

AperTO - Archivio Istituzionale Open Access dell'Università di Torino

## What drives European beech (*Fagus sylvatica* L.) mortality after forest fires of varying severity?

### This is the author's manuscript

*Original Citation:*

*Availability:*

This version is available <http://hdl.handle.net/2318/1565756> since 2016-06-13T10:17:29Z

*Published version:*

DOI:10.1016/j.foreco.2016.03.008

*Terms of use:*

Open Access

Anyone can freely access the full text of works made available as "Open Access". Works made available under a Creative Commons license can be used according to the terms and conditions of said license. Use of all other works requires consent of the right holder (author or publisher) if not exempted from copyright protection by the applicable law.

(Article begins on next page)

Manuscript Number: FORECO16157

Title: What drives European beech (*Fagus sylvatica* L.) to death after forest fires of varying severities?

Article Type: FLA Full Length Article

Keywords: beech fire ecology; fungal infestation; wood decay; Southern European Alps

Corresponding Author: Mrs. Janet Maringer,

Corresponding Author's Institution: University of Stuttgart

First Author: Janet Maringer

Order of Authors: Janet Maringer; Davide Ascoli, PhD; Nicolas Küffer, PhD; Sebastian Schmidlein, Prof. ; Marco Conedera, PhD

Abstract: Predicting the timing and the amount of tree mortality after a forest fire is of paramount importance for forest management decisions related to post-fire measures. Such knowledge is particularly needed in mountainous regions where forest stands often serve as protection against natural hazards. In this paper, we focus on drivers and temporal trends in mortality processes of fire-injured beeches (*Fagus sylvatica* L.). We studied beech forests in the Southern European Alps, which burnt between 1970 and 2012. The results show that beech trees—which obviously lack fire-resistance traits—experience increased mortality within the first two decades post-fire after fires of moderate and high burn severity. Beech mortality was ubiquitous in high severity sites, whereas taller beeches had a higher survivability after fires of moderate severity, representing a crucial ecological legacy for natural beech regeneration. In contrast, mortality remains low and at a level similar to unburnt beech forests for low burn severity sites. Among mortality drivers, fungal infestation and the presence of fungal sporulation in particular was found to be most significant. Interestingly, fungi initiating post-fire tree mortality appeared to differ from fungi occurring after mechanical injury. From a practical point of view, insight gained regarding post-fire mortality is of great help in planning post-fire silvicultural measures.

Suggested Reviewers: Stefanie Gärtner PhD  
stefanie.gaertner@waldbau.uni-freiburg.de

Tibor Standovár PhD  
standy@ludens.elte.hu

Francisco Moreira PhD  
fmoreira@isa.ulisboa.pt

Tara L. Keyser  
tkeyser@fs.fed.us



Janet Maringer  
Institute of Landscape planning and ecology  
Keplerstr. 11  
D- 70174 Stuttgart  
Germany  
jm@ilpoe.uni-stuttgart.de

Stuttgart, 17<sup>th</sup> October 2015

Dear Editor,

We wish to submit an original research article entitled “What drives lead to European beech (*Fagus sylvatica* L.) mortality after forest fires of varying severities?” for consideration by the journal Forest Ecology and Management.

We confirm that this work is original and has not been published elsewhere, nor is it currently under consideration for publication elsewhere. The article has been read and approved by all authors.

In this paper, we report on the timing and drivers of beech mortality after forest fires of varying severity. This is of importance with regards to forest management decisions related to post-fire measures. Such knowledge is particularly needed in mountainous regions where forest stands often serve as protection against natural hazards.

We believe this topic is appropriate for publication by Forest Ecology and Management because it links fire-ecological processes with forest management decisions, and makes further contributions to a broader understanding of beech fire ecology. A parallel study focusing on post-fire regeneration processes after a mast year has recently been published in the journal of Forest Ecology and Management (doi:10.1016/j.foreco.2015.05.031).

We have no conflicts of interest to disclose. If you find that the manuscript is appropriate for your journal, we suggest the following reviewers:

- Dr. Stefanie Gärtner, [stefanie.gaertner@waldbau.uni-freiburg.de](mailto:stefanie.gaertner@waldbau.uni-freiburg.de)
- Prof. Dr. Tibor Standovár, [standy@ludens.elte.hu](mailto:standy@ludens.elte.hu)
- Francisco Moreira, [fmoreira@isa.ulisboa.pt](mailto:fmoreira@isa.ulisboa.pt)
- Tara L. Keyser, [tkeyser@fs.fed.us](mailto:tkeyser@fs.fed.us)

Please address all correspondence concerning this manuscript to me at [jm@ilpoe.uni-stuttgart.de](mailto:jm@ilpoe.uni-stuttgart.de)

I thank you and the reviewers in advance for your efforts in evaluating and reviewing this manuscript. I look forward to your review.

Sincerely,

Janet Maringer



21 **Abstract**

22 Predicting the timing and the amount of tree mortality after a forest fire is of  
23 paramount importance for forest management decisions related to post-fire measures.  
24 Such knowledge is particularly needed in mountainous regions where forest stands  
25 often serve as protection against natural hazards. In this paper, we focus on drivers  
26 and temporal trends in mortality processes of fire-injured beeches (*Fagus sylvatica*  
27 L.). We studied beech forests in the Southern European Alps, which burnt between  
28 1970 and 2012. The results show that beech trees—which obviously lack fire-  
29 resistance traits—experience increased mortality within the first two decades post-fire  
30 after fires of moderate and high burn severity. Beech mortality was ubiquitous in high  
31 severity sites, whereas taller beeches had a higher survivability after fires of moderate  
32 severity, representing a crucial ecological legacy for natural beech regeneration. In  
33 contrast, mortality remains low and at a level similar to unburnt beech forests for low  
34 burn severity sites.

35 Among mortality drivers, fungal infestation and the presence of fungal sporulation in  
36 particular was found to be most significant. Interestingly, fungi initiating post-fire tree  
37 mortality appeared to differ from fungi occurring after mechanical injury. From a  
38 practical point of view, insight gained regarding post-fire mortality is of great help in  
39 planning post-fire silvicultural measures.

40 **Keywords:** beech fire ecology; fungal infestation; wood decay; Southern European  
41 Alps

42

## 43 **1 Introduction**

44 Climate change and the related predictions of a warmer and drier climate (IPCC,  
45 2014) lead to increasing concerns about the future impact of wildfires on forest  
46 resistance and resilience in both fire-prone and less fire-prone forest ecosystems  
47 (Bachelet et al., 2007; Fischer et al., 2010; Schumacher and Bugmann, 2006). In fire-  
48 prone regions, the size and intensity of wildfires have already increased in recent  
49 decades (e.g., Westerling et al., 2006; Sullivan et al., 2011; Sarris et al., 2014). This  
50 has raised questions on how to predict fire-injured tree-mortality rates in the  
51 framework of planning post-fire restoration measures (Brown et al., 2003; Ledgard  
52 and Davis, 2004; Kobziar et al., 2006; Keyser et al., 2008; Moreira et al., 2012).  
53 Models explaining driving factors and predicting post-fire tree mortality have been  
54 developed mainly for fire-prone ecosystems (e.g., McHugh and Kolb, 2003; Ledgard  
55 and Davis, 2004; Rigolot, 2004; Kobziar et al., 2006; Sieg et al., 2006; Hood et al.,  
56 2007; Fernandes et al., 2008; Stevens-Rumann et al., 2012). Comparatively little  
57 attention has been paid to tree species that dominate in less fire-prone regions. From a  
58 forest management perspective, one of the major problems arises from the lack of  
59 data and experience regarding the vulnerability and resilience of such forest stands  
60 under increasing fire disturbances.

61 European beech (*Fagus sylvatica* L.), for example, represents a tree species with high  
62 economic and ecological value in Europe whose forest stands are usually considered  
63 less prone to fire (Pezzatti et al., 2013). However, during the exceptional drought in  
64 2003 (e.g. Beniston, 2004), beech stands in the Southern Alps experienced numerous  
65 and atypically large forest fires, indicating a potential shift in fire regime together  
66 with climate change (Valese et al., 2014). To date, the species' post-fire survival  
67 strategies are poorly understood. Mature beeches are generally considered highly

68 susceptible to fire because of a lack of fire resistance or fire-adaptation traits, such as  
69 thick bark or persistent resprouting capability (Peters, 1997; Packham et al., 2012).  
70 Furthermore, beech relies on gravity and rodent seed dispersal, which makes its  
71 regeneration limited by the distance to the nearest seed-bearing tree or by the distance  
72 to the forest edge (Wagner et al., 2010; van Couwenberghe et al., 2010).  
73 Consequently, the greater the burnt area and the further away an off-site seed source  
74 is, the more limited natural beech regeneration may be. Recent studies suggest,  
75 however, that beech stands exhibit surprisingly high resilience after single fire events  
76 (Ascoli et al., 2013; Maringer et al., *subm.*). This fire surviving strategy is mainly  
77 based on rapid *in situ* seed production when mast years coincide with suitable  
78 germination conditions in the post-fire environment (e.g., improved light conditions  
79 and reduced litter cover on the soil, Ascoli et al., 2015). Thus, post-fire density and  
80 spatial distribution of surviving seed trees are of paramount importance for new  
81 cohort recruitments and represent the basis for the rapid recovery of beech forests.

82 It is well known that the timing of post-fire beech mortality depends on fire intensity.  
83 Beech mortality may occur immediately after very severe fires or be delayed by  
84 several years after low to moderate fire severity (Conedera et al., 2007; Ascoli et al.,  
85 2013). There is, however, a lack of knowledge regarding factors that drive such  
86 delayed post-fire beech mortality, and with respect to the predictability of its timing.  
87 Such information would support forest managers in making decisions related to post-  
88 fire measures, such as salvage logging, and fuel treatment (Ascoli et al., 2013).

89 In the present study, we focus on the key questions of detecting the major drivers  
90 influencing delayed post-fire beech mortality. In particular we ask:

91 (1) Are there any temporal trends in fire-caused beech mortality?



92 (2) Which tree-specific traits (e.g., tree size) enhance the survivability of fire-injured  
93 beech trees?

94 (3) What are the main factors associated with beech mortality after fire disturbance?

## 95 **2 Materials and methods**

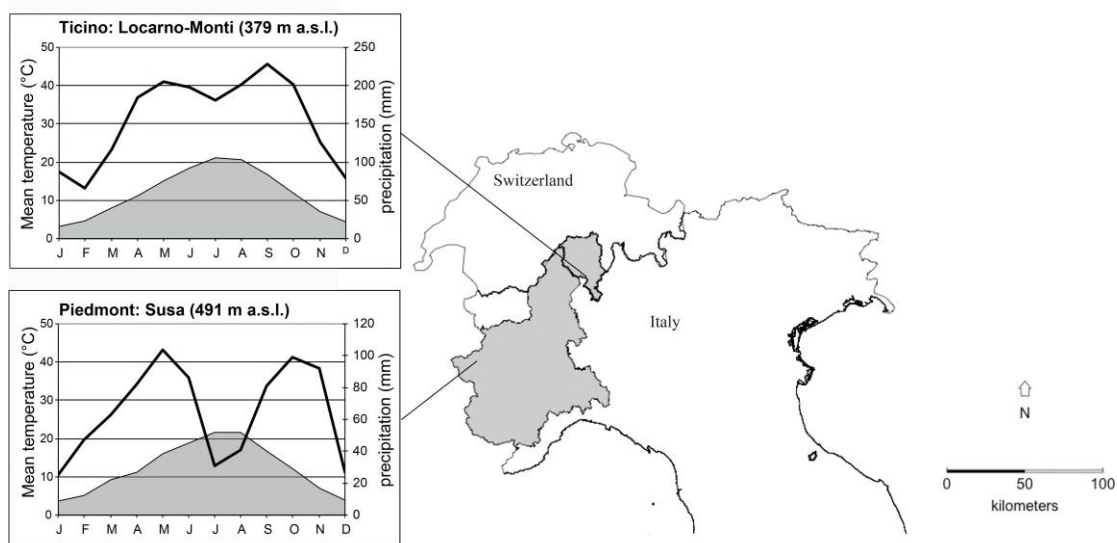
### 96 **2.1 Study area**

97 The present study was conducted in the neighboring regions of Piedmont (Italy) and  
98 Ticino (Switzerland) located in the Southwestern European Alps (Figure 1). Both  
99 regions are characterized by a marked elevation gradient along which forest  
100 vegetation types are distributed. Beech dominated forests occupy the intermediate  
101 elevation belt ranging from 600-1,000 m a.s.l. to 1,300-1,700 m a.s.l. depending on  
102 the locality and aspect (Camerano et al, 2004; Ceschi, 2006). These forests are mostly  
103 in the process of transformation from former unevenly-aged coppice management to  
104 high-stand forests (Nocentini, 2009).

