

*Nutrient enrichment diminishes plant diversity and density, and alters long-term ecological trajectories, in a biodiverse forest restoration*

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Accepted Version

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Daws, M. I., Walters, S. J., Harris, R. J., Tibbett, M. ORCID: <https://orcid.org/0000-0003-0143-2190>, Grigg, A. H., Morald, T. K., Hobbs, R. J. and Standish, R. J. (2021) Nutrient enrichment diminishes plant diversity and density, and alters long-term ecological trajectories, in a biodiverse forest restoration. *Ecological Engineering*, 165. 106222. ISSN 0925-8574 doi: 10.1016/j.ecoleng.2021.106222 Available at <https://centaur.reading.ac.uk/97548/>

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To link to this article DOI: <http://dx.doi.org/10.1016/j.ecoleng.2021.106222>

Publisher: Elsevier

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1 **Nutrient enrichment diminishes plant diversity and density, and alters long-term**  
2 **ecological trajectories, in a biodiverse forest restoration**

3

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20

21

## 22 ABSTRACT

23 Nutrient enrichment can negatively affect natural plant communities and result in the loss of  
24 species diversity and productivity. Despite this, fertiliser (especially phosphorus) is typically  
25 applied to restore highly biodiverse communities. Long-term effects of nutrient addition to  
26 restored plant communities, particularly those adapted to inherently low nutrient soils, have  
27 received little attention. We report results of a large-scale 20-year field experiment  
28 established in West Australian jarrah forest restored after bauxite mining. Three P-application  
29 rates were applied (0, 80 and 120 kg ha<sup>-1</sup>) once at the beginning of the experiment, and plant  
30 communities monitored after 1, 6, 13 and 20 years. One year after the onset of restoration,  
31 native plant species richness and plant density was highest at 80 and 120 kg P ha<sup>-1</sup>.  
32 Subsequently, native species richness, plant density, and the richness and density of seeder  
33 and slow-growing resprouter species were highest without fertilisation, establishing the  
34 negative impact of P enrichment on plant community and ecosystem development in P  
35 impoverished soils. Total plant cover was similar for all P treatments across the  
36 chronosequence which, when combined with higher stem densities at zero P, suggests zero P  
37 favoured smaller, slower growing species. Applied-P initially favoured weeds and  
38 ephemerals and, while these species declined over time, other species were lost from these  
39 plots. The similarity of the restored communities to unmined reference jarrah forest increased  
40 over time and was consistently highest at in the absence of P fertiliser. Jarrah forest  
41 restoration is assumed to follow the initial floristic model of plant succession. However, we  
42 question this assumption and instead suggest that successional outcomes are contingent on P  
43 fertilisation rather than initial floristics per se., Applied P retarded recruitment of resprouter  
44 species that were present at zero P, debunking the assumption under IFM that these species  
45 do not disperse to restored areas. Consequently, based on the most comprehensive long-term  
46 study of P-fertilisation in the context of restoration of P-impoverished ecosystems yet

47 reported, we propose that P limitation is important for the recreation of species diversity in  
48 inherently P impoverished forests. These results highlight the necessity of long-term  
49 experiments for understanding forest successional dynamics and implications for restoration  
50 practices.

