181 binoculars. A total of 255 Maritime pines and 843 Scots pines were inspected in the 12 pairs of plots.



182  
 183  
 184  
 185  
 186  
 187

**Table 1.** Compositional and structural characteristics of the mixed and pure plots. For each tree species, mean (minimum, maximum) values per plot are given for the number of trees (expressed per plot and per hectare), DBH (Diameter at Breast Height), tree height, basal area and % trees calculated for the total number of trees (N) or basal area (BA).

	mixed plots (N=12)		pure <i>P. sylvestris</i> plots (N=12)	
	<i>P. sylvestris</i>	<i>P. pinaster</i>	<i>P. sylvestris</i>	<i>P. pinaster</i>
number of trees / plot	26.3 (14, 42)	19.7 (9, 36)	45.3 (26, 76)	1.6 (0, 6)
number of trees / ha	372.5 (198.1, 594.2)	278.2 (127.3, 509.3)	640.2 (367.8, 1075.2)	22.4 (0.0, 84.9)
DBH (cm)	29.6 (20.2, 40.3)	37.5 (23.5, 47.7)	30.3 (20.4, 39.8)	42.0 (26.0, 56.7)
tree height (m)	19.3 (14.0, 24.7)	20.4 (14.9, 26.9)	20.1 (14.8, 24.5)	21.2 (15.3, 26.3)
basal area (m <sup>2</sup> /ha)	26.2 (13.0, 45.9)	30.8 (11.1, 48.7)	45.8 (29.3, 59.1)	2.6 (0.0, 7.1)
% trees (N)	57.4 (36.8, 76.9)	42.6 (23.1, 63.2)	97.0 (90.9, 100.0)	3.0 (0.0, 9.1)
% trees (BA)	46.3 (31.9, 71.4)	53.7 (28.6, 68.1)	94.4 (85.0, 100.0)	5.6 (0.0, 15.0)

188  
 189

## 190 2.2 Statistical analyses

191 Data were analyzed at the plot level and at the individual tree level. At the plot level, we analyzed the  
 192 proportion of Scots pine trees infested with *V. album* using three sets of explanatory variables. In the  
 193 first model, we used plot type (pure vs. mixed plots) as explanatory categorical variable. Since a  
 194 possible effect of plot type may be driven by either a dilution of Scots pine and/or an increase in  
 195 Maritime pine, we ran two other models by substituting plot type by i) the basal area of Scots pine +  
 196 the basal area of Maritime pine + their interaction or ii) the number of Scots pine trees + the number  
 197 of Maritime pine trees + their interaction. Combining both Scots pine and Maritime pine abundance  
 198 in the same model allowed addressing both the effect of host concentration (here Scots pine) and  
 199 the pure effect of the associated species (here the abundance of Maritime pine). For these three  
 200 models we used generalized linear mixed models (GLMM) with a binomial error and a *logit* link  
 201 function on a response variable consisting of the number of infested Scots pine trees vs. the number

202 of non-infested Scots pine trees per plot. To take into account the structure of the dataset with  
203 paired plots we used Pair identity (12 pairs of plots) as a random factor.

204 We used the same general approach to analyze the probability of mistletoe infestation at the level of  
205 individual trees, but further accounted for tree-level covariates. For the analyses at the tree level we  
206 first estimated the individual relative tree height ( $\Delta H$ ), which indicates how much taller or lower a  
207 tree is as compared to its neighbors (Castagneyrol et al., 2013; Damien et al., 2016). We calculated  
208 for each Scots pine tree its  $\Delta H$  by subtracting from the height of the tree the mean height of the trees  
209 in the corresponding plot. As such,  $\Delta H > 0$  indicates that a tree is higher than the mean canopy  
210 height. In order to verify if  $\Delta H$  was independent of the sampling design, we first tested if the height  
211 and  $\Delta H$  of Scots pine trees differed between mixed and pure plots using linear mixed models (LMM)  
212 with plot type as explanatory variable. Next we analyzed the probability of a Scots pine tree being  
213 infested by *V. album* by using three sets of explanatory variables. In the first model, we analyzed the  
214 effect of  $\Delta H$ , plot type (pure vs. mixed) and their interaction on the presence/absence of *V. album* on  
215 individual Scots pine trees using a GLMM with binomial error and a *logit* link function. As for the  
216 analyses at the plot level, we replaced plot type by i) the basal area of Scots pine, of Maritime pine  
217 and their interaction and ii) the number of trees of each species and their interaction. For all models  
218 at the tree level we used as random factors Plot identity nested within Pair identity to account for  
219 the nested structure of the dataset where trees were incorporated in a plot, that belonged to a pair  
220 of plots (Schielzeth and Nakagawa, 2013).

221 For all models, both at the plot and tree level, we applied a model simplification procedure by  
222 comparing nested models, with vs. without the variable of interest. We sequentially removed  
223 predictors, starting with the least significant, while applying marginality principle where the principal  
224 effects were not removed if involved in a significant interaction. Significance of effects was tested by  
225 comparing models with and without the term with type II Wald chi-square tests on log likelihood  
226 ratios. For model validation we visually checked model residuals. For the simplified models,  $R^2$  values  
227 were calculated to estimate the variance explained by fixed effects (marginal  $R^2$ ,  $R^2_m$ ), and by fixed

228 plus random effects (conditional  $R^2$ ,  $R^2_c$ )(Nakagawa and Schielzeth, 2013). Variables were scaled  
229 before analyses.  
230 All analyses were carried out in R (R Core Team, 2019). The following functions and libraries were  
231 used: glmer function from lme4 package (Bates et al., 2015), r.squaredGLMM from MuMin package  
232 (Barton, 2018), Anova from car package (Fox and Weisberg, 2011) and simulateResiduals from  
233 DHARMA package (Hartig, 2019) for residual plots.

234

### 235 **3. Results**

236 The overall *V. album* infestation level was 35.8 % for Scots pine trees (n= 843 trees) and 1.2 % for  
237 Maritime pine trees (n= 255 trees), confirming that Maritime pine can be considered a non-host  
238 species for *V. album* ssp. *austriacum* in the study area.