105 The area of investigation is characterized by a gradient of precipitation that ranges  
106 from the drier Piedmont region with an annual precipitation of 778 mm and a mean  
107 annual temperature of 12.3°C (Susa meteorological station: 07°3'0"E, 45°08'0"N;  
108 Arpa, Piedmont) to the wetter Canton Ticino, with an annual precipitation of 1,897  
109 mm and a similar mean annual temperature of 12.4°C (Locarno-Monti meteorological  
110 station: 08°47'43"E, 46°10'12"N; observation period 1981-2010; MeteoSwiss, 2015).

111 In winter and early spring, northern foehn winds cause episodic relative humidity of  
112 below 20% accompanied by significant temperature rises (Isotta et al., 2014). These  
113 conditions favor winter surface fires, which are mostly induced by humans. Such fires  
114 usually start at the wildland-urban interface (Conedera et al., 2015) and episodically  
115 spread into more highly elevated beech forests (800-1,500 m a.s.l.). Total winter

116 (DJF) precipitation ranges from 158 mm (Piedmont) to 495 mm (Ticino) (Arpa  
 117 Piedmont; MeteoSwiss, 2015). The generally dry winters contrast with the humid  
 118 summers (JJA) where dry spells normally do not last longer than thirty consecutive  
 119 days (Isotta et al., 2014). Therefore, summer fires rarely occur in climatically average  
 120 years, but may ignite (by both lightning and humans) and spread with particular  
 121 intensity during extraordinary and prolonged drought, such as in the summer 2003  
 122 (Ascoli et al., 2013; Valesse et al., 2014).



123  
 124 **Figure 1: The study region on the southern slope of the Alps located in Canton Ticino (Switzerland) and**  
 125 **Piedmont Region (Italy) marked in grey with representative climate diagrams.**

126

## 127 2.2 Selection of fire sites

128 We examined the Swiss forest fire database (Pezzatti et al., 2010) and those of the  
 129 Italian State Forestry Corps (Ufficio Territoriale per la Biodiversità di Verona Centro  
 130 Nazionale Biodiversità Forestale di Peri) for the purpose of identifying fire sites that  
 131 burnt after 1970. In order to keep the sample uniform, we limited our selection to  
 132 beech forests on crystalline bedrock. To this end, we overlaid the obtained fire  
 133 perimeters with detailed regional forest and geological maps (Ceschi, 2006;

134 Camerano et al., 2004) in a geographical information system (GIS) (version 10.0; ©  
135 ESRI). In total, we obtained 94 potential beech stands affected by individual fires that  
136 we inspected in summer 2011 to select fire sites that matched all of the following  
137 criteria: (i) pre-fire stands dominated by beech with >95% of their stems, (ii) burnt  
138 area within the beech forest of >0.25 ha, (iii) no signs of additional fires during the  
139 last 50 years, (iv) no evidences of pre-fire wood pasture, and (v) no post-fire artificial  
140 regeneration (plantations). From the examined 94 fire sites, 36 satisfied all of the  
141 selection criteria and were retained for the field survey in the years 2012 and 2013  
142 (Appendix A).

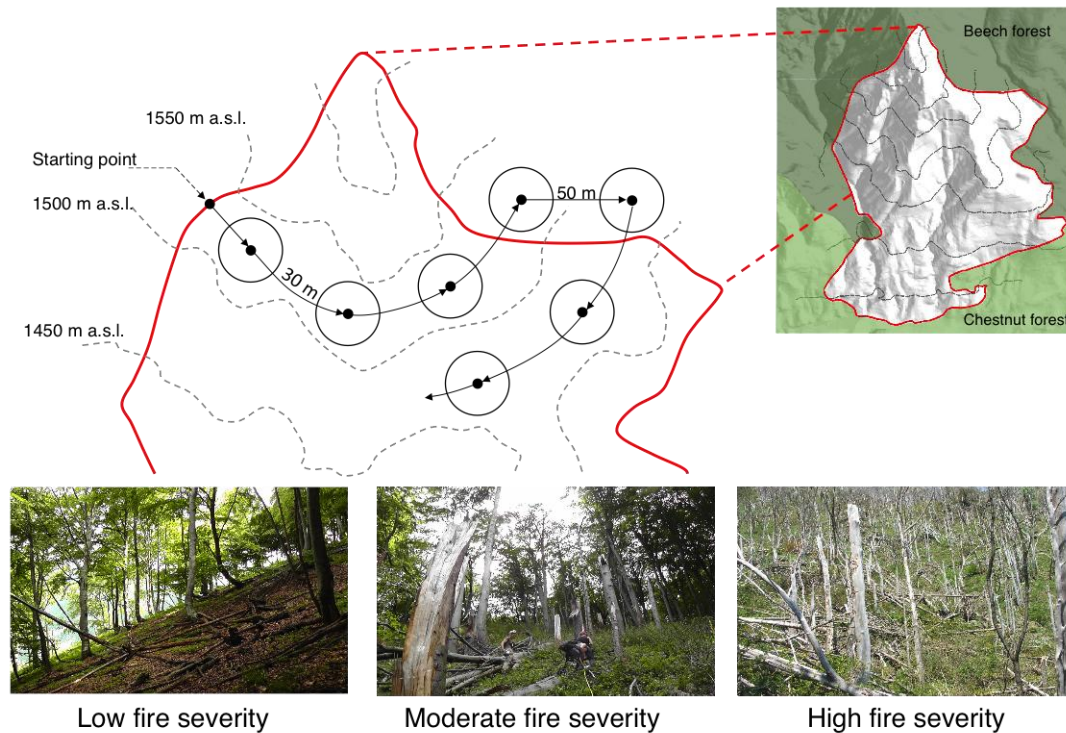
### 143 **2.3 Data collection**

#### 144 *Sampling design*

145 We placed between one and three transects in each fire site, spaced 50 m apart in  
146 elevation and following the contour lines (see Figure 2). Along the transects, circle  
147 plots of 200 m<sup>2</sup> were placed 30 m apart, starting at a distance of 10 m from the  
148 unburnt forest. Wherever possible, a minimum of one and a maximum of 4 control  
149 plots per fire site were placed within the adjacent, unburnt beech forest (see Figure 2).  
150 Fieldwork was conducted between July 2012 and September 2013, and a total of 234  
151 and 39 plots were assessed in burnt and unburnt beech forests, respectively (Appendix  
152 A).

153

154



155

156 **Figure 2: Sampling design in the upper part of the burnt beech forest. Circular plots of 200 m<sup>2</sup> were placed**  
 157 **30 m apart along horizontal transects from the burnt into the unburnt beech forest (figure left). Each plot is**  
 158 **further characterized in terms of burn severity as a function of the portion of dead beeches (photographs).**

159

160 *Field measurements: plot characteristics and stand structure*

161 Each 200-m<sup>2</sup> plot was characterized by its slope, aspect, elevation, and micro-  
 162 topography (concave, plane, convex). Every pre-fire tree was classified as **alive or**  
 163 **dead**. Dead individuals were further distinguished as dead standing trees (dead  
 164 standing tree with crown portions but without visible green foliage and snags) and  
 165 dead fallen trees (logs). We attempted to identify each tree (alive, dead) with diameter  
 166 at breast height (DBH)  $\geq 8$  cm at the species level, which was sometimes not possible  
 167 because of the progressed wood decay stage. DBH was measured to the nearest cm  
 168 for all standing trees, and the average diameter was recorded for logs. For standing  
 169 beech individuals, data collection further included growing habitus (monocormic,  
 170 polycormic), tree height, percentage of crown volume killed, decay stage of the wood,

171 height of the fire scar, proportion of damaged bark, and fungal activity. Tree growing  
172 habitus was defined as polycormic if two or more resprouts grew out of the same  
173 stool. The percentage of crown volume killed was visually estimated by the  
174 volumetric proportion of crown killed compared to the space occupied by the pre-fire  
175 crown volume (Hood et al., 2007). In order to assess the contribution of fungi  
176 infestation to the mortality process (Conedera et al., 2007; Conedera et al., 2010),  
177 fungal fructification (fruit bodies) was assessed quantitatively by assigning one of  
178 four abundance classes (none, few, partial, mass) to each beech tree. A subset of the  
179 fungal specimens was collected, put in paper bags, and transported to the laboratory  
180 for species determination according to Krieglsteiner (2000), Gerhard (2005) and Klug  
181 and Lewald-Brudi (2012). For all dead standing trees and logs, the decay stage of the  
182 wood was classified according to following classes: (1) cambium still fresh, (2) knife  
183 penetrates low, cambium disappeared, (3) knife penetrates into the fiber direction, but  
184 not transversely, or (4) knife penetrates in both directions.

#### 185 *Assessment of climatic variables*

186 Precipitation and temperature can influence tree mortality (Lines et al., 2010) and  
187 both variables may occur as secondary stressors in fire-injured trees. Therefore,  
188 precipitation and air temperature data was obtained for each fire site from the  
189 WorldClim Database (Hijmans et al., 2005). Yearly precipitation and temperature  
190 averages refer to the period 1950–2010.

#### 191 *Assessment of burn severity*

192 In accordance with a parallel study by Maringer et al. (subm.), we assessed burn  
193 severity by calculating the basal area of post-fire/pre-fire living trees. For old (>10  
194 years post-fire) fire sites, pre-fire conditions were assessed exclusively from the

195 control plots, while in recently burnt areas ( $\leq 10$  years), pre-fire stand characteristics  
196 were determined by the number of visible dead trees in burnt plots. We defined low  
197 burn severity in plots with less than 5% crown volume loss and less than 20% basal  
198 area loss. In contrast, high burn severity was indicated by extensive crown loss ( $>$   
199 50%) and basal area killed ( $> 60\%$ ). All plots with intermediate losses in terms of  
200 crown and basal area were assigned to the moderate severity class.