51

52 Key words: competition, initial floristics model, legume, nitrogen, rehabilitation

53

## 54 **1. Introduction:**

55 Studies in a range of native ecosystems have shown long-term negative effects of nutrient  
56 enrichment on plant community composition, species richness and ecosystem productivity  
57 (Wheeler and Shaw, 1991; Wassen et al., 2005; Isbell et al., 2013). Understanding this  
58 relationship is key to restoring historic plant communities because soil nutrients are often  
59 modified by human activity, either enriched in the case of old-fields (e.g., Standish et al.,  
60 2006; Tibbett et al., 2019a) or depleted in the case of mine sites (e.g. Prematuri et al., 2020).  
61 Consequently, adding fertiliser to restore nutrients lost during mining is generally viewed as a  
62 key step in restoring native plant communities (e.g., EPA, 1996; Bell, 2001; Tibbett, 2010;  
63 DFAT, 2016). Fertiliser addition can also increase under- and over-storey plant growth  
64 thereby having a range of potential benefits such as a reduced risk of soil erosion (Ward et  
65 al., 1990) and increased carbon sequestration and timber production (Brancalion et al., 2019).  
66 However, longer term effects of fertiliser addition on community composition in restored  
67 mine sites have received little attention. Long-term data are needed to establish this  
68 relationship, particularly for projects with goals focused on restoring community composition  
69 and species richness similar to undisturbed reference communities.

70       There is increasing evidence in post-mining restoration that fertiliser application may  
71 be detrimental, at least in the short-term, to re-establishing diverse native vegetation  
72 communities. For example, in three-year-old restored sites in the Cape Floristic Region,  
73 South Africa, NPK fertiliser increased weed growth, but decreased survival of proteaceous  
74 shrubs (Holmes, 2001) that are sensitive to high P (Stock and Allsopp, 1992). Similarly, up to  
75 five years following restoration, fertiliser-P addition increased the growth of weeds and  
76 native ephemerals in restored jarrah forest in Western Australia, while decreasing the  
77 abundance of long-lived resprouter species (Daws et al., 2013; Daws et al., 2015; Tibbett et  
78 al. 2020). Resprouters are a diverse and dominant component of undisturbed jarrah forest

79 (Norman et al., 2006a; Koch, 2007). Nitrogen fertiliser did not have the same detrimental  
80 short-term effects (Daws et al., 2013) perhaps because excess N-fertiliser can be rapidly lost  
81 from soils and instead, nitrogen is provided by early-successional legumes (Grant et al.,  
82 2007). In contrast, P-fertiliser can persist in restored soils for at least 20 years (Banning et al.,  
83 2008; Standish et al., 2008; Spain et al., 2018; Daws et al., 2019a) and therefore potentially  
84 affects vegetation dynamics in both the short and longer terms.

85         The jarrah forest occurs within one of the world's biodiversity hotspots (Hopper and  
86 Gioia, 2004) with 300–400 understorey plant species in areas where mining occurs (Koch,  
87 2007). Jarrah forest soils are highly weathered and naturally P-impooverished (Hopper, 2009).  
88 Consequently, many understorey species have specialised adaptations for P-acquisition  
89 including cluster roots, mycorrhizal symbioses and exudation of carboxylases (Lambers et al.,  
90 2008). These species can exhibit toxicity to applied-P at elevated external concentrations due  
91 to a limited ability to regulate P uptake (e.g., de Campos et al., 2013). Many Western  
92 Australian resprouter species also have low growth rates (Pate et al., 1990; Bowen, 1991;  
93 Bowen and Pate, 1993). Thus, there are two possible, non-exclusive hypotheses for negative  
94 effects of applied-P on restored jarrah forest communities. Firstly, effects on composition  
95 may result from species-specific direct negative effects of elevated P on plant survival (e.g.,  
96 Holmes, 2001). Secondly, since the growth of many resprouter species to applied-P is  
97 conservative compared with that of weeds, ephemerals and N<sub>2</sub>-fixing legumes (Daws et al.,  
98 2015; Standish et al., 2008). they may be simply out-competed in high P soils.

99         Jarrah forest restoration is assumed to follow the initial floristics model of plant  
100 succession (Norman et al., 2006a; Koch, 2007) whereby the composition of the developing  
101 vegetation reflects the initial community that establishes (Egler, 1954). To date, the  
102 assumption has been that this model holds because late-successional species do not arrive or  
103 arrive but fail to establish. Empirical support for the model comes from restored jarrah-forest

104 trajectories up to 14 years of age that received P-fertiliser at the onset (Norman et al., 2006a;  
105 Koch, 2007). Plant available soil-P can remain elevated for at least 20 years in restored jarrah  
106 forest (Banning et al., 2008; Daws et al. 2019a) following a single initial P-fertiliser  
107 application. Consequently, legacy effects of P-fertiliser could help explain why late arriving  
108 species do not establish: they may be outcompeted by species that respond to, and continue to  
109 benefit from, increased soil P. These possibilities have not been explored as long-term  
110 effects of P-fertiliser on plant community composition in restored jarrah forest sites have  
111 received little attention.