239 The infestation level of Scots pines was almost twice as high in pure plots as in mixed plots ( $X^2 = 37.2$ ,  
240  $df = 1$ ,  $P < 0.001$ ), with a mean infestation level of  $45.1 \pm 8.4$  % ( $\pm$  SE) in pure plots vs.  $25.4 \pm 6.8$  % in  
241 mixed plots (**Fig. 2**). However, plot type *per se* only explained a limited amount of variance in *V.*  
242 *album* infestation ( $R^2_m = 0.062$ ,  $R^2_c = 0.390$ ).

243

244 For the model using basal area of both tree species as explanatory variables, only the basal area of  
245 Scots pine trees was selected in the final model, showing an increase in infestation level with  
246 increasing Scots pine basal area (**Table 2**). On the contrary, for the model using number of trees only  
247 the number of Maritime pine trees was selected, showing an increase in infestation level with  
248 decreasing number of Maritime pine trees (**Table 2**). Therefore, although they did not retain the  
249 same variables as significant predictors, both models yielded consistent results whereby mistletoe  
250 infestation was higher where host-trees were more abundant and where non-host trees were less  
251 abundant. The model using basal area of Scots pines as an explanatory variable explained more  
252 variance in *V. album* infestation ( $R^2_m = 0.14$ , Table 2) than the model using the number of Maritime  
253 pine trees ( $R^2_m = 0.07$ , Table 2).

254

255 **Table 2.** Summary of models testing the effects of basal area (BA) and tree number (N) of *P. sylvestris* and *P.*  
 256 *pinaster* on *V. album* infestation level of *P. sylvestris* at the plot level. Explanatory variables in bold had a  
 257 significant effect (at  $P < 0.05$ ).  $R^2m$  and  $R^2c$  are marginal and conditional  $R^2$ , respectively, and are calculated for  
 258 the final model resulting from model simplification.

Response	Predictors	Estimate ( $\pm$ SE)	$\chi^2$	Df	P-value	$R^2m$ ( $R^2c$ )
infestation level	<b>BA <i>P. sylvestris</i></b>	1.32 ( $\pm$ 0.36)	13.48	1	< 0.001	0.14 (0.41)
	BA <i>P. pinaster</i>	0.40 ( $\pm$ 0.29)	1.76	1	0.185	
	BA <i>Ps</i> x BA <i>Pp</i>	0.21 ( $\pm$ 0.26)	0.68	1	0.411	
infestation level	N <i>P. sylvestris</i>	0.38 ( $\pm$ 0.23)	1.61	1	0.204	
	<b>N <i>P. pinaster</i></b>	-0.43 ( $\pm$ 0.16)	14.43	1	< 0.001	0.07 (0.40)
	N <i>Ps</i> x N <i>Pp</i>	0.25 ( $\pm$ 0.25)	1.04	1	0.307	

259

260

261 The height of Scots pine trees was not statistically different between mixed and pure plots ( $\chi^2 = 1.43$ ,  
 262  $df = 1$ ,  $P = 0.230$ ). However,  $\Delta H$  (i.e. the difference between individual Scots pine tree height and  
 263 mean plot height) was slightly, but significantly lower in mixed plots than in pure plots ( $\chi^2 = 8.62$ ,  $df =$   
 264  $1$ ,  $P = 0.003$ ), with a mean  $\Delta H$  of -0.45 m in mixed plots and -0.003 m in pure plots, indicating that  
 265 Scots pines were on average lower than Maritime pines in mixed plots.

266

267 At the individual tree level, both  $\Delta H$  and plot type had significant and independent effects on *V.*  
 268 *album* infestation probability (**Table 3**). The probability of individual Scots pines being infested  
 269 increased with increasing  $\Delta H$  and was higher in pure than in mixed plots (**Fig. 3**). The fact that plot  
 270 type remained significant after the effect of  $\Delta H$  was accounted for, and conversely, indicates that  
 271 factors other than those related to relative tree height additionally contributed to the effect of plot  
 272 type on *V. album* infestation probability. When plot type was replaced by the basal area of the two  
 273 tree species,  $\Delta H$  and basal area of Scots pine were selected in the final model (**Table 3**). The  
 274 infestation probability increased with  $\Delta H$  and with the basal area of Scots pines in the plot. For the

275 model including  $\Delta H$  and the number of trees of each species,  $\Delta H$  and the number of Maritime pines  
 276 were selected (**Table 3**), leading to a higher infestation probability with increasing  $\Delta H$  and decreasing  
 277 number of Maritime pine trees per plot.

278  
 279

280 **Table 3.** Summary of models testing the effects of the individual relative tree height of *P. sylvestris* ( $\Delta H$ ) and  
 281 plot composition on infestation probability by *Viscum album* of individual *P. sylvestris* trees. The effect of the  
 282 following predictors on *V. album* infestation probability of individual *P. sylvestris* trees were tested in separate  
 283 models: 1) relative tree height ( $\Delta H$ ), plot type (pure or mixed) and their interaction, 2)  $\Delta H$ , basal area (BA) of *P.*  
 284 *sylvestris* and of *P. pinaster* and their interaction and 3)  $\Delta H$ , tree number (N) of *P. sylvestris* and of *P. pinaster*  
 285 and their interaction. Explanatory variables in bold characters had a significant effect (at  $P < 0.05$ ).  $R^2_m$  and  $R^2_c$   
 286 are marginal and conditional  $R^2$ , respectively, and are calculated for the final model resulting from model  
 287 simplification.