#### 201 **2.4 Data aggregation**

202 To describe temporal trends in fire-caused beech mortality, sites were categorized  
203 according to the number of years passed since the fire event, i.e., “ $\leq 9$  years”, “10–15  
204 years”, “16–21 years”, “22–32 years”, and “32–43 years”, always including the  
205 corresponding control plot in the unburnt beech forest. Plots were additionally  
206 classified according to their proportion of poly- to monocormic trees: (i) stands  
207 evolved into high forests ( $< 33\%$  polycormic trees), (ii) stands in the transition phase  
208 between former unevenly-aged coppices and high forests (33–66% polycorm trees),  
209 and (iii) unevenly-aged coppices ( $> 66\%$  polycorm trees). Finally, standing pre-fire  
210 beeches were grouped into four DBH-classes (8–12 cm, 12–24 cm, 24–36 cm, and  $\geq 36$   
211 cm; Frehner et al., 2005). For all DBH-classes, stem density ( $N\ ha^{-1}$ ) and basal area  
212 ( $m^2\ ha^{-1}$ ) were calculated separately for living trees and dead standing trees,  
213 respectively.

#### 214 **2.5 Beech mortality model construction**

215 We assessed the risk of beech mortality using mixed-logit models (GLMM). Since  
216 models were individual tree-based using the vitality status (alive or dead) of the  
217 standing beeches as a response variable, within-plot data violated the rule of  
218 independency. Thus, plots were included as a random effect for intercepts, in addition

219 to potential risk factors (explanatory variables) characterizing site-, plot- and  
220 individual-trees. These included total annual precipitation (**PREC**) and annual mean  
221 temperature (**TEMP**) at site level, micro-topography (**TOPO**), slope (**SLO**), elevation  
222 (**ELE**), and aspect (**ASP**) at plot level, as well as tree size (**DBH**, **HEIGHT**), growth  
223 habitus (**POLY**), and fungi fructification (**FUNGI**) at tree level. Continuous risk  
224 factors were standardized to calculate the mixed-logit models, which related the  
225 probability  $\pi_{ijk}$  of mortality for an individual beech tree  $j$  in a particular plot  $i$  over the  
226 number of years post-fire (YPF)  $k$  to the mentioned risk factors ( $X_1, \dots, X_n$ ) as follows:

$$227 \log(\pi_{ijk}/1 - \pi_{ijk}) = \beta_0 + \text{offset}(\log(\text{YPF}_{ik})) + \beta_1 X_{(\text{treeIndex})ij} + \dots + \beta_n X_{(\text{siteIndex})i} + y_i$$

228 where  $\beta_0$  represents the overall intercept,  $\beta_1$  to  $\beta_n$  the regression parameters for the  
229 corresponding variables, and  $y_i$  the random effect. The offset function corrects the  
230 number of mortality events for different YPF values (Boeck et al., 2014). Continuous  
231 predictors ( $X$ ) were visualized and afterwards implemented in the models as linear  
232 and/or quadratic terms.

233 As a general rule, values referring to tree characteristics such as stem diameter and  
234 height were recalculated based on the year of fire. Average annual growth rates  
235 (Z'Graggen, 1992; Eidg. Anstalt für das forstliche Versuchswesen (EAFV), 1983)  
236 were subtracted from **DBH** and **HEIGHT** for all years post-fire (Table 1). Fungi  
237 infestation normally starts within the second year post-fire (Conedera et al., 2007;  
238 Conedera et al., 2010) and was therefore regarded as an initial parameter. In contrast,  
239 proportions of bark damage and the length of the fire scar were excluded from the  
240 modeling approach, because immediate fire effects were impossible to reconstruct for  
241 older fire events due to the rapid progression of wood decay.

242 Assuming that factors affecting beech mortality might alter their influence as a  
243 function of burn severity, we performed models separately for low, moderate, and

244 high burn severities (hereafter referred to as low-model, moderate-model, high-  
245 model). To validate the influence of fire on beech mortality, a separate model was  
246 conducted for the unburnt forests (*control*). Data exploration followed the guidelines  
247 of Zuur et al. (2010), which suggest detecting collinearity among variables by using  
248 Pearson's correlation coefficient and the variance inflation factor (VIF). After  
249 excluding **HEIGHT** ( $r^2 > 0.8$  with **DBH**) from all models and **TEMP** ( $r^2 > -0.7$  with  
250 **PREC**) from the low-severity model, all VIFs were below 3, indicating the absence of  
251 any critical collinearity.

## 252 **2.6 Model performance and selection**

253 By choosing a GLMM, the data assumed a two-level hierarchical structure with pre-  
254 fire trees at level 1 nested within plots at level 2. Hence, variables were categorized at  
255 level 1 and 2, and model selection started by considering only standardized level 1  
256 variables.

257 After finding significant explanatory variables at level 1, level 2 variables were then  
258 included in models and both were tested for interactions. During this process, low  
259 variations were found for the estimated values for **FUNGI** with four expressions  
260 (none, low, few, high). Consequently, this variable was converted into a dummy  
261 variable (0/1).

262 GLMM model selection referred to the lowest information-theoretic approach based  
263 on the correct Akaike information criterion (AIC; Venables and Ripley, 1999), and  
264 explanatory variables were retained if significantly different from zero ( $p \leq 0.05$ ).  
265 Model diagnostics checked for the best-fitting models based on deviance residuals  
266 that were plotted against the fitted values and all variables included and not included  
267 in the model to detect unusual patterns in residuals (Zuur et al., 2010).



268 All analyses were performed using R statistics software (R Development Core Team,  
269 2014). Logistic regression models were fitted and validated using the lme4 (Pinheiro  
270 et al., 2015) and VGAM (Yee et al., 2015) packages. Graphical outputs were mainly  
271 produced using the packages lattice (Deepayan, 2008) and ggplot2 (Wickham and  
272 Chang, 2015), and maps were created using map and GIS tools (Brownrigg, 2015;  
273 Brunsdon and Chen, 2015).  
274

275 **Table 1: Risk factors included (•) and excluded (–) in the calculated mixed-logit models (GLMM) for burnt**  
 276 **(B)<sup>1</sup> and unburnt (UB) plots.**

Variables	Abbreviation	Unit	Models	
			B	UB
<i>response variable</i>				
beech living status	STATUS	0=alive, 1=dead	•	•
<i>topography</i>				
slope	SLOPE	%	•	•
aspect	ASP	°	•	•
elevation	ELE	m a.s.l.	•	•
micro-topography	TOPO	factor	•	•
<i>climate</i>				
temperature	TEMP	°C	• <sup>2</sup>	•
precipitation	PREC	mm	•	•
<i>tree characteristics</i>				
diameter at breast height	DBH	cm	•	•
height	HEIGHT	m	--	--
growth habitus	POLY	0/1	•	•
fungi cover	FUNGI	0/1	•	•

277 <sup>1</sup> calculated separately for low, moderate, and high burn severity

278 <sup>2</sup> not used in the low-model

## 279 **3 Results**

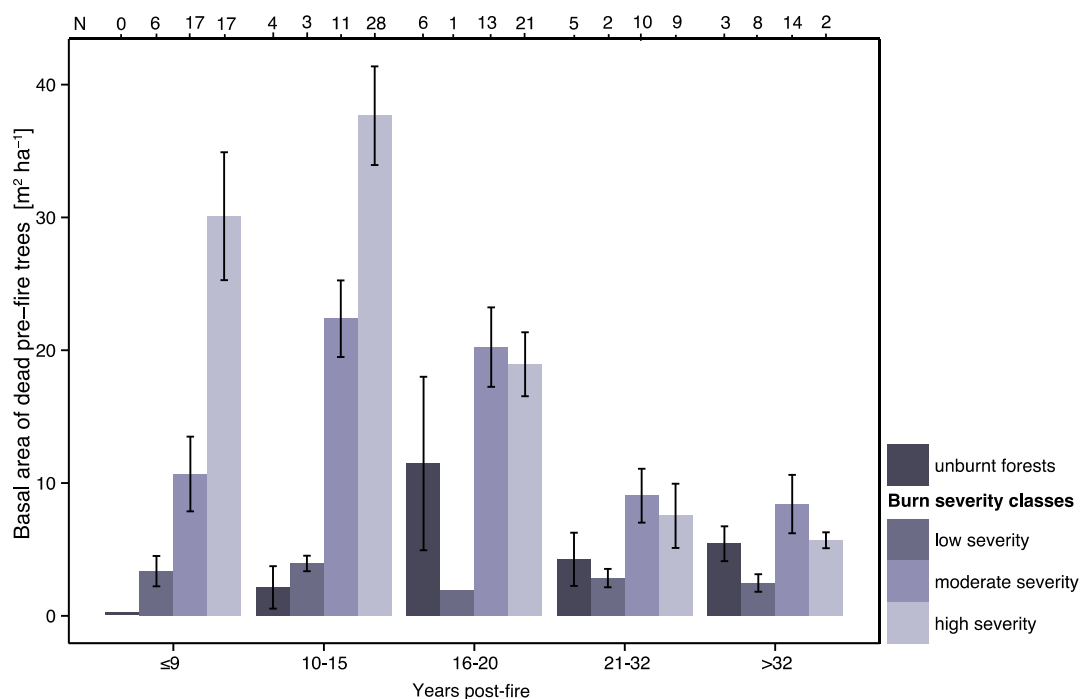
### 280 **3.1 Forest structure**

281 Most (61%) of the burnt forest stands were classified as high stand forests, a minority  
 282 (16%) as unevenly-aged coppices, with the remainder within the transition phase. In  
 283 total, 3,504 mature trees were recorded, of which beech consisted of 88% and 93% in  
 284 the burnt and unburnt forests, respectively. Other tree species rarely (< 4%) grew  
 285 within the pure beech stands (Appendix B).

### 286 **3.2 Post-fire beech mortality**

287 From the overall number of beech trees assessed in burnt plots (N = 2,845), half died  
 288 due to fire impact, whereas in unburnt forests, only 10% of the recorded trees were  
 289 classified as dead. Fungi infestation occurred in 23% of survived beeches, and 72% of  
 290 dead trees. We found at least 10 different fungal species on the stems of fire-injured

291 beeches (see Table 3). As reported in Figure 3, the average basal area of standing  
 292 dead beeches in burnt forests was  $14.1 \pm 0.95 \text{ m}^2 \text{ ha}^{-1}$ , ranging from  $1.9 \text{ m}^2 \text{ ha}^{-1}$  to  
 293  $37.6 \text{ m}^2 \text{ ha}^{-1}$  over the years post-fire. Tree mortality in low burn severity sites is quite  
 294 similar to that in unburnt forests, while tree mortality increases with burn severity and  
 295 peaks 10 to 15 years post-fire. The highest overall basal area losses (up to 85% of the  
 296 initial value) occurred in high severity sites, followed by moderate severity sites (up  
 297 to 63%).  
 298