112         In this study, we report results from a long-term (20-year) experiment of the effects of a  
113 single initial application of P-fertiliser, at rates of 0, 80 or 120 kg P ha<sup>-1</sup>, on vegetation  
114 responses in restored jarrah forest. Plots were monitored 1, 6, 13 and 20 years after initial  
115 restoration. Specifically, we explore the effects of fertility (applied-P) on diversity and  
116 development of the understory community to assess whether fertility affects the diversity  
117 and trajectory of the developing community.

118

## 119 2. Materials and methods

### 120 2.1. Study site

121 The experiment was established within the Alcoa of Australia Ltd. bauxite mining lease in  
122 the northern jarrah forest, located 60 to 100 km south-east of Perth, Western Australia  
123 (Standish et al., 2015). The area has a Mediterranean-type climate with cool, wet winters and  
124 hot, dry summers. Annual rainfall is ~1,200 mm with average summer maximum  
125 temperatures of 29 °C and average winter minimum temperatures of 6 °C (Australian Bureau  
126 of Meteorology, 2015).

127 The forest overstorey vegetation comprises the dominant species *Eucalyptus*  
128 *marginata* (jarrah) and *Corymbia calophylla* (marri). There is a mid-storey layer dominated  
129 by *Banksia grandis*, *Allocasuarina fraseriana* and *Xanthorrhoea preisii* with woody shrubs  
130 of *Bossiaea aquifolium*, various *Acacia* species, and a diverse understorey (Koch, 2007).

131 Jarrah forest soils are gravelly with low concentrations of available N, P and K (Hingston et  
132 al., 1989; Tibbett et al., 2020) with high rates of phosphorus fixation due to the presence of  
133 amorphous iron and aluminium oxides. Generically these soils are classified as lateritic  
134 oxisols (USDA, 1999) or ferralsols (FAO, 2012).

135

### 136 2.2. Experimental design

137 A large-scale field experiment was established on recently mined pits. Restoration activities  
138 comprise pit landscaping, deep ripping, topsoil return, contour ripping and native flora  
139 seeding (Standish et al., 2015). Treatment plots of 25 m × 25 m were established in four  
140 replicate mine pits using a randomised complete block design. Plots were seeded during  
141 summer 1994 (January to April) with 1.44 kg ha<sup>-1</sup> jarrah seed, 0.60 kg ha<sup>-1</sup> marri seed and an  
142 understorey seed mix containing 0.72kg ha<sup>-1</sup> of legume species and 0.20 kg ha<sup>-1</sup> of non-  
143 legume species. Following seeding, fertiliser was applied at 0, 80 or 120 kg ha<sup>-1</sup> for P (as

144 double superphosphate) and 80 kg ha<sup>-1</sup> for N (as ammonium sulphate) (Lockley and Koch,  
145 1996).

146

### 147 *2.3. Vegetation monitoring*

148 In September 1995, one year after the establishment of the experiment, a 20 m × 20 m plot  
149 was established within the centre of each 25 m × 25 m plot and subdivided into twenty 2 m ×  
150 2 m quadrats with a total of 80 m<sup>2</sup> sampled per plot. Species identity and density were  
151 recorded for all shrub and understorey species in each quadrat with density summed for the  
152 entire plot. Plots were re-monitored in spring 2000, 2007 and 2014, i.e. at 6, 13 and 20 years  
153 of age, except that percentage cover (a visual estimate that included overhang) was also  
154 recorded for each species. Cover estimates accounted for vertical structuring of vegetation,  
155 such that the sum of all cover estimates could exceed 100%.