Model tested	Predictors	Estimate ( $\pm$ SE)	$X^2$	Df	P-value	$R^2_m$ ( $R^2_c$ )
Model 1	$\Delta H$	0.96 ( $\pm$ 0.19)	87.07	1	< 0.001	0.23 (0.58)
	<b>plot type</b>	1.17 ( $\pm$ 0.32)	13.88	1	< 0.001	
	$\Delta H \times$ plot type	0.31 ( $\pm$ 0.25)	1.64	1	0.201	
Model 2	$\Delta H$	1.14 ( $\pm$ 0.12)	85.26	1	< 0.001	0.30 (0.57)
	<b>BA <i>P. sylvestris</i></b>	1.31 ( $\pm$ 0.43)	10.39	1	0.001	
	BA <i>P. pinaster</i>	0.50 ( $\pm$ 0.33)	1.52	1	0.217	
	BA <i>Ps</i> $\times$ BA <i>Pp</i>	0.38 ( $\pm$ 0.30)	1.61	1	0.205	
Model 3	$\Delta H$	1.18 ( $\pm$ 0.12)	90.05	1	< 0.001	0.25 (0.58)
	N <i>P. sylvestris</i>	0.17 ( $\pm$ 0.33)	0.01	1	0.931	
	<b>N <i>P. pinaster</i></b>	-0.33 ( $\pm$ 0.32)	11.15	1	< 0.001	
	N <i>Ps</i> $\times$ N <i>Pp</i>	0.51 ( $\pm$ 0.37)	1.89	1	0.169	

288

289

## 290 4. Discussion

291 We showed that the infestation level of Scots pines by *V. album* was almost twice as high in pure  
292 Scots pine plots compared to mixed plots of Scots pine and Maritime pine. Our study is the first to  
293 reveal and quantify associational resistance to a plant parasite in mixed vs. pure forest stands.  
294 Despite the correlative nature of our study, we can speculate that mechanisms proposed to explain  
295 tree diversity effects on resistance to insects and pathogens also apply to plant parasites. The  
296 observation that tree diversity reduces *V. album* infestation level may be related to i) changes in  
297 behavior or abundance of seed dispersing birds and/or to ii) changes in *V. album*-tree interactions.  
298 Both processes are potentially influenced by the density of the host tree (Scots pine) and the density  
299 of the associated, non-host species (Maritime pine).

300

#### 301 **4.1 Effect of host and non-host densities on mistletoe infestation**

302 We analyzed in the same model the effect of Scots pine and Maritime pine abundance on *V. album*  
303 infestation level. The use of number of trees indicated a pure associational effect of Maritime pine  
304 whereby *V. album* infestation decreased with increasing abundance of the non-host species, whereas  
305 analysis based on host and non-host basal area suggested an effect of host cover, whereby *V. album*  
306 infestation increased with increasing Scots pine basal area. These findings suggest that both the  
307 proportion and density of the host tree can account for the effect of mixture on *V. album* infestation.  
308 However, because the experimental plots were based on a replacement of one species by the other  
309 and the number of trees or the basal area of the two species were correlated negatively, we could  
310 not quantify the relative importance of these two mechanisms. To demonstrate a pure associational  
311 effect one should compare plots with the same Scots pine density but with absence or presence of  
312 Maritime pines (Damien et al., 2016; Hambäck et al., 2014). Concerning the effect of Scots pine  
313 abundance in our plots it seems that the *V. album* infestation depended more on Scots pine basal  
314 area than on the number of Scots pine trees, possibly because seed dispersing birds, such as Mistle  
315 trushes, may react more to the species space occupancy in the stand (particularly crown surface  
316 where birds land) than to the number of trees. Kolodziejek and Kolodziejek (2013) observed in

317 Poland, in pure Scots pine stands, a higher prevalence of *V. album* in low density stands compared to  
318 high density stands. This pattern corresponds to the resource dilution hypothesis where infestations  
319 are more concentrated on a more diluted resource of host trees (Otway et al., 2005). Likewise,  
320 Mellado and Zamora (2016) showed an increase in visits of frugivorous birds and *V. album* seed  
321 abundance in lower density *Pinus nigra* stands. However, effects of tree density, basal area, crown  
322 cover and tree height may have been confounded in this or other studies, which complicates their  
323 interpretation (Donohue, 1995; Kolodziejek and Kolodziejek, 2013). Low tree density can correspond  
324 to taller trees with a larger crown affecting possibly bird behavior. Moreover, Kolodziejek and  
325 Kolodziejek (2013) studied pure Scots pine stands whereas our results are based on pure and mixed  
326 stands, explaining that we did not observe the same pattern. We therefore encourage future studies  
327 to uncouple the effects of stand density and tree dimensions to move the understanding of  
328 associational effects on mistletoe toward a more mechanistic framework.

329

#### 330 **4.2 Mistletoe infestation increased with relative host size**

331 We showed that Scots pine trees that were higher than the surrounding trees had a higher *V. album*  
332 infestation probability, both in mixed and pure plots. Many studies have shown a higher infestation  
333 by mistletoe species in taller trees (Aukema and Martínez del Rio, 2002; Donohue, 1995; Kolodziejek  
334 and Kolodziejek, 2013; Norton et al., 1997; Roxburgh and Nicolson, 2008; Shaw et al., 2005; Smith  
335 and Reid, 2000; Teodoro et al., 2010) and some could attribute this effect to preferences of birds for  
336 visiting taller trees, either in open landscape or forest (Aukema and Martínez del Rio, 2002; Monteiro  
337 et al., 1992; Roxburgh and Nicolson, 2008). In our mixed plots, Scots pines were slightly lower than  
338 Maritime pines, making them possibly less attractive for birds and thus leading to a lower seed  
339 deposition on Scots pines in mixed stands compared to pure stands. Reduced host apparency is a  
340 pure associational effect that has been found to diminish insect attacks on trees (Castagneyrol et al.,  
341 2013; Damien et al., 2016; Dulaurent et al., 2012) and can thus likewise reduce *V. album* seed  
342 deposition by birds on partially hidden trees. Taller trees may not only be more apparent to birds,

343 they also offer a larger crown surface to land on, which could increase their infestation probability.