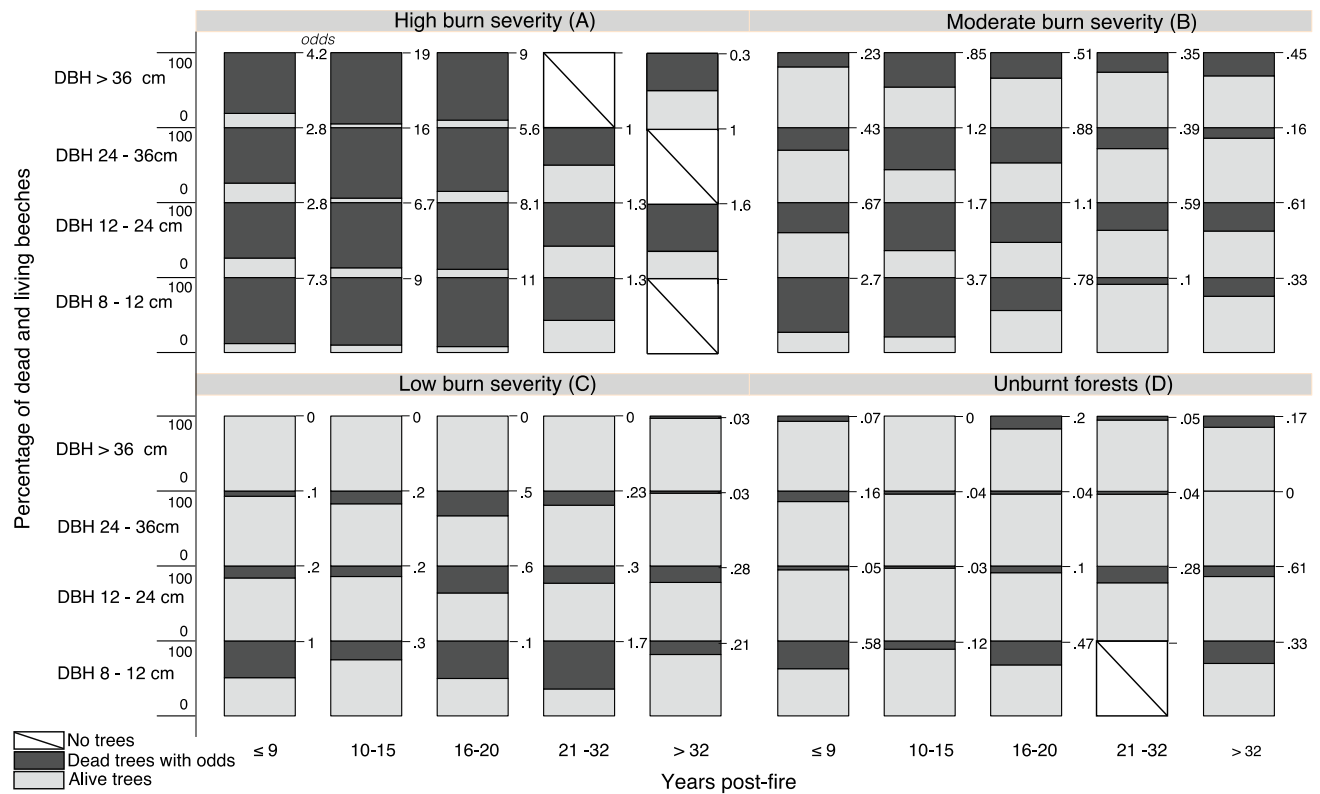


299  
 300 **Figure 3: Mean ( $\pm$ SE) basal area of standing dead pre-fire beeches in low-, moderate-, and high severity**  
 301 **sites, and the corresponding unburnt plots as a function of years post-fire.**

302 Taking unburnt forests as the reference, the odds of beech mortality (i.e., the ratio of  
 303 the probability of dying vs. surviving) was 42, 5, and 2.3 times greater in high-,  
 304 moderate-, and low severity sites, respectively. Within the burn severity classes, the  
 305 extent and timing of beech mortality varied as a function of tree size. In low-severity  
 306 sites, tree mortality was usually limited to small beeches, whereas in the case of

307 moderate severity, intermediate-sized beeches were also affected. Finally, beech  
308 mortality was high and affected all tree sizes in high severity plots. There, mortality  
309 started immediately after fire disturbance and continued up to two decades post-fire  
310 with mortality odds ratios always greater than 2.8 (Figure 4 A). In contrast, in  
311 moderate severity plots, the odds of mortality for small (DBH < 12 cm) beeches were  
312 two to six times higher than for intermediate-sized, and four to eleven times higher  
313 than for large and tall beeches (DBH > 36 cm). These differences in the mortality rate  
314 are clear within the first 15 years post-fire, when mortality is higher than in  
315 subsequent years (Figure 4 B). Similar patterns were observed in low severity sites,  
316 where the odds of death for small beeches were generally higher than for tall and  
317 large beeches (Figure 4 C). In these fire sites, the probability of large-beech mortality  
318 was near zero, while intermediate-sized beech mortality ranged between 0.03 and  
319 0.56 throughout the whole observation period.

320



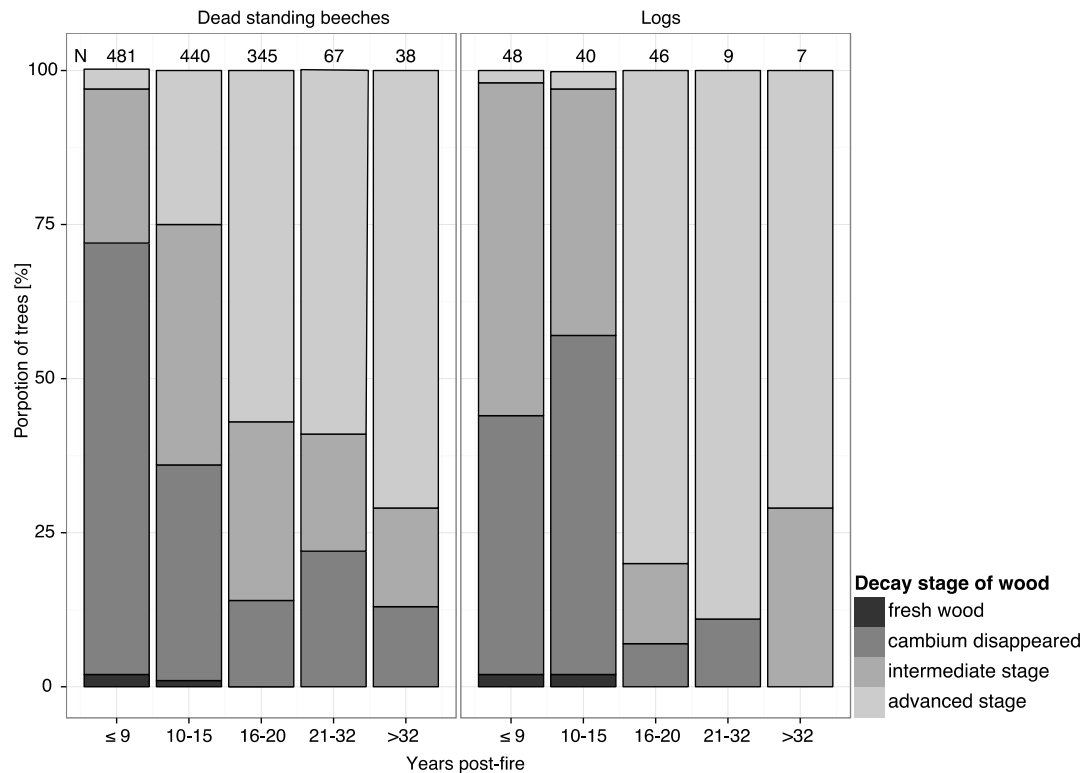
321  
322

323 **Figure 4: Percentage of survived (light grey) and dead (dark grey) beeches for small (DBH < 12 cm),**  
 324 **intermediate-sized (DBH = 12 – 24 cm), large (DBH = 24 – 36 cm), and very large (DBH > 36 cm)**  
 325 **individuals, separated for different burn severities and years post-fire. The odds ratios of mortality are also**  
 326 **shown to the right of each column.**

327

### 328 3.3 Deadwood characteristics

329 In the first decade post-fire, the majority of standing dead trees (70%) and logs (54%)  
 330 showed slight to intermediate decomposition (Figure 5). Over the years post-fire, the  
 331 proportion of intermediate and advanced wood decay stages steadily increased, finally  
 332 ranging between 64% (10-15 years post-fire) and 87% (> 32 years post-fire) for  
 333 standing dead trees, and exceeding 90% for logs 16 years post-fire. As a result, in the  
 334 first two decades post-fire, the number of standing dead trees was ten times higher  
 335 than logs. After the second decade post-fire, both the number of standing dead trees  
 336 and logs decreased by approximately 80%. Figure 5 clearly illustrates that dead  
 337 standing trees appear to dominate in the first 20 years post-fire.



338

339 **Figure 5: Decay stages of wood for standing dead trees and logs as a function of years post-fire.**

340

### 341 **3.4 Drivers of beech mortality**

342 The best beech mortality models clearly describe the mortality rate for fire-injured  
 343 beeches, with FUNGI (fungi fruitbodies), DBH, and ELE (elevation) as common  
 344 factors (Table 2). FUNGI had a positive and highly significant ( $p < 0.001$ ) effect on  
 345 beech mortality in all three burn severity models, indicating an increased risk of  
 346 mortality after visible fungal activity in terms of fruit body formation. The odds ratio  
 347 of beech mortality after fungi fructification was 14.8 in the moderate-severity model,  
 348 which was four times that of the low-model and double that of the high-severity  
 349 model.

350 In addition to fungi fructification, DBH was significantly and negatively correlated  
 351 with beech mortality in the low- and moderate-models, indicating a consistently  
 352 decreasing risk of mortality with increasing DBH. The odds of tall beech survival

353 after a low severity fire was four-times higher than for moderate burn severity,  
354 whereas no detectable correlation existed between mortality and DBH in the high-  
355 model. The correlation between beech mortality and the quadratic term of DBH was  
356 found to be positive in addition to the linear predictor in the control-model (unburnt  
357 forests), indicating increased mortality for tall and thin beeches. Polycormic growth  
358 habitus (POLY) reduced the mortality risk factor in moderate- and high-models, but  
359 not in the low-model.

360 Site factors only marginally correlated with beech mortality, except for ELE. The  
361 linear and quadratic terms of ELE were weakly but significantly and positively  
362 correlated with beech mortality in all three fire-severity models. The explanatory  
363 power of the low-model containing all three variables (DBH, FUNGI, ELE) was 38%.  
364 The quadratic term of TEMP weakly and negatively correlated with beech mortality  
365 in the moderate- and high-models, respectively. Furthermore, positive correlations  
366 with beech mortality were found for PREC in the moderate-model and ASP in the  
367 high-model. In summary, beech mortality was explained by six (FUNGI, DBH,  
368 POLY, ELE, TEMP, PREC) variables with an explanatory power of 32% in the  
369 moderate-model, and by five (FUNGI, POLY, ELE, TEMP, ASP) variables with an  
370 explanatory power of 17% in the high-model.

371 From the above-mentioned variables, the linear and quadratic terms of DBH had the  
372 most explanatory power in the control-model. With DBH as the only highly  
373 significant variable, the model had an explanatory power of 66%, whereas the weakly  
374 (0.1%-level) correlated variables ELE and ASP only marginally improved the model.

Table 2: Results of the mixed-logit models for the burnt and unburnt forests separated for low (low-model), moderate (moderate-model) and high (high model) burn severities.