156

### 157 *2.4. Soil sampling*

158 In April 2014 six soil samples were collected from random locations, within each 20 m × 20  
159 m plot. Samples were collected at 0–10 cm depth from the furrows of the rip-lines and mixed.  
160 0–10 cm depth was chosen because jarrah forest soils have high rates of P fixation on  
161 amorphous iron and aluminium oxides. As a result, limited downward movement of P occurs  
162 through the soil profile (e.g. Tibbett et al., 2020). Soil was air-dried at 26 °C, gravel removed  
163 using a 2 mm sieve (Rayment and Higginson, 1992) and samples analysed at a commercial  
164 laboratory (CSBP Soil and Plant Laboratories, Bibra Lake, Perth, Australia) assessing  
165 ammonium (NH<sub>4</sub><sup>+</sup>), nitrate (NO<sub>3</sub><sup>-</sup>), Colwell(available)-P and pH (in water).

166

### 167 *2.5. Statistical analysis*

168 Each plant species was assigned to one of four categories reflecting hypothesised responses  
169 to P. We predicted that weeds and native ephemerals would respond positively to fertiliser  
170 application (Prober and Wiehl, 2012), resprouter species to respond negatively (Lambers et  
171 al., 2008) and that seeders would be mixed in their responses. Seeders must re-establish  
172 through germination and seedling establishment, whereas resprouters can re-establish by  
173 sprouting from surviving underground structures (Bell, 2001; Clarke et al. 2015). In Bell's  
174 categorisation, used for fire response, ephemerals and weeds are also seeders; whereas the  
175 seeder category used here consisted only of longer-lived species. Species were classified  
176 based on published literature (Bellairs and Bell, 1990; Bell et al., 1993; Ward et al., 1997;  
177 Smith et al., 2000; Norman et al., 2006b; Burrows et al., 2008; Western Australian  
178 Herbarium, 2012).

179 Analyses for the effects of P-application rate and restoration age on (1) species  
180 richness, (2) density, and (3) cover for all species combined, and the four growth forms were  
181 undertaken using linear mixed effects models in the lme4 package using the *lmer* function  
182 (Bates et al., 2011) for R (R Development Core Team 2009). P-application rate and site age  
183 were treated as fixed effects and site (pit) and subject (individual plot) as random effects.

184 To assess which species were associated with unfertilised plots after 20 years, species  
185 occurring more than four times in zero P plots and fewer than four times in 80 and 120 kg P  
186 ha<sup>-1</sup> plots were determined. For species associated with fertilised plots, the species that  
187 occurred more than four times in 80 and 120 kg P ha<sup>-1</sup> plots and fewer than four times in the  
188 zero P plots were determined (Standish et al., 2008).

189 A dissimilarity matrix using Sorensens dissimilarity was created based on species  
190 abundance data for the 18 reference forest plots using PC-ORD v. 6.0 (MjM Software,  
191 Gleneden Beach, OR, US), resulting in an overall forest plot: forest plot similarity value (1 –  
192 dissimilarity). Subsequently, for each experimental plot, pair-wise similarity values were

193 determined by comparison with each and every forest reference plot resulting in 18 pair-wise  
194 similarity values per experimental plot; the average of these values was then expressed as a  
195 percentage of the overall forest plot: forest plot similarity value.

196

197 **3. Results:**

198 *3.1. Soil analysis*

199 Twenty years after P-application, soil (Colwell) P concentrations were significantly higher as  
200 P-application rates increased (Table 1,  $P = 0.001$ ). P-application had no effect on soil  $\text{NH}_4^+$   
201 concentrations ( $P = 0.496$ ), and  $\text{NO}_3^-$  concentrations were lower than the detection limits of  
202 the analysis ( $\leq 1 \text{ mg kg}^{-1}$ ). Soil pH was significantly lower at 80 and 120 kg P  $\text{ha}^{-1}$  compared  
203 with zero P ( $P < 0.01$ , Table 1).