344 Taller trees may also be a more suitable host for *V. album*, which is a light demanding species (Zuber,  
345 2004). *V. album* survival may be thus better in dominant, sun-exposed trees. Taller trees, in the same  
346 taxon, may also provide a more reliable water supply because of their deeper rooting system and  
347 thereby offering a higher survival to mistletoe species (Norton et al., 1997; Roxburgh and Nicolson,  
348 2008).

349 Additionally, we showed that for the same relative tree height in a considered stand, individual Scots  
350 pine trees had a lower infestation probability in mixed stands than in pure stands, indicating that  
351 other mechanisms than relative tree height play a role for the observed lower infestation level in  
352 mixed stands. In mixed stands birds may land on Scots pines and Maritime pines and a part of the  
353 seeds will be dropped and thus lost on Maritime pine. Future studies on bird behavior in relation to  
354 host proportion may show if this mechanism is important. Pure Scots pine stands also represent  
355 areas with higher *V. album* densities for birds, as host tree density is higher and trees have a higher  
356 infestation level than in mixed stands. Birds feeding on *V. album* may stay longer or be more  
357 abundant in pure Scots pine stands where they can find a higher amount of resources (Skórka and  
358 Wójcik, 2005; Snow and Snow, 1984; Telleria et al., 2008;2014), thereby increasing seed deposition in  
359 already infested stands. Aukema (2003) and Martinez del Rio et al. (1996), documented a local  
360 aggregation of mistletoe for respectively a desert mistletoe in North America and a cactus mistletoe  
361 in Chili. They could link this pattern with bird behavior as the percentage of non-parasitized hosts  
362 receiving seeds increased with the percentage of mistletoe-infested hosts in the neighborhood. This  
363 created a positive feedback as infected neighborhoods become even more heavily infected. Likewise,  
364 Morales et al. (2012) showed, for a mistletoe species dispersed by a marsupial, a reduction in seed  
365 dispersal distances when the neighborhood had a high mistletoe density.

366 Altogether, it seems probable that the observed higher mistletoe infestation in pure Scots pine  
367 stands is related to bird preferences for apparent trees and for areas with higher infestation levels.  
368 However, we cannot exclude that physiological (like chemical defenses, e.g. Lazaro-Gonzalez et al.



369 2019) or anatomical traits of Scots pines (e.g. bark thickness) may be different between pure and  
370 mixed stands and that these traits could explain the rate of mistletoe establishment and growth.  
371 Further research is therefore needed to determine which mechanism is most important.

372

### 373 **4.3 Consequences for forest management**

374 Since high densities of *V. album* reduce tree growth and contribute to tree mortality, different  
375 methods for controlling this parasitic plant have been proposed. The most effective one is  
376 mechanical control, such as pruning of infested branches, or removing infested trees (Varga et al.,  
377 2012). This may be applicable in infested orchards, but seems less applicable in extensive forests with  
378 tall trees. Moreover, removing infested trees may render remaining host trees more prone to  
379 infestation (Vallauri, 1998). However, this may not be the case in our mixed stands where lower host  
380 abundance seems to decrease infestation level. We showed that in mixed pine stands the infestation  
381 level of Scots pine was on average 44 % lower compared to pure stands. Conservation pest  
382 management, that is the use of tree diversity to keep *V. album* infestation at a low level, has to our  
383 knowledge only been tested by Oliva & Colinas (2010), who showed that *Abies* stands with a low  
384 level of *V. album* infestation had a higher proportion of accompanying tree species than stands with  
385 a high infestation level. However, they observed no differences between highly infested and non-  
386 infested stands, probably because of confounding factors for the non-infested stands.

387 Management of tree species diversity in forest stands for associational resistance shows several  
388 advantages. It not only allows diminishing the negative effects of *V. album* on tree growth and  
389 mortality, but may also permit an overall higher stand productivity (Riofrío et al., 2016; 2017). Lower  
390 *V. album* infestation can be even one of the factors related to higher productivity in mixed stands  
391 and would merit further research.

392

### 393 **4.4 Conclusion and perspectives**

394 We showed that tree diversity can reduce Scots pine infestation by a plant parasite, the mistletoe *V.*  
395 *album*. Although literature on mixed forest resistance to herbivorous insects and fungal pathogens  
396 may help to identify possible mechanisms underlying mixed forest resistance to this plant parasite,  
397 further dedicated research is needed to clarify them. In particular, mistletoe is actively dispersed by  
398 birds. Studies on bird behavior and abundance in relation to stand composition and *V. album*  
399 infestation level may allow to precise their role in the observed reduced infestation in mixed stands.  
400 Moreover, it will be useful to evaluate the effect of different tree species mixtures on *V. album*  
401 infestation levels and in different regions, as host preferences of *V. album* may vary regionally. This  
402 will also allow generalizing our results and recommendations to other forest systems and ecological  
403 conditions.

404

405 **Funding:** This work was supported by the Spanish Ministry of Economy and Competitiveness  
406 (projects PCIN-2017-026 and PCIN-2017-027) and the French National Research Agency (ANR-16-  
407 SUMF-0003-01) by funding the national contribution to the Sumforest ERA-net project REFORM  
408 (Resilience of Forest Mixtures).

409

## 410 **Author Contributions**

411 IVH and HJ conceived the mistletoe study. MdR and FB set up the forest triplets, and CO measured  
412 the trees. IVH, HJ and LP performed the mistletoe field survey, IVH and BC analyzed the data, IVH  
413 drafted the first version of the manuscript. All authors contributed to the writing of the present  
414 version of the manuscript.