Models	Burnt forests			Unburnt forests
	<i>Low-model</i>	<i>Moderate-model</i>	<i>High-model</i>	<i>Control</i>
	Odds ratio [conf. interval]	Odds ratio [conf. interval]	Odds ratio [conf. interval]	Odds ratio [conf. interval]
<b><i>fixed term</i></b>				
Intercept	34 <sup>-1***</sup> [86 <sup>-1</sup> -15 <sup>-1</sup> ]	53 <sup>-1***</sup> [93 <sup>-1</sup> - 40 <sup>-1</sup> ]	1.5 <sup>-1</sup> [3.3 <sup>-1</sup> -1.8]	51 <sup>-1***</sup> [176 <sup>-1</sup> -22 <sup>-1</sup> ]
FUNGI	4.6 <sup>***</sup> [2.2-9.9]	14.8 <sup>***</sup> [9.8-21.5]	7.4 <sup>***</sup> [3.8-14.3]	ns
DBH	5 <sup>-1***</sup> [8.3 <sup>-1</sup> -3.1 <sup>-1</sup> ]	1.7 <sup>-1***</sup> [1.8 <sup>-1</sup> -1.3 <sup>-1</sup> ]	ns	4.7 <sup>-1***</sup> [8.2 <sup>-1</sup> -2.8 <sup>-1</sup> ]
DBH <sup>2</sup>	ns	ns	ns	1.7 <sup>***</sup> [1.4-2.1]
POLY	ns	2.0 <sup>-1***</sup> [2.9 <sup>-1</sup> -1.3 <sup>-1</sup> ]	2.2 <sup>-1*</sup> [4.5-1.3]	ns
ELE	2.8 <sup>*</sup> [1.1-6.7]	ns	2.1 <sup>-1*</sup> [4.1 <sup>-1</sup> -1.2]	1.9 <sup>-1*</sup> [4 <sup>-1</sup> -1]
ELE <sup>2</sup>	2.3 <sup>*</sup> [1.1-4.7]	1.3 <sup>**</sup> [1.1-1.5]	ns	ns
TEMP <sup>2</sup>	ns	1.2 <sup>-1*</sup> [1.5 <sup>-1</sup> -1]	1.8 <sup>-1**</sup> [2.7 <sup>-1</sup> -1.2 <sup>-1</sup> ]	ns
PREC	ns	1.6 <sup>***</sup> [1.2-2.2]	ns	ns
PREC <sup>2</sup>	ns	ns	ns	ns
ASP	ns	ns	1.7 <sup>*</sup> [1-2.9]	1.8 <sup>*</sup> [1.1 <sup>-1</sup> -4.3]
<b><i>random term</i></b>				
Plot	Var (x)[SD]	Var (x)[SD]	Var (x)[SD]	Var (x)[SD]
	1.7[1.3]	0.5[0.6]	1.1[1]	2.6[1.6]
D <sup>2</sup> <sub>fixed effects</sub>	38%	32%	17%	66%
D <sup>2</sup> <sub>fixed + random effects</sub>	55%	38%	35%	80%

376 Signif. codes: '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 'ns' 1. Abbreviations see Table 1



## 378 4 Discussion

### 379 4.1 Post-fire stand dynamics

380 The selected stands showed typical beech forest structural characteristics for the  
381 Southern Alps, with interlocking transition stages from unevenly-aged coppices to  
382 high forest stands (Nocentini, 2009; Ascoli et al., 2013). In these stands, fires of  
383 mixed severity induced significant forest structure changes by causing mortality in  
384 half of the pre-fire beeches. In general, most fire-caused beech mortality occurs  
385 within the first two decades post-fire. Such short-term mortality has also been  
386 observed in studies of other fire-injured broadleaf stands (Harrod et al., 2000;  
387 Ledgard et al., 2004; Shafiei et al., 2010; Catry et al., 2010; Bravo et al., 2014).  
388 Moreover, our findings are in accordance with a companion study (Maringer et al.,  
389 *subm.*) that observed a post-fire regeneration window spanning the first two decades  
390 post-fire, when the regeneration takes advantages of progressive canopy openings and  
391 particularly suitable germination beds.

392 As already reported in numerous studies for other tree species (e.g., Keyser et al.,  
393 2008; van Mantgem et al., 2013) the extent and pace of beech mortality highly depend  
394 on tree size. We observed a decreasing risk of mortality for beeches from thinner  
395 (DBH < 12 cm) to tall and large (DBH > 36 cm) individuals. With increasing time  
396 since fire (> 20 years post-fire), the mortality rate for beeches decreases toward a  
397 nearly natural level (control plots). Similar observations are reported for Oriental  
398 beech (*Fagus orientalis* LIPSKY) forests 37 years after a fire (Shafiei et al., 2010; Adel  
399 et al., 2013).

400 Standing dead beeches, snags in particular, accounted for the majority of the resulting  
401 dead wood, which is typical for burnt forests (Lampainen et al., 2004; Brown et al.,

402 2013). The number of dead standing trees and especially the amount of logs—which  
403 usually have contact with the soil surface—rapidly decrease with time since fire, due  
404 to the fast decomposition rate of beech wood relative to standardized time intervals  
405 (Kahl, 2008; Lombardi et al., 2008).

#### 406 **4.2 Triggers of post-fire beech mortality**

407 Among the detected drivers of post-fire beech mortality, the presence of visible fungal  
408 activity in terms of fruit body formation revealed to be the most relevant. The  
409 accelerating effect of secondary fungal activity in the dieback process of fire-injured  
410 beeches is well known, not only for European beech (Conedera et al., 2007; Conedera  
411 et al., 2010; van Gils et al., 2010) but also for American beech (*Fagus grandifolia*  
412 Ehrh.) (Tubbs and Houston, 1990). Thin bark is one characteristic of the genus *Fagus*  
413 that renders beech species particularly susceptible to fire scar formation (Tubbs and  
414 Houston, 1990; Peters, 1997; Hicks, 1998; Packham et al., 2012) and thus potentially  
415 to secondary fungal infestation. Fungi infestation causes rapid decay in the exposed  
416 wood tissues in fire scars, reducing trunk mechanical stability and, in turn, the time  
417 and likelihood of successful scar overgrowth and healing.

418 This is likely the reason why beech diameter as a risk factor was found to play only a  
419 secondary role in the modeling of post-fire beech mortality patterns, although tree size  
420 is linearly and negatively correlated to beech mortality in low- and moderate-models.  
421 The fact that tree mortality caused by fire is higher for thinner trees than for larger  
422 ones has often been stated in the literature for different tree species (McHugh and  
423 Kolb, 2003; Kobziar et al., 2006; Shafiei et al., 2010; Brando et al., 2012). Most  
424 studies relate the higher mortality rate of thinner trees to bark thickness and tree  
425 height, which usually increase with tree age (Gutsell and Johnson, 1996; Lawes et al.,  
426 2013). However, to date little is known about the relationship between European

427 beech bark thickness and tree size. For Oriental beech, a close relative to the target  
428 species, Bonyad et al. (2012) discovered a strong positive correlation between DBH  
429 and bark thickness. Shekholeslami et al. (2011) investigated Oriental beech bark  
430 thickness along the trunk and found thicker bark on the bole of the trees, a trait that is  
431 evident also for larger beeches in the Alps (Figure 6). Studies on large European  
432 beeches have found more structured, creviced, and rough bark (Russo et al., 2010;  
433 Dymytriva et al., 2014), characteristics which have been found to increase thermal  
434 insulation and thus resistance to fire injury (Fahnestock and Hare, 1964; Nicolai,  
435 1986; Bauer et al., 2010; Odhiambo et al., 2014). Also considering that larger beeches  
436 have large crowns, low limbs, and limited litter yield around their boles (Yaussy et  
437 al., 2004), it is understandable that survivability increases for large individuals after  
438 surface fires of low to moderate intensity.



439

440 **Figure 6: Old beech with structured, creviced, and rough bark (left figure) and a cross section with a thick**  
441 **and relatively rough bark (right figure; Piedmont, Italy; D. Ascoli)**

442

443 In summary, the combination of the portion of woody tissue exposed through fire  
444 scars and the likelihood of being colonized by active and fructifying fungi appear to  
445 be crucial in inducing the mortality process in fire-injured beeches. In this respect, the  
446 results of our modeling approach confirm the findings of Conedera et al. (2010),  
447 indicating that the moderate-burn severity class is the most sensitive to secondary

448 fungal infestation (odds ratios four and two times higher with respect to low and high  
449 severity, respectively).

450 Our sampling design does not permit a conclusive analysis of the role played by fungi  
451 species and their particular timing of infestation. Once colonized, the visible  
452 abundance of fungal fructifications does not appear to be an indicator of enhanced  
453 risk of mortality, as revealed by our modeling approach. Similarly, neither the season  
454 of the fire event (vegetation rest or growing season) nor the fungal species concerned  
455 play an apparent role, given that the lethal issue of fungal activity is not dependent on  
456 the fire date (data not shown) or the fungal species.

457 Interestingly, in this study, the assessed fungal species colonizing fire scars appear to  
458 differ from those infesting mechanically injured beeches (Table 3; see Standovár and  
459 Kenderes (2003) for a review and Conedera et al. (2007) for fire related fungi). The  
460 existence of specific, fire-related fungi infesting beeches is thus confirmed. The  
461 question remains, however, regarding the specific ecological conditions needed at  
462 colonization time, and the different impact of each fungal species in the mortality  
463 process of beech trees. Even though we did not directly measure the abundance of  
464 fungal sporulation in the present study, we postulate that the sheer presence of fungal  
465 fruit bodies may be more important than the amount of fungal fruit bodies. This  
466 relationship is yet to be tested in further field work.

Species	Short biological description
<b>Fungi on fire-injured beeches</b>	
<i>Armillaria spec.</i> (Fries) Staude	----
<i>Cerrena cf. unicolor</i> (Bull.) Murrill	Spaced forest stands on humid soils. <sup>d</sup>
<i>Daldinia concentrica</i> (Bolton) Cest. & de Not.	Specifically adapted to wildfire and can be invisible for many decades. <sup>j</sup>
<i>Inonotus nodulosus</i> (Fr.) P. Karst	Usually occurs on humid soil during late successional forest stages. <sup>d</sup>
<i>Irpex lacteus</i> (Fr.)	Usually abundant in fire scars where it causes white rot finally causing the stem to break. <sup>j</sup>
<i>Oudemansiella mucida</i> (Schrad.) Höhn.	Sabrobiont, occurs in the early wood-decay stage on dead standing beeches or on living beeches. Especially in regions with high humidity. <sup>d</sup>
<i>Schizophyllum commune</i> (Fr.)	Often occur after “sun burn” on broadleaf trees. <sup>j</sup>
<i>Stereum hirsutum</i> (Willd.) Pers.	Pioneer species fruits often after fires in deciduous forests. <sup>j</sup>
<i>Trametes hirsuta</i> (Wulfen) Pilát	Occurs on injured beeches, which are exposed to light. Sabrobiont on dead standing or lying trees, which still have pieces of bark. <sup>d</sup>
<i>Fomes fomentarius</i> (L. ex Fr.) Gill.	White rot of beech wood and other broadleaf species; occurs on living, standing beeches and leads to progressive wood decay. The infested tree normally breaks at its weakest point. <sup>h</sup>
<b>Fungi on mechanically-injured beeches</b>	
<i>Daedalea quercina</i> (L.) Fr.	Causing brown rot often leading to huge wood loss inside the stem. <sup>a</sup>
<i>Fomitopsis pinicola</i> (Sw. ex Fr.) Gill.	Unable to invade living sapwood, but wounded trees are easily colonized. <sup>b</sup>
<i>Ganoderma applanatum</i> (Pers.) Pat.	Causes white heart rot and is dispersed by a specialized mycophagous fly. <sup>c</sup>
<i>Inonotus radiatus</i> (Sw. ex Fr.) Karst.	The main host is alder ( <i>Alnus</i> sp.) but also occurs on dying beeches. <sup>d</sup>
<i>Laetiporus sulphureus</i> (Bull.) Bond. Ex Sinq.	Intensive wood decay in the tree’s trunk causing static problems mainly in the root crown. <sup>e</sup>
<i>Inonotus cuticularis</i> (Bull.) P. Karst.	Causes brown rot, mainly on beeches in barely disturbed forests. <sup>d</sup>
<i>Inonotus obliquus</i> (Pers.) Pilát	Cause white heart rot. The fungus penetrates the tree through poorly-healed wounds. <sup>f</sup> Decay may continue for 10–80+ years inside a living host tree. <sup>d</sup>
<i>Meripilus giganteus</i> (Pers.) P. Karst	Causes intensive white rot, mainly on beech and oak wood. <sup>d</sup>
<i>Nectria galligena</i> Bres.	Causes cancer disease. Entry of the pathogen is facilitated by the beech scale insect ( <i>Cryptococcus fagisuga</i> ). <sup>d</sup>
<i>Nectria ditissima</i> Tul.	Similar to <i>N. galligena</i> . <sup>d</sup>
<i>Oxyporus populinus</i> (Fr.) Donk.	Causes white heart rot, especially in the basal part of the stem. <sup>d</sup>
<i>Pholiota squarrosa</i> Huds. ex Fr.	Infests weakened beeches. <sup>g</sup>
<i>Polyporus squamosus</i> Huds. ex Fr.	Cause white rot, often along the wounds where spores colonized the stem. <sup>d</sup>
<i>Pleurotus ostreatus</i> (Jacq. ex Fr.) Kummer	Often found on dying or dead standing deciduous broadleaf trees. <sup>d</sup>



### 470 4.3 **Limits of the study**

471 The retrospective approach of this study combined with the resulting rapidity of post-  
472 fire beech mortality may have caused a bias in the possibility and precision of  
473 assessing pre-fire stand characteristics and tree decaying-related variables, especially  
474 for fire sites older than 20 years post-fire. This may account for the decreasing  
475 explanatory power of models with increasing rapidity of post-fire stand dynamics,  
476 even though we discarded problematic variables such as the proportion of bark  
477 damage by fire and fire scar height. In fact, while 38% of the variance in tree  
478 mortality was explained in the low-model, explanatory power decreased to 32% in the  
479 moderate-model and dropped to 17% in the high-model. In contrast, the control-  
480 model reached an explanatory power of 66%.