204

205 *3.2. Effects of applied-P and time since restoration on plant species richness*

206 After one-year, native plant species richness was lower at zero P than 80 or 120 kg P  $\text{ha}^{-1}$ .  
207 Subsequently, species richness increased at zero P, whilst generally declining in the 80 and  
208 120 kg P  $\text{ha}^{-1}$  treatments. By 6 years of age, richness was highest at zero P. This was  
209 maintained up to 20 years of age (Fig. 2A): after 20 years, there were 33.5 species at zero P  
210 compared with 20.4 and 24.5 at 80 and 120 kg P  $\text{ha}^{-1}$ , respectively. The effects of applied-P,  
211 time since restoration and the  $P \times \text{Time}$  interaction on species numbers were significant  
212 (Table 2).

213 After one year there were, on average, an additional 5.5 weed species at 80 and 120 kg P  $\text{ha}^{-1}$   
214 compared with zero P (Fig. 3A). For the remainder of the experiment, the number of weed  
215 species at 80 and 120 kg P  $\text{ha}^{-1}$  declined to be similar to those observed at zero P (Fig 3A),  
216 although even after 20 years three of the nine species associated with fertilised plots were  
217 weeds (Table 3). The effects of applied-P, time since restoration and the  $\text{Time} \times P$  interaction  
218 on the number of weed species were significant (Table 2). Ephemeral species exhibited a  
219 similar trend to weeds (Fig 3B). Time since restoration and the  $\text{Time} \times P$  interaction had a  
220 significant effect on the number of ephemeral species (Table 2): the effect of P was not  
221 significant (Table 2).

222           The number of seeder species was initially similar at all three P-application rates (Fig.  
223 3C). From six years onwards, the number of seeders declined, although this decline was least  
224 pronounced at zero P. As a result, species richness of this group at the 6, 13 and 20-year  
225 monitoring intervals was highest at zero P (Fig. 3C). The effects of time and P-application  
226 rate on species richness of seeders were significant (Table 2).

227           Species richness of resprouters was initially similar at all three P-application rates.  
228 Subsequently, the number of resprouters increased, particularly at zero P where the number  
229 of species more than doubled over the duration of the experiment (Fig. 3D). After 20 years,  
230 15 of the 19 species associated with the unfertilised plots were resprouters (Table 3). The  
231 effects of time since restoration, applied P and the Time  $\times$  P interaction on the number of  
232 resprouter species were significant (Table 2).

233

### 234 *3.3. Effects of applied-P and time since restoration on stem density and cover*

235           The number of individual native plants was initially (after one year) highest at 80 and 120 kg  
236 P ha<sup>-1</sup> (Figure 2B). Subsequently stem density increased at zero P while declining in the P  
237 addition treatments: after 20-years, stem density was nearly twice as high at zero P compared  
238 with either 80 or 120 kg P ha<sup>-1</sup>. This cross-over in density related to P-application rate was  
239 reflected in a significant Time  $\times$  P interaction (Table 2). The effect of time since restoration  
240 on the density of native species was also significant (Table 2).

241           Weed density exhibited little change over time (Fig 4A): neither the effects of  
242 applied-P nor time were significant (Table 2). After one year, applied-P resulted in ~2  
243 additional ephemeral plants m<sup>-2</sup> compared with zero P (Fig. 4B). Subsequently, the density of  
244 ephemerals increased slightly at zero P and declined in the two P-addition treatments. This  
245 result was supported by a significant effect of time and a significant Time  $\times$  P interaction  
246 (Table 2).