415

## 416 **Appendix A. Supplementary material**

417 Supplementary material to this article can be found online at .....

418

## 419 **References**

- 420 Aukema, J.E., 2003. Vectors, viscin, and Viscaceae: mistletoes as parasites, mutualists, and resources.  
421 Front. Ecol. Environ. 1, 212–219. [https://doi.org/10.1890/1540-](https://doi.org/10.1890/1540-9295(2003)001[0212:VVAVMA]2.0.CO;2)  
422 [9295\(2003\)001\[0212:VVAVMA\]2.0.CO;2](https://doi.org/10.1890/1540-9295(2003)001[0212:VVAVMA]2.0.CO;2)
- 423 Aukema, J.E., Martínez del Rio, C., 2002. Where does a fruit-eating bird deposit mistletoe seeds?  
424 Seed deposition patterns and an experiment. Ecology 83, 3489–3496.  
425 [https://doi.org/10.1890/0012-9658\(2002\)083\[3489:WDAFEB\]2.0.CO;2](https://doi.org/10.1890/0012-9658(2002)083[3489:WDAFEB]2.0.CO;2)
- 426 Barbosa, P., Hines, J., Kaplan, I., Martinson, H., Szczepaniec, A., Szendrei, Z., 2009. Associational  
427 Resistance and Associational Susceptibility: having right or wrong neighbors. Annu. Rev. Ecol.  
428 Evol. Syst. 40, 1–20. <https://doi.org/10.1146/annurev.ecolsys.110308.120242>.
- 429 Barton, K., 2018. MuMIn: Multi-Model Inference. R package version 1.42.1. [https://CRAN.R-](https://CRAN.R-project.org/package=MuMIn)  
430 [project.org/package=MuMIn](https://CRAN.R-project.org/package=MuMIn)
- 431 Bates, D., Mächler, M., Bolker, B., Walker, S., 2015. Fitting Linear Mixed-Effects Models using lme4. J.  
432 Stat. Softw. 67, 1–48. <https://doi.org/10.18637/jss.v067.i01>
- 433 Briggs, J., 2011. Mistletoe (*Viscum album*): A brief review of its local status with recent observations  
434 on its insects associations and conservation problems. Proc Cotteswold Nat. Fld Club 45,  
435 181–193.
- 436 Brockerhoff, E.G., Barbaro, L., Castagneyrol, B., Forrester, D.I., Gardiner, B., González-Olabarria, J.R.,  
437 Lyver, P.O., Meurisse, N., Oxbrough, A., Taki, H., Thompson, I.D., van der Plas, F., Jactel, H.,  
438 2017. Forest biodiversity, ecosystem functioning and the provision of ecosystem services.  
439 Biodivers. Conserv. 26, 3005–3035. <https://doi.org/10.1007/s10531-017-1453-2>
- 440 Castagneyrol, B., Giffard, B., Péré, C., Jactel, H., 2013. Plant apparency, an overlooked driver of  
441 associational resistance to insect herbivory. J. Ecol. 101, 418–429.  
442 <https://doi.org/10.1111/1365-2745.12055>

443 Castagneyrol, B., Jactel, H., Moreira, X., 2018. Anti-herbivore defences and insect herbivory:  
444 Interactive effects of drought and tree neighbours. *J. Ecol.* 106, 2043-2057.  
445 <https://doi.org/10.1111/1365-2745.12956>

446 Castagneyrol, B., Jactel, H., Vacher, C., Brockerhoff, E.G., Koricheva, J., 2014. Effects of plant  
447 phylogenetic diversity on herbivory depend on herbivore specialization. *J. Appl. Ecol.* 51,  
448 134–141. <https://doi.org/10.1111/1365-2664.12175>

449 Damien, M., Jactel, H., Meredieu, C., Régolini, M., van Halder, I., Castagneyrol, B., 2016. Pest damage  
450 in mixed forests: Disentangling the effects of neighbor identity, host density and host  
451 apparency at different spatial scales. *For. Ecol. Manag.* 378, 103–110.  
452 <https://doi.org/10.1016/j.foreco.2016.07.025>

453 Dobbertin, M., Hilker, N., Rebetez, M., Zimmermann, N.E., Wohlgemuth, T., Rigling, A., 2005. The  
454 upward shift in altitude of pine mistletoe (*Viscum album* ssp. *austriacum*) in Switzerland—the  
455 result of climate warming? *Int. J. Biometeorol.* 50, 40–47. [https://doi.org/10.1007/s00484-](https://doi.org/10.1007/s00484-005-0263-5)  
456 [005-0263-5](https://doi.org/10.1007/s00484-005-0263-5)

457 Dobbertin, M., Rigling, A., 2006. Pine mistletoe (*Viscum album* ssp. *austriacum*) contributes to Scots  
458 pine (*Pinus sylvestris*) mortality in the Rhone valley of Switzerland. *For. Pathol.* 36, 309–322.

459 Donohue, K., 1995. The Spatial Demography of Mistletoe Parasitism on a Yemeni Acacia. *Int. J. Plant*  
460 *Sci.* 156, 816–823. <https://doi.org/10.1086/297305>

461 Dulaurent, A.-M., Porté, A.J., van Halder, I., Vétillard, F., Menassieu, P., Jactel, H., 2012. Hide and seek  
462 in forests: colonization by the pine processionary moth is impeded by the presence of  
463 nonhost trees. *Agric. For. Entomol.* 14, 19–27. [https://doi.org/10.1111/j.1461-](https://doi.org/10.1111/j.1461-9563.2011.00549.x)  
464 [9563.2011.00549.x](https://doi.org/10.1111/j.1461-9563.2011.00549.x)

465 Durand-Gillmann, M., Cailleret, M., Boivin, T., Nageleisen, L.-M., Davi, H., 2014. Individual  
466 vulnerability factors of Silver fir (*Abies alba* Mill.) to parasitism by two contrasting biotic  
467 agents: mistletoe (*Viscum album* L. ssp. *abietis*) and bark beetles (Coleoptera: Curculionidae:

468 Scolytinae) during a decline process. *Ann. For. Sci.* 71, 659–673.  
469 <https://doi.org/10.1007/s13595-012-0251-y>

470 Forrester, D.I., Bauhus, J., 2016. A review of processes behind diversity—productivity relationships in  
471 forests. *Curr. For. Rep.* 2, 45–61. <https://doi.org/10.1007/s40725-016-0031-2>

472 Fox, J., Weisberg, S., 2011. *An R Companion to Applied Regression*, Second Edition. Thousand Oaks  
473 CA: Sage.

474 Frochot, H., Sallé, G., 1980. Modalités de dissémination et d’implantation du gui. *R.F.F.* 32, 505–519.

475 Grossiord, C., 2019. Having the right neighbors: how tree species diversity modulates drought  
476 impacts on forests. *New Phytol.* <https://doi.org/10.1111/nph.15667>

477 Hambäck, P.A., Inouye, B.D., Andersson, P., Underwood, N., 2014. Effects of plant neighborhoods on  
478 plant–herbivore interactions: resource dilution and associational effects. *Ecology* 95, 1370–  
479 1383.