481 Furthermore, fungi were not systematically assessed at the species level on each tree,  
482 which hampered our ability to provide a detailed analysis of the timing and impact of  
483 single species in the decay process, and the role of species occurrence under specific  
484 post-fire environmental conditions.

## 485 **5 Conclusions**

486 In this study, we used a retrospective approach for reconstructing the post-fire  
487 dynamics and main drivers of fire-related beech mortality in 36 differently-aged fire  
488 sites in the Southern Alps. Despite the discussed methodological limits of the  
489 approach, we were able to answer the main questions of the study, providing  
490 important new insights regarding beech fire ecology and post-fire beech mortality in  
491 particular.

492 Fire-induced beech mortality was found to be very high within the first two decades  
493 post-fire. Thus, the process can be explained in terms of a dynamic phase (< 20 years

494 post fire), when beeches mostly die, and a more stable phase (> 20 years post-fire)  
495 when nearly no so-far-surviving beeches die as a consequence of fire impact.  
496 Consequently, gap formation is gradual, providing a 20-year window for beech  
497 regeneration under increasing light conditions. This finding is useful for the  
498 development of ecologically-based silvicultural post-fire treatments.

499 Major mortality drivers were found to be the combination of the proportion of woody  
500 tissue exposed as a consequence of fire scars and the likelihood of being colonized by  
501 active (fructifying) fungi. In the case of low to moderate burn severity, tree size may  
502 contribute to a reduction in the proportion of stems exposed to fungal infestation.  
503 Unfortunately, our research did not focus specifically on the role of single fungal  
504 species in the dieback process of fire-injured beeches. Further research is therefore  
505 needed on the temporal occurrence and ecology of specific post-fire fructifying fungi  
506 in beech forests.

507 This study demonstrates that beech can persist in a mixed-severity fire regime. Tall  
508 and large beeches survive for at least several years after low to moderate severe fires.  
509 These conditions mostly contribute to seed production and dispersal in mast years  
510 (e.g., Nilsson and Wastljung, 1987) and thus immediately provide seeds in the  
511 environment. At this stage, the seedbed is favorable to beech seed germination and  
512 seedling establishment (Ascoli et al., 2015).

513 From a practical point of view, it should be possible for forest managers to assess the  
514 burn severity class (ratio of dead to living trees) and related stand mortality dynamics  
515 within the third year post-fire. Target parameters for the evaluation of the mortality  
516 process are the diameters of survived trees in relation to the burn severity and  
517 evidence of open bark, which serves as entry points for fungal infection. Furthermore,  
518 our results show that very tall and large beeches mostly survive even after high severe



519 fires at least for several years. These remnant beeches are important as seed source for  
520 emerging regeneration. Therefore, they have to preserve when planning post-fire  
521 management measures.

## 522 **Acknowledgements**

523 This study was partially supported by the Swiss Federal Office for the Environment  
524 (FOEN). Fieldwork assistance was carried out with the support of Franco Fibbioli,  
525 Simone Giavi, Marianne Steffen, Lisa Berghäuser, Jordi Murgadas from the Swiss  
526 Federal Institute for Forest, Snow and Landscape Research, and Sven Hofmann from  
527 the University of Karlsruhe (Germany). Thanks go to Curtis Gautschi (professional  
528 linguist of the Swiss Federal Institute for Forest, Snow and Landscape Reseach  
529 (WSL)), who took care of the English version.

530

## 531 Appendix A

532 Table A.1: Investigated fire sites sorted by region (Piedmont, Ticino) and the date of fire. Items listed: years  
 533 post-fire (age), UTM coordinates (WGS84),  $\bar{\theta}$  annual temperature (T),  $\Sigma$  annual precipitation (P) (T and P:  
 534 WorldClim data base; Hijmans et al., 2005), and the number of plots investigated in the burnt ( $N_b$ ) and  
 535 unburnt beech forests ( $N_c$ ).

Regions	Municipality	Date of fire	Age	E	N	T [°C]	P [mm]	$N_b/N_c$
Piedmont								
	Sparone	28.12.80	34	382545	5030710	6	1109	16/1
	Rosazza	19.01.90	24	418645	5058661	5.8	1195	5/0
	Corio	15.02.90	24	385562	5021543	7.5	989	10/2
	Arola	04.06.97	16.5	449208	5074546	7.9	1172	13/0
	Dissimo	06.04.03	11	466503	5111215	5	1402	5/1
	Varallo	11.08.03	10.5	442360	5078456	7.2	1186	11/1
	Villadossola	16.03.05	9	440231	5098748	5.6	1305	11/1
	Condove	01.03.08	7	364870	5000781	7.4	979	11/1
	Giaglione	03.03.12	2	341650	5001664	6.4	1067	8/1
	Druogno	26.03.12	2	453207	5110682	4.8	1394	12/1
Ticino								
	Indemini	07.08.70	42.5	488196	5105864	6.4	1349	3/1
	Minusio	04.11.71	41	484123	5116368	4.7	1415	2/1
	Gordevio	09.03.73	40	482190	5116678	6.5	1355	1/0
	Moghegno	27.11.73	39	492538	5101434	8.3	1310	3/1
	Gordola	28.03.76	37	490491	5116753	6.0	1365	2/1
	Arbedo	20.03.76	37	506667	5116933	7.1	1290	13/1
	Astano	01.01.81	32	485796	5096454	8.2	1304	2/1
	Indemini	01.01.81	32	484488	5104578	5.5	1376	12/1
	Intragna	04.01.87	27	477570	5112256	7.6	1318	3/0
	Aurigeno	01.08.89	23.5	478824	5118037	8.2	1308	2/1
	Mugena	23.03.90	23	492683	5105828	7.1	1330	6/1
	Novaggio	10.03.90	23	486829	5098133	5.4	1371	2/1
	Avegno	05.05.90	23	482007	5116521	6.5	1355	2/0
	Pollegio	09.04.95	18	492574	5139100	5.3	1391	5/2
	Tenero	21.04.96	17	487212	5116007	8.5	1315	3/0
	Ronco s.A.	15.03.97	16	477225	5110649	6.6	1349	6/1
	Magadino	15.04.97	16	491560	5107650	6.9	1335	26/3
	Sonvico	03.04.97	16	501239	5101934	8.8	1300	5/2
	Arbedo	14.11.98	14	506770	5115571	8.5	1302	3/2
	Indemini	19.12.98	14	488487	5106098	6.6	1347	1/1
	Gordevio	24.04.02	11	482190	5116678	6.5	1355	13/4
	Maggia	12.03.02	11	477394	5124084	5.7	1388	3/1
	Bodio	18.03.03	10	495105	5136703	4	1436	5/1
	Someo	06.08.03	9.5	475281	5126733	5.6	1395	3/1
	Cugnasco	03.04.06	7	494084	5114855	9.4	1317	4/1
	Ronco s.A.	23.04.07	6	477225	5110649	6.6	1349	2/1

## 536 Appendix B

537 Table B.1: Distribution of mature tree species in the burnt and unburnt forests sorted by the target species  
 538 (beech), and trees showing wind-, gravity-, and animal seed dispersal strategies. The amount of dead trees  
 539 related to the total number of trees of a particular species ( $\Sigma N$ ) is expressed in the proportion of mortality  
 540 (mort. [%]). Species proportion [%] indicates the proportion of particular species out of the total number of  
 541 trees.

Species	Burnt forests			Unburnt forests		
	$\Sigma N$	Mort. [%]	Species proportion [%]	$\Sigma N$	Mort. [%]	Species proportion [%]
<b>Target species</b>						
<i>Fagus sylvatica</i> L.	2845	53	88	887	13	93
<b>Pioneers with wind-dispersal seeds</b>						
<i>Betula pendula</i> ROTH	129	44	4	20	30	2
<i>Larix decidua</i> MILL.	66	62	2	14	36	1
<i>Sorbus aria</i> (L.) CRANTZ	16	69	1	5	0	<1
<i>Alnus glutinosa</i> (L.) GAERTN.	4	75	<1	0	100	0
<i>Corylus avellana</i> L.	1	0	<1	0	100	0
<i>Populus tremula</i> L.	1	0	<1	0	100	0
<i>Sorbus aucuparia</i> L.	1	0	<1	0	0	0
<i>Laburnum alpinum</i> FABR.	0	0	0	14	64	1
<b>Trees with gravity- /animal-dispersal seeds</b>						
<i>Castanea sativa</i> MILL.	57	70	2	11	9	1
<i>Quercus petraea</i> (MATTUSCHKA)	30	40	1	2	0	<1
<i>Fraxinus excelsior</i> L.	6	33	<1	0	100	0
<i>Picea abies</i> (L.) H.KARST.	6	0	>1	0	100	0
<i>Pinus sylvestris</i> L.	3	0	<1	0	100	0
<i>Prunus avium</i> L.	2	0	<1	4	75	<1
<i>Taxus baccata</i> L.	2	0	<1	0	100	0
<i>Acer pseudoplatanus</i> L.	1	0	<1	1	0	<1
<i>Quercus pubescens</i> WILLD.	1	100	<1	0	0	0

542

543

544

545 **References**

546

547 Adel, M.N., Pourbabaei, H., Omid, A., Dey, D.C., 2013. Forest structure and woody  
548 plant species composition after a wildfire in beech forests in the north of Iran. *Journal*  
549 *of Forestry Research* 24, pp. 255–262. DOI: 10.1007/s11676-012-0316-7.

550 Ascoli, D., Castagneri, D., Valsecchi, C., Conedera, M., Bovio, G., 2013. Post-fire  
551 restoration of beech stands in the Southern Alps by natural regeneration. *Ecological*  
552 *Engineering* 54, pp. 210–217. DOI: 10.1016/j.ecoleng.2013.01.032.

553 Ascoli, D., Vacchiano, G., Maringer, J., Bovio, G., Conedera, M., 2015. The  
554 synchronicity of masting and intermediate severity fire effects favors beech  
555 recruitment. *Forest Ecology and Management* 353, pp. 126–135. DOI:  
556 10.1016/j.foreco.2015.05.031.