247 At 80 and 120 kg P ha<sup>-1</sup>, the density of seeder species exhibited little change over the  
248 20-year experiment (Fig. 4C). However, at zero P, seeder density increased significantly  
249 between six and 13 years, thereafter declining (Fig. 4C). The effects of both time and applied-  
250 P on seeder density were significant (Table 2). At all three P-application rates, stem density  
251 of resprouter species increased with time: the effect of time was highly significant (Table 2).  
252 However, the relative increase was greatest at zero P, where the resprouter density was  
253 initially lowest: density increased more than three-fold in this treatment (Fig 4D) and after 20  
254 years resprouter density was highest at zero P. Both the main effect of applied-P and the  
255 Time × P interaction were significant (Table 2).

256 Between 6 and 13 years, total plant cover approximately doubled in all three P  
257 treatments, before declining at 20 years (Figure 2C). After six years cover was lowest at zero  
258 P but after 20 years this treatment resulted in the highest cover: the Time × P interaction was  
259 significant (Table 2), although the main effect of P was not (Table 2). The change in  
260 percentage cover over time was also highly significant (Table 2).

261 Percentage cover of weeds and ephemerals was low throughout the experiment,  
262 neither time nor applied-P had a significant effect on percentage cover (Table 2; Fig 5AB).  
263 The cover of seeder species changed significantly over time (Table 2; Figure 5C), peaking at  
264 13 years before declining by 20 years. There was no effect of P-treatment on cover associated  
265 with seeders (Table 2).

266 Percentage cover of resprouters changed significantly over time (Table 2). Cover  
267 increased between 6 and 13 years and then declined slightly (Fig 5D). Cover was initially  
268 (after 6 years) higher at zero P than either 80 or 120 kg P ha<sup>-1</sup> and this difference increased  
269 over time: after 20 years, cover at zero P was more than twice that in the P-addition  
270 treatments (Fig. 5D). This was supported by a significant main effect of P-treatment and a  
271 significant Time × P interaction (Table 2).

272

273 *3.4. Changes in community composition with time since restoration*

274 Applied-P had a significant effect on the similarity of the restored communities to reference

275 unmined forest plots: similarity was consistently highest at zero P (General Linear Mixed

276 Model,  $P < 0.05$ ; Fig. 6). In addition, the similarity of the restored communities to the forest277 sites increased significantly over time (General Linear Mixed Model,  $P < 0.001$ ).

278

#### 279 **4. Discussion:**

280 Fertiliser application is broadly advocated as best practice for re-establishing native plant  
281 communities in post-mining restoration (e.g., EPA, 1996; Minerals Council of South Africa,  
282 2007; Tibbett, 2010; DFAT, 2016). However, we found that P-fertiliser had significant, long-  
283 term effects both above- and below-ground on restored jarrah forest. In particular, we report  
284 negative impacts of P-fertilisation on long-term soil-P concentrations, species richness and  
285 community composition. Negative impacts likely result from direct effects of elevated P (i.e.  
286 toxicity; Lambers et al., 2008; Williams et al., 2019) and facilitation of competitive plant  
287 species (e.g., legumes and overstorey trees; Daws et al., 2015). Our findings have significant  
288 implications for restoring nutrient-limited systems world-wide.

289         Following a single application of P-fertiliser at the outset of restoration, available soil  
290 P remained elevated after 20 years compared with the zero P treatment (this study), and  
291 concentrations measured in undisturbed jarrah forest (Ward, 2000; Standish et al. 2008;  
292 Tibbett et al., 2020). In Brazilian Atlantic rainforest restored after bauxite mining Bizuti et al.  
293 (2020) found that, soil P pools remained lower than unmined forest, even with fertiliser  
294 inputs. However, previous studies in jarrah forest and other restored *Eucalyptus* forests  
295 reported that a single P-fertiliser application elevated soil-P concentrations for at least 20  
296 years (Spain et al., 2018; Banning et al., 2008). These differences may result from the jarrah  
297 forest soils having soil nutrient levels that are low by global standards and, at least for the  
298 jarrah forest, suggests P application may have long-term impacts on plant responses.