480 Hartig, F., 2019. DHARMA: residual diagnostics for hierarchical (multi-level/mixed) regression models.  
481 R package version 0.2.4. <https://CRAN.R-project.org/package=DHARMA>

482 Jactel, H., Bauhus, J., Boberg, J., Bonal, D., Castagneyrol, B., Gardiner, B., Gonzalez-Olabarria, J.R.,  
483 Koricheva, J., Meurisse, N., Brockerhoff, E.G., 2017. Tree diversity drives forest stand  
484 resistance to natural disturbances. *Curr. For. Rep.* 3, 223–243.  
485 <https://doi.org/10.1007/s40725-017-0064-1>

486 Jactel, H., Gritti, E., Drössler, L., Forrester, D., L. Mason, W., Morin, X., Pretzsch, H., Castagneyrol, B.,  
487 2018. Positive biodiversity–productivity relationships in forests: Climate matters. *Biol. Lett.*  
488 14, 20170747. <https://doi.org/10.1098/rsbl.2017.0747>

489 Jactel, H., Brockerhoff, Eckehard G., 2007. Tree diversity reduces herbivory by forest insects. *Ecol.*  
490 *Lett.* 10, 835–848. <https://doi.org/10.1111/j.1461-0248.2007.01073.x>

491 Kolodziejek, J., Kolodziejek, A., 2013. The spatial distribution of pine mistletoe *Viscum album ssp.*  
492 *austriacum* (Wiesb.) Volmann in a scots pine (*Pinus sylvestris* L.) stand in Central Poland. *Pol.*  
493 *J. Ecol.* 61, 705–714.

494 Lázaro-González, A., Hódar, J.A., Zamora, R., 2017. Do the arthropod communities on a parasitic plant  
495 and its hosts differ? *Eur. J. Entomol.* 114, 215–221.

496 Lázaro-González, A., Hódar, J.A., Zamora, R., 2019. Mistletoe Versus Host Pine: Does Increased  
497 Parasite Load Alter the Host Chemical Profile? *J. Chem. Ecol.* 45, 95–105.  
498 <https://doi.org/10.1007/s10886-018-1039-9>.

499 Martín, S., Díaz-Fernández, P., de Miguel, J., 1998. Regiones de procedencia de especies forestales  
500 españolas. Organismo Autónomo Parques Nacionales.

501 Martínez del Río, C., Silva, A., Medel, R., Hourdequin, M., 1996. Seed dispersers as disease vectors:  
502 bird transmission of mistletoe seeds to plant hosts. *Ecology* 77, 912–921.

503 Mellado, A., Zamora, R., 2016. Spatial heterogeneity of a parasitic plant drives the seed-dispersal  
504 pattern of a zoochorous plant community in a generalist dispersal system. *Funct. Ecol.* 30,  
505 459–467. <https://doi.org/10.1111/1365-2435.12524>

506 Mellado, A., Zamora, R., 2014. Generalist birds govern the seed dispersal of a parasitic plant with  
507 strong recruitment constraints. *Oecologia* 176, 139–147. [https://doi.org/10.1007/s00442-](https://doi.org/10.1007/s00442-014-3013-8)  
508 014-3013-8

509 Monteiro, R.F., Martins, R.P., Yamamoto, K., 1992. Host specificity and seed dispersal of  
510 *Psittacanthus robustus* (Loranthaceae) in south-east Brazil. *J. Trop. Ecol.* 8, 307–314.  
511 <https://doi.org/10.1017/S026646740000657X>

512 Morales, J.M., Rivarola, M.D., Amico, G., Carlo, T.A., 2012. Neighborhood effects on seed dispersal by  
513 frugivores: testing theory with a mistletoe–marsupial system in Patagonia. *Ecology* 93, 741–  
514 748. <https://doi.org/10.1890/11-0935.1>

515 Mutlu, S., Osmar, E., İlhan, V., Turkoglu, H.I., Atici, O., 2016. Mistletoe (*Viscum album*) reduces the  
516 growth of the Scots pine by accumulating essential nutrient elements in its structure as a  
517 trap. *Trees* 30, 815–824. <https://doi.org/10.1007/s00468-015-1323-z>

518 Nakagawa, S., Schielzeth, H., 2013. A general and simple method for obtaining  $R^2$  from generalized  
519 linear mixed-effects models. *Methods Ecol. Evol.* 4, 133–142. <https://doi.org/10.1111/j.2041->  
520 [210x.2012.00261.x](https://doi.org/10.1111/j.2041-210x.2012.00261.x)

521 Noetzli, K.P., Müller, B., Sieber, T.N., 2003. Impact of population dynamics of white mistletoe (*Viscum*  
522 *album* ssp. *abietis*) on European silver fir (*Abies alba*). *Ann. For. Sci.* 60, 773–779.  
523 <https://doi.org/10.1051/forest:2003072>

524 Norton, D.A., Ladley, J.J., Owen, H.J., 1997. Distribution and population structure of the  
525 loranthaceous mistletoes *Alepis flavida*, *Peraxilla colensoi*, and *Peraxilla tetrapetala* within  
526 two New Zealand *Nothofagus* forests. *N. Z. J. Bot.* 35, 323–336.  
527 <https://doi.org/10.1080/0028825X.1997.10410158>

528 Oliva, J., Colinas, C., 2010. Epidemiology of *Heterobasidion abietinum* and *Viscum album* on silver fir  
529 (*Abies alba*) stands of the Pyrenees. *For. Pathol.* 40, 19–32. <https://doi.org/10.1111/j.1439->  
530 [0329.2009.00603.x](https://doi.org/10.1111/j.1439-0329.2009.00603.x)

531 Otway, S.J., Hector, A., Lawton, J.H., 2005. Resource dilution effects on specialist insect herbivores in  
532 a grassland biodiversity experiment. *J. Anim. Ecol.* 74, 234–240.