557 Bachelet, D., Lenihan, J.M., Neilson, R.P., 2007. Wildfires and global climate  
558 change: The importance of climate change for future wildfire scenarios in the Western  
559 United States. *Pew Center on Global Climate Change*, Arlington (USA)

560 Bauer, G., Speck, T., Blömer, J., Bertling, J., Speck, O., 2010. Insulation capability of  
561 the bark trees with different fire adaption. *Journal of Materials Science* 45, pp. 5950–  
562 5959. DOI: 10.1007/s10853-010-4680-4.

563 Beniston, M., 2004. The 2003 heat wave in Europe: A shape of things to come? An  
564 analysis based on Swiss climatological data and model simulations. *Geophysical*  
565 *Research letters* 31, pp. L02202. DOI: 10.1029/2003GL018857.

566 Boeck, A., Dieler, J., Biber, P., Pretzsch, H., Ankerst, D.P., 2014. Predicting tree  
567 mortality for European beech in Southern Germany using spatially explicit  
568 competition indices. *Forest Science* 60, pp. 613–622. DOI: 10.5849/forsci.12-133.

569 Bonyad, A.E., Sima, A., Bakhshandeh, A., Dadras, H., 2012. Evaluation of non-  
570 destructive Meyer method for determination of bark volume of beech (*Fagus*  
571 *orientalis* Lipsky) in different geographical aspects. Caspian Journal of  
572 Environmental Science 10, pp. 67–73.

573 Brando, P.M., Nepstad, D.C., Balch, J.K., Bolker, B., Christman, M.C., Coe, M. et al,  
574 2012. Fire-induced tree mortality in a neotropical forest: the roles of bark traits, tree  
575 size, wood density and fire behavior. Global Change Biology 18, pp. 630–641. DOI:  
576 10.1111/j.1365-2486.2011.02533.x.

577 Brown, J., Reinhard, M., Kramer, K.A., 2003. Coarse woody debris: Managing  
578 benefits and fire hazard in the recovering forest. USDA Forest Service – Rocky  
579 Mountain research Station. General Technical Report RMRS-GTR-105. URL: <  
580 [http://www.fs.fed.us/rm/pubs/rmrs\\_gtr105.pdf](http://www.fs.fed.us/rm/pubs/rmrs_gtr105.pdf)> (26.10.2015).

581 Brown, M.J., Kertis, J., Huff, M.H., 2013. Natural tree regeneration and coarse woody  
582 debris dynamics after a forest fire in the Western Cascade Range, Research Paper  
583 PNW-RP 592, Portland. URL: < [http://www.fs.fed.us/pnw/pubs/pnw\\_rp592.pdf](http://www.fs.fed.us/pnw/pubs/pnw_rp592.pdf)>.  
584 (26.10.2015).

585 Brownrigg, R., 2015. Draw Geographical Maps. Version 3.0.0-2. R Core Team. URL:  
586 <https://cran.r-project.org/web/packages/maps/maps.pdf>. (26.10.2015).

587 Brunsdon, C., Chen, H., 2015. Package `GISTools'. Version 0.7-4. R Core Team.  
588 URL: <https://cran.r-project.org/web/packages/GISTools/index.html>. (26.10.2015).

589 Camerano, P., Gottero, F., Terzuolo, P., Varese, P., 2004. Tipi forestali del Piemonte.  
590 Blu Edizioni, Torino.

591 Catry, F.X., Rego, F., Moreira, F., Fernandes, P.M., Pausas, J.G., 2010. Post-fire tree  
592 mortality in mixed forests of central Portugal. *Forest Ecology and Management* 7, pp.  
593 1184–1192. DOI: 10.1016/j.foreco.2010.07.010.

594 Ceschi, I., 2006. *Il bosco nel Canton Ticino*. Armando Dadó Editore, Locarno.

595 Conedera, M., Lucini, L., Holdenrieder, O., 2007. Pilze als Pioniere nach Feuer. *Wald*  
596 *und Holz* 11, pp. 45–48.

597 Conedera, M., Lucini, L., Valse, E., Ascoli, D., Pezzatti, G. (Eds.), 2010. Fire  
598 resistance and vegetative recruitment ability of different deciduous tree species after  
599 low- to moderate-intensity surface fires in southern Switzerland. In: Viegas, D.X. (ed)  
600 VI International Conference on Forest Fire Research, 15-18 November 2010,  
601 Coimbra, Portugal. [CD-ROM]. Portugal, ADAI/CEIF University of Coimbra.

602 Conedera, M., Tonini, M., Oleggini, L., Orozco, C.V., Leuenberger, M., Pezzatti, G.,  
603 2015. Geospatial approach for defining the Wildland-Urban Interface in the Alpine  
604 environment. *Computers, Environment and Urban Systems* 52, pp. 10–20.

605 Deepayan, S., 2008. *Lattice: multivariate data visualization with R*. Springer, New  
606 York (USA).

607 Dymytrova, L., Nadyeina, O., Hobi, M.L., Scheidegger, C., 2014. Topographic and  
608 forest-stand variables determining epiphytic lichen diversity in the primeval beech  
609 forest in the Ukrainian Carpathians. *Biodiversity Conservation* 23, pp. 1367–1394.  
610 DOI: 10.1007/s10531-014-0670-1.

611 Eidg. Anstalt für das forstliche Versuchswesen (EAFV). 1983. *Ertragstabellen Buche*.  
612 EAFV (Eds.), Birmensdorf (Switzerland).

613 ESRI. *ArcGIS Desktop: Release 10*. (Environmental Systems Research Institute:  
614 Redlands, CA).

615 Fernandes, P.M., Vega, J.A., Jiménez, E., Rigolot, E., 2008. Fire resistance of  
616 European pines. *Forest Ecology and Management* 256, pp. 246–255. DOI:  
617 10.1016/j.foreco.2008.04.032.

618 Fischer, R., Lorenz, M., Granke, O., Mues, V., Iost, S., van Dobben, H. et al, 2010.  
619 Forest condition in Europe, 2010 Technical Report of ICP Forests, Hamburg.

620 Frehner, M., Wasser, B., Schwitter, R., 2005. Nachhaltigkeit und Erfolgskontrolle im  
621 Schutzwald. Wegleitung für Pflegemassnahmen in Wäldern mit Schutzfunktion.  
622 Bundesamt für Umwelt, Wald, Landschaft: Bern.

623 Gerhardt, E. 2005. Guida ai funghi. Zanichelli: Bologna.

624 Gutsell, S.L., Johnson, E.A., 1996. How fire scars are formed: coupling a disturbance  
625 process to its ecological effect. *Can. J. For. Res.* 26, pp. 166–174. DOI: 10.1139/x26-  
626 020.

627 Harrod, J.C., Harmon, M.E., White, P., 2000. Post-fire succession and 20th century  
628 reduction in fire frequency and xeric southern Appalachian sites. *Journal of*  
629 *Vegetation Science* 4, pp. 465–472. DOI: 10.2307/3246576.

630 Hicks, R.R., 1998. Ecology and Management of central hardwood forests. John Wiley  
631 & Sons, New York (USA).

632 Hijmans, R.J., Cameron, S.E., Parra, J.L., Jones, P.G., Jarvis, A., 2005. Very high  
633 resolution interpolated climate surfaces for global land areas. *International Journal of*  
634 *Climatology* 25, pp. 1965–1978. DOI: 10.1002/joc.1276.

635 Hood, S.M., Smith, S.L., Cluck, D.R., 2007. Delayed conifer tree mortality following  
636 fire in California. USDA Forest Service, pp. 261–283. URL: <  
637 [http://www.fs.fed.us/psw/publications/documents/psw\\_gtr203/psw\\_gtr203\\_019hood.](http://www.fs.fed.us/psw/publications/documents/psw_gtr203/psw_gtr203_019hood.pdf)  
638 [pdf](http://www.fs.fed.us/psw/publications/documents/psw_gtr203/psw_gtr203_019hood.pdf). > (26.10.2015).

639 IPCC 2014. Pachauri R.K., Allen M.R., Barros V.R., Broome J., Cramer W., Christ  
640 R., Church J.A., Clarke L., Dahe Q., Dasgupta P., Dubash N.K., Edenhofer O.,  
641 Elgizouli I., Field C.B., Forster P., Friedlingstein P., Fuglestvedt J., Gomez-Echeverri  
642 L., Hallegatte S., Hegerl G., Howden M., Jiang K., Cisneros B.J., Kattsov V., Lee H.,  
643 Mach K.J., Marotzke J., Mastrandrea M.D., Meyer L., Minx J., Mulugetta Y., O'Brien  
644 K., Oppenheimer M., Pereira J.J., Pichs-Madruga R., Plattner G.-K., Pörtner H.-O.,  
645 Power S.B., Preston B., Ravindranath N.H., Reisinger A., Riahi K., Rusticucci M.,  
646 Scholes R., Seyboth K., Sokona Y., Stavins R., Stocker T.F., Tschakert P., van  
647 Vuuren D., van Ypersele J.-P. (2014) *Climate Change 2014: Synthesis Report*. (Eds  
648 R.K. Pachauri, L. Meyer). Cambridge University Press: Cambridge (UK), New York  
649 (USA).

650 Isotta, F.A., Frei, C., Weilguni, V., Perc'ec Tadić, M., Lassègues, P., Rudolf, B. et al,  
651 2014. The climate of daily precipitation in the Alps: development and analysis of a  
652 high-resolution grid dataset from pan-Alpine rain-gauge data. *International Journal of*  
653 *Climatology* 34, pp. 1657–1675. DOI: 10.1002/joc.3794

654 Kahl, T., 2008. Kohlenstofftransport aus dem Totholz in den Boden. PhD-Thesis.  
655 Universität Freiburg, Freiburg i. Br.

656 Keyser, T.L., Lentile, L.B., Smith, F.W., Shepperd, W.D., 2008. Changes in forest  
657 structure after a large, mixed-severity wildfire in Ponderosa Pine Forests of the Black  
658 Hills, South Dakota, USA. *Forest Science* 54, pp. 328–338.

659 Klug, P., Lewald-Brudi, M. 2012. *Holzzersetzende Pilze*. Arbus-Verlag: Bad-Boll.

660 Kobziar, L., Moghaddas, J., Stephens, S.L., 2006. Tree mortality patterns following  
661 prescribed fires in a mixed conifer forest. *Can. J. For. Res.* 36, pp. 3222–3238. DOI:  
662 10.1139/x06-183.



663 Krieglsteiner, G.J. (Ed.), 2000. Die Großpilze Baden-Württembergs. Erwin Ulmer  
664 Verlag, Stuttgart.

665 Lampainen, J., Kuuluvainen, T., Wallenius, T., Karjalainen, L., Vanha-Majamaa, I.,  
666 2004. Long-term forest structure and regeneration after wildfire in Russian Karelia.  
667 Journal of Vegetation Science 15, pp. 245–256. DOI: 10.1111/j.1654-  
668 1103.2004.tb02259.x.

669 Lawes, M.J., Midgley, J.J., Clarke, P.J., 2013. Costs and benefits of relative bark  
670 thickness in relation to fire damage: a savanna/forest contrast. Journal of Ecology  
671 101, pp. 517–524. DOI: 10.1111/1365-2745.12035.

672 Ledgard, N., Davis, M., 2004. Restoration of mountain beech (*Nothofagus solandri*  
673 var. *cliffortioides*) forest after fire. New Zealand Journal of Ecology 28, pp. 125–135.