299         P-fertiliser addition initially benefitted weeds and native ephemerals. This effect of P-  
300 addition has been reported previously in both restored jarrah forest and other restored  
301 vegetation communities (e.g., Prober and Wiehl 2002; Holmes, 2001; Daws et al., 2013).  
302 However, as the establishing vegetation developed, these species declined in abundance as  
303 also reported by Norman et al. (2006a). However, given their decline and the low percentage

304 cover associated with these two groups, they are unlikely to be responsible for driving the  
305 negative long-term responses to P observed for other understorey groups (e.g. resprouters).

306 Species richness and density of seeder species was generally highest in one-year old  
307 restored sites, declining thereafter. Conversely, percentage cover of this group of species  
308 reached a maximum after 13 years before declining. Many seeders are N<sub>2</sub>-fixing legumes that  
309 exhibit significant initial growth to applied-P (e.g., Daws et al., 2013, 2015) before  
310 declining over time as individuals senesce (Grant et al., 2007). Apart from year one, both the  
311 number of species and stem density of seeders was highest at zero P suggesting this group  
312 generally benefitted from a less competitive environment. Similarly, Daws et al. (2015)  
313 reported that among P-responsive legumes, while total cover responded positively to applied  
314 P, species richness was reduced at either 20 or 80 kg P ha<sup>-1</sup> compared with zero P.

315 For resprouter species, including many cluster-root forming species, richness, stem  
316 density and cover were highest at zero P throughout the 20-year duration of this experiment.  
317 One explanation for this response is that reduced competition at zero P benefitted  
318 establishment of these slow-growing species. Competition was likely to be more intense in  
319 the applied-P treatments due to increased growth of reseeders (see above) as well as  
320 potentially increased competition from the establishing overstorey layer. In newly established  
321 jarrah stands, tree growth responds positively to P-application, although these benefits of  
322 fertiliser application largely disappear by 15 years of age (Daws et al., 2019b). Furthermore,  
323 studies in a range of forested systems have shown negative effects of intense overstorey  
324 competition on understorey species richness (e.g. Chan et al., 2006). Alternatively, there may  
325 have been direct negative effects of applied-P on growth and survival in the 80 and 120 kg P  
326 ha<sup>-1</sup> treatments. Since P-toxicity has been observed for various Western Australian species  
327 (de Campos 2013; Williams et al., 2019), further experiments are required to disentangle

328 relative effects of competition and P-toxicity on the (re-)establishment of resprouter species  
329 in jarrah forest restoration.

330 Resprouter species are under-represented in restored jarrah forest compared with  
331 reference forest (Norman et al., 2006a; Koch, 2007). Our data suggest that their low  
332 abundance may partially reflect the routine application of P-fertiliser (80 kg ha<sup>-1</sup> prior to 2004  
333 and 40 kg P ha<sup>-1</sup> from 2004 onwards; Standish et al., 2015). Since many species associated  
334 with zero P plots were resprouters, this finding has significant practical implications: Alcoa  
335 propagates and plants seven resprouter species (listed exclusively as zero P species in Table  
336 3) into restored sites to increase their abundance (A.H. Grigg pers. comm.). A low P-fertiliser  
337 regime may potentially increase the success of planting these species.

338 Applied P reduced the overall similarity of the composition of restored forest to  
339 reference forest sites. Similarly, P-application rates as low as 20 kg P ha<sup>-1</sup> reduced the  
340 similarity of 2.5-year-old restored sites compared with reference forest sites (Daws et al.  
341 2013). In addition, the composition of all restored sites became more similar over time to the  
342 reference sites, although this was most pronounced at zero P. Increasing similarity was likely  
343 driven by, (1) fewer weeds and ephemerals at zero P, (2) a higher number of resprouter  
344 species at zero P, with this number increasing over time, and (3) a decrease in abundance of  
345 seeders over time in all three treatments.

346 Consistent with previous studies