533 R Core Team, 2019. R: A language and environment for statistical computing. R Foundation for  
534 Statistical Computing, Vienna, Austria. URL <https://www.R-project.org/>.

535 Rigling, A., Eilmann, B., Koechli, R., Dobbertin, M., 2010. Mistletoe-induced crown degradation in  
536 Scots pine in a xeric environment. *Tree Physiol.* 30, 845–852.  
537 <https://doi.org/10.1093/treephys/tpq038>.

538 Riofrío, J., del Río, M., Bravo, F., 2016. Mixing effects on growth efficiency in mixed pine forests.  
539 *Forestry* 2016, 90, 381-392.

540 Riofrío, J., del Río, M., Pretzsch, H., Bravo, F., 2017. Changes in structural heterogeneity and stand  
541 productivity by mixing Scots pine and Maritime pine. *For. Ecol. Manag.* 405, 219–228.  
542 <https://doi.org/10.1016/j.foreco.2017.09.036>

543 Root, R.B., 1973. Organization of a plant-arthropod association in simple and diverse habitats: the  
544 fauna of collards (*Brassica oleracea*). Ecol. Monogr. 43, 95–124.

545 Rosado-Sánchez, S., Parra-Tabla, V., Betancur-Ancona, D., Moreira, X., Abdala-Roberts, L., 2018. Tree  
546 species diversity alters plant defense investment in an experimental forest plantation in  
547 southern Mexico. Biotropica 50, 246–253.

548 Roxburgh, L., Nicolson, S.W., 2008. Differential dispersal and survival of an African mistletoe: does  
549 host size matter? Plant Ecol. 195, 21–31. <https://doi.org/10.1007/s11258-007-9295-8>

550 Sangüesa-Barreda, G., Linares, J.C., Julio Camarero, J., 2013. Drought and mistletoe reduce growth  
551 and water-use efficiency of Scots pine. For. Ecol. Manag. 296, 64–73.  
552 <https://doi.org/10.1016/j.foreco.2013.01.028>

553 Schielzeth, H., Nakagawa, S., 2013. Nested by design: model fitting and interpretation in a mixed  
554 model era. Methods Ecol. Evol. 4, 14–24.

555 Shaw, D.C., Chen, J., Freeman, E.A., Braun, D.M., 2005. Spatial and population characteristics of  
556 dwarf mistletoe infected trees in an old-growth Douglas-fir western hemlock forest. Can. J.  
557 For. Res. 35, 990–1001.

558 Skórka, P., Wójcik, J.D., 2005. Population Dynamics and Social Behavior of the Mistle Thrush *Turdus*  
559 *viscivorus* During Winter. Acta Ornithol. 40, 35–42. <https://doi.org/10.3161/068.040.0109>

560 Smith, M.S., Reid, N., 2000. Population dynamics of an arid zone mistletoe (*Amyema preissii*,  
561 Loranthaceae) and its host *Acacia victoriae* (Mimosaceae). Aust. J. Bot. 48, 45–58.  
562 <https://doi.org/10.1071/bt97076>

563 Snow, B.K., Snow, D.W., 1984. Long-term defence of fruit by Mistle Thrushes (*Turdus viscivorus*). Ibis  
564 126, 39–49.

565 Telleria, J.L., Ramirez, A., Perez Tris, J., 2008. Fruit tracking between sites and years by birds in  
566 Mediterranean wintering grounds. Ecography 31:381–388. doi:10.1111/j.0906-  
567 7590.2008.05283.x



568 Tellería, J.L., Carrascal, L.M., Santos, T., 2014. Species abundance and migratory status affects large-  
569 scale fruit tracking in thrushes (*Turdus* spp.). *J. Ornithol.* 155, 157–164.  
570 <https://doi.org/10.1007/s10336-013-0997-5>

571 Teodoro, G.S., van den Berg, E., de Castro Nunes Santos, M., de Freitas Coelho, F., 2010. How does a  
572 *Psittacanthus robustus* Mart. population structure relate to a *Vochysia thyrsoidea* Pohl. host  
573 population? *Flora - Morphol. Distrib. Funct. Ecol. Plants* 205, 797–801.  
574 <https://doi.org/10.1016/j.flora.2010.04.013>

575 Tsopelas, P., Angelopoulos, A., Economou, A., Soulioti, N., 2004. Mistletoe (*Viscum album*) in the fir  
576 forest of Mount Parnis, Greece. *For. Ecol. Manag.* 202, 59–65.  
577 <https://doi.org/10.1016/j.foreco.2004.06.032>

578 Vallauri, D., 1998. Dynamique parasitaire de *Viscum album* L. sur pin noir dans le bassin du Saignon  
579 (préalpes françaises du sud). *Ann. Sci. For.* 55, 823–835.  
580 <https://doi.org/10.1051/forest:19980706>

581 Varga, I., Taller, J., Baltazár, T., Hyvönen, J., Poczai, P., 2012. Leaf-spot disease on European mistletoe  
582 (*Viscum album*) caused by (*Phaeobotryosphaeria visci*): a potential candidate for biological  
583 control. *Biotechnol. Lett.* 34, 1059–1065. <https://doi.org/10.1007/s10529-012-0867-x>

584 Watson, D.M., 2009. Determinants of parasitic plant distribution: the role of host quality. *Botany* 87,  
585 16–21. <https://doi.org/10.1139/B08-105>

586 Watson, D.M., 2001. Mistletoe—a keystone resource in forests and woodlands worldwide. *Annu.*  
587 *Rev. Ecol. Syst.* 32, 219–249.

588 Zuber, D., 2004. Biological flora of Central Europe: *Viscum album* L. *Flora* 199, 181–203.  
589 <https://doi.org/10.1078/0367-2530-00147>

590 Zuber, D., Widmer, A., 2009. Phylogeography and host race differentiation in the European mistletoe  
591 (*Viscum album* L.). *Mol. Ecol.* 18, 1946–1962. [https://doi.org/10.1111/j.1365-](https://doi.org/10.1111/j.1365-294X.2009.04168.x)  
592 [294X.2009.04168.x](https://doi.org/10.1111/j.1365-294X.2009.04168.x)

593

594 **Figure captions:**

595

596 **Fig. 1.** Possible effects of tree diversity on mistletoe (*Viscum album*) infestation. The solid arrows  
597 represent the mistletoe cycle (seed consumption and dispersion by birds, seed germination and plant  
598 establishment on the host tree, here Scots pine). The dashed arrows represent possible effects of a  
599 non-host tree species (here Maritime pine) on mistletoe infestation of Scots pines growing in a mixed  
600 stand.

601

602 **Fig. 2.** Mean percentage ( $\pm$ SE) of Scots pines infested with *V. album* in mixed vs. pure plots.

603

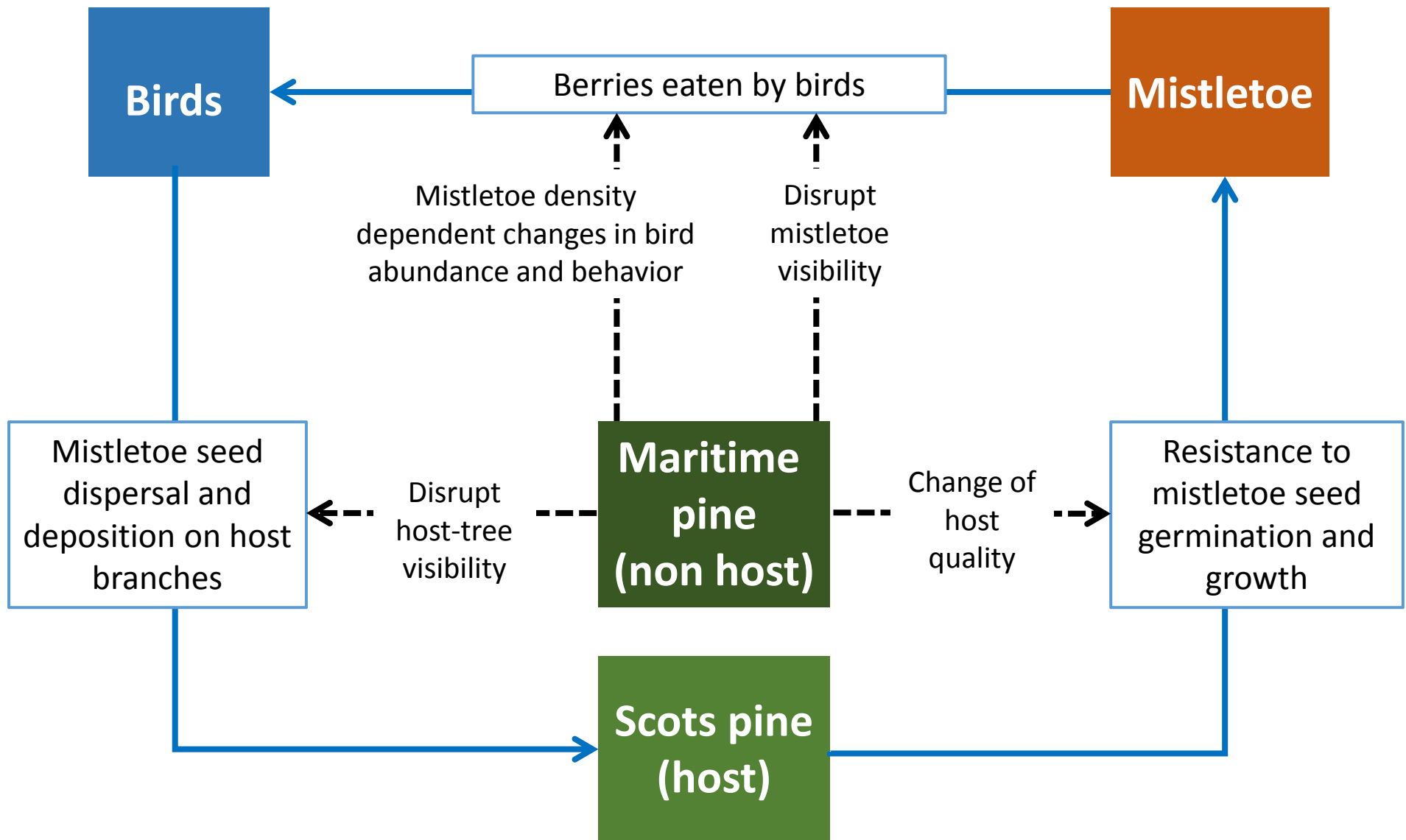
604

605 **Fig. 3.** Effect of individual relative tree height ( $\Delta H$ ), which indicates how much taller or lower an  
606 individual *P. sylvestris* tree is as compared to its neighbors within the plot, in mixed and pure plots,  
607 on the probability of individual *P. sylvestris* trees being infested by *Viscum album* (i.e. model 1 of  
608 Table 3). The dashed vertical line at  $\Delta H = 0$  indicates the cases in which *P. sylvestris* are on average as  
609 tall as the other trees in the plot. Light green and dark green vertical bars at  $y = 0$  and  $y = 1$  represent  
610 observed  $\Delta H$  in mixed and pure stands, respectively. Logistic curves represent predictions from  
611 models (solid lines) and their standard errors (dashed lines).

612

613

614



% infested *P. sylvestris* per plot