674 Lee, M.-W., Hyeon-Hur, Chang, K.-C., Lee, T.-S., Ka, K.-H., Jankovsky, L., 2008.  
675 Introduction to distribution and ecology of Sterile Conks of *Inonotus obliquus*.  
676 Mycobiology 36, pp. 199–202. DOI: 10.4489/MYCO.2008.36.4.199.

677 Lines, E.R., Coomes, D.A., Purves, D.W., 2010. Influences of forest structure, climate  
678 and species composition on tree mortality across the Eastern US. PlosOne 5(10). Pp.  
679 e13212. DOI:10.1371/journal.pone.0013212.

680 Lombardi, F., Cherubini, P., Lasserre, B., Tognetti, R., Machetti, M., 2008. Tree rings  
681 used to assess time since death of deadwood of different decay classes in beech and  
682 silver fir forests in the central Apennines (Molise, Italy). Can. J. For. Res. 38, pp.  
683 821–833. DOI: 10.1139/X07-195.

684 Maringer, J., Conedera, M., Ascoli, D., Schmatz, D.R., Wohlgemuth, T., subm.  
685 Resilience of European beech forests (*Fagus sylvatica* L.) after fire in a global change  
686 context. International Journal of Wildland Fire.

687 McHugh, C.W., Kolb, T.E., 2003. *Ponderosa pine* mortality following fire in northern  
688 Arizona. *International Journal of Wildland Fire* 12, pp. 7–12. DOI:  
689 10.1071/WF02054.

690 Moreira, F., Arianoutsou, M., Corona, P., De las Heras, J., 2012. Post-fire  
691 management and restoration of Southern European forests. Springer, Netherlands.

692 Nicolai, V., 1986. The bark of trees: thermal properties, microclimate and fauna.  
693 *Oecologia* 69, pp. 148–160.

694 Nilsson, S., Wastljung, U., 1987. Seed predation and cross-pollination in mast-  
695 seeding beech (*Fagus sylvatica*) patches. *Ecology* 68, pp. 260–265.

696 Nocentini, S., 2009. Structure and management of beech (*Fagus sylvatica* L.) forests  
697 in Italy. *iForest* 2, pp. 105–113. DOI: 10.3832/ifor0499-002.

698 Odhiambo, B., Meincken, M., Seifert, T., 2014. The protective role of bark against  
699 fire damage: a comparative study on selected introduced and indigenous tree species  
700 in the Western Cape, South Africa. *Trees* 28, pp. 555–565. DOI: 10.1007/s00468-  
701 013-0971-0

702 Packham, J.R., Thomas, P., Atkinson, M., Degen, T., 2012. Biological flora of the  
703 British Isles: *Fagus sylvatica*. *Journal of Ecology*, pp. 1557–1608. DOI:  
704 10.1111/j.1365-2745.2012.02017.x.

705 Peters, R., 1997. *Beech forests*. Kluwer, Dordrecht.

706 Pezzatti, G., Reinhard, M., Conedera, M., 2010. Swissfire: Die neue schweizerische  
707 Waldbranddatenbank. *Swiss Forestry Journal* 161, pp. 465–469.

708 Pezzatti, G., Zumbrunnen, T., Bürgi, A., Ambrosetti, P., Conedera, M., 2013. Fire  
709 regime shifts as a consequence of fire policy and socio-economic development: An

710 analysis based on the change point approach. *Forest Policy and Economics* 29, pp. 7–  
711 18. DOI: 10.1016/j.forpol.2011.07.002.

712 Pinheiro, J., Bates, D., DebRoy, S., Sarkar, D., EISPACk, 2015. *Linear and*  
713 *Nonlinear Mixed Effects Models*.

714 R Development Core Team, 2014. *R: A language and environment for statistical*  
715 *computing*, Vienna, Austria.

716 Reinartz, H., Schlag, M., 2002. *Baumschädigende Pilze: Wirkungs- und*  
717 *Ausbreitungsweise baumschädigender Pilze zur Einschätzung der Höhe von*  
718 *Wertminderungen im Rahmen der Gehölzwertermittlung*, Köln.

719 Rigolot, E., 2004. Predicting post-fire mortality of *Pinus halepensis* Mill. and *Pinus*  
720 *pinea* L. *Plant Ecol* 171, pp. 139–151.

721 Russo, D., Cistrone, L., Garonna, A.P., 2010. Habitat selection by the highly  
722 endangered long-horned beetle *Rosalia alpina* in Southern Europe: a multiple spatial  
723 scale assessment. *Journal of Insect Conservation* 15, pp. 685-693. DOI:  
724 10.1007/s10841-010-9366-3.

725 Sarris, D., Christopoulou, A., Angelonidi, E., Koutsias, N., Fulé, P., Arianoutsou, M.,  
726 2014. Increasing extremes of heat and drought associated with recent severe wildfires  
727 in southern Greece. *Regional Environmental Change* 14, pp. 1257–1268. DOI:  
728 10.1007/s10113-013-0568-6.

729 Schumacher, S., Bugmann, H., 2006. The relative importance of climatic effects,  
730 wildfires and management for future forest landscape dynamics in the Swiss Alps.  
731 *Global Change Biology* 12, pp. 1435–1450. DOI: 10.1111/j.1365-2486.2006.01188.x.

732 Schwarze, F., Baum, S., 2000. Mechanisms of reaction zone penetration by decay  
733 fungi in wood of beech (*Fagus sylvatica*). *New Phytologist* 146, pp. 129–140. DOI:  
734 10.1046/j.1469-8137.2000.00624.x.

735 Shafiei, A., Akbarinia, M., Jalali, G., Hosseini, M., 2010. Forest fire effects in beech  
736 dominated mountain forest of Iran. *Forest Ecology and Management* 259, pp. 2191–  
737 2196. DOI: 10.1016/j.foreco.2010.02.025.

738 Shekholeslami, A., Kazemnezhad, F., Akhshabi, S., 2011. Bark measurement of  
739 beech (*Fagus orientalis* Lipsky.) in Tosakoti - Hyrcanian Forest. *International Journal*  
740 *of Forest, Soil and Erosion* 1, pp. 1–4.

741 Shigo, A.L., 1970. Growth of *Polyporus glomeratus*, *Poria obliqua*, *Fomes igniarius*,  
742 and *Pholiota squarrosa-adiposa* in media amended with manganese, calcium, zinc,  
743 and iron. *Mycologia* 62, pp. 604–607. DOI: 10.2307/3757536.

744 Sieg, C.H., McMillin, J.D., Fowler, J.F., Allen, K.K., Negron, J.F., Wadleigh, L.L. et  
745 al, 2006. Best predictor for post-fire mortality of Ponderosa Pine Trees in the  
746 Intermountain West. *Forest Science* 52, pp. 718–728.

747 Standovár, T., Kenderes, K., 2003. A review on natural stand dynamics in  
748 beechwoods of East Central Europe. *Applied Ecology and Environmental Research* 1,  
749 pp. 19–46.

750 Stevens-Rumann, C., Sieg, C.H., Hunter, M.E., 2012. Ten years after wildfires: How  
751 does varying tree mortality impact fire hazard and forest resiliency? *Forest Ecology*  
752 *and Management* 267, pp. 199–208. DOI: 10.1016/j.foreco.2011.12.003.

753 Sullivan, B.W., Kolb, T.E., Hart, S.C., Kaye, J.P., Hungate, B.A., Dore, S. et al, 2011.  
754 Wildfire reduces carbon dioxide efflux and increases methane uptake in ponderosa

755 pine forest soils of the southwestern USA. *Biogeochemistry* 104, pp. 251–265. DOI:  
756 10.1007/s10533-010-9499-1

757 Tubbs, C.H., Houston, D., 1990. American Beech (*Fagus grandifolia* Ehrh.), *Silvics*  
758 of North America. Vol. 2 Hardwoods. Agric. Handbook 654.

759 Valese, E., Conedera, M., Held, A.C., Ascoli, D., 2014. Fire, humans and landscape  
760 in the European Alpine region during the Holocene. *Anthropocene* 6, pp. 63–74. DOI:  
761 dx.doi.org/10.1016/j.ancene.2014.06.006.

762 van Couwenberghe, R., Collet, C., Lacombe, E., Pierrat, J.-C., Gégout, J.-C., 2010.  
763 Gap partitioning among temperate tree species across a regional soil gradient in  
764 windstorm-disturbed forests. *Forest Ecology and Management* 260, pp. 146–154.  
765 DOI: 10.1016/j.foreco.2010.04.013.

766 van Mantgem, P.J., Nesmith, J., Keifer, M.B., Brooks, M., 2013. Tree mortality  
767 patterns following prescribed fire for *Pinus* and *Abies* across the southwestern United  
768 States. *Forest Ecology and Management* 289, pp. 463–469. DOI:  
769 10.1016/j.foreco.2012.09.029.

770 Venables, W.N., Ripley, B.D., 1999. *Modern Applied Statistics with S-Plus*. Springer,  
771 New York (USA).

772 Wagner, S., Collet, C., Madsen, P., Nakashizuka, T., Nyland, R., Sagheb-Talebi, K.,  
773 2010. Beech regeneration research: from ecological to silvicultural aspects. *Forest*  
774 *Ecology and Management* 259, pp. 2172–2182. DOI: 10.1016/j.foreco.2010.02.029.

775 Webster, J., Weber, R., 2007. *Introduction to fungi*, 3rd. Cambridge University Press,  
776 Cambridge (UK).

777 Westerling, A.L., Hidalgo, H.G., Cayan, D.R., Swetnam, T.W., 2006. Warming and  
778 earlier spring increase Western U.S. forest wildfire activity. *New Series* 313, pp. 940–  
779 943. DOI: 10.1126/science.1128834

780 Wickham, H., Chang, W., 2015. Package `ggplot2`. URL: < [https://cran.r-](https://cran.r-project.org/web/packages/ggplot2/ggplot2.pdf)  
781 [project.org/web/packages/ggplot2/ggplot2.pdf](https://cran.r-project.org/web/packages/ggplot2/ggplot2.pdf).> (15.10.2015).

782 Yaussy, D.A., Hix, D.M., Long, R.P., Goebel, C.P. (Eds.), 2004. Prescribed surface-  
783 fire tree mortality in Southern Ohio: equations based on the thermocouple probe  
784 temperatures, Newtown Square, PA.

785 Yee, T.W., Stoklosa, J., Huggins, R.M., 2015. The VGAM package for capture-  
786 recapture data using the conditional likelihood. *Journal of Statistical Software*, pp. 1–  
787 33.

788 Z'Graggen, S., 1992. Dendrohistometrisch-klimatologische Untersuchung an Buchen  
789 (*Fagus sylvatica* L.). Universität Basel, Basel.

790 Zarzyński, P., 2007. The range of trophic preferences of oak mazelgill (*Daedalea*  
791 *quercina* (L.): Fr.) isolate examined in Vitro. *Acta Sci. Pol. Silv. Colendar. Rat. Ind.*  
792 *Lignar* 6, pp. 112–118.

793 Zuur, A., Ieno, E.N., Elphick, C.S., 2010. A protocol for data exploration to avoid  
794 common statistical problems. *Methods in Ecology and Evolution*, pp. 3–14. DOI:  
795 10.1111/j.2041-210X.2009.00001.x.

796 **Web references**

797 MeteoSwiss (2015) Swiss climate. Federal Office of Meteorology and Climatology.  
798 Zürich, Switzerland. <<http://www.meteoschweiz.admin.ch/home.html?tab=overview>,  
799 updated on 2015>, (accessed 1.02.15).

800 Arpa Piemonte. <<http://www.arpa.piemonte.it/banca-dati-meteorologica>>

801 (accessed 1.02.15).

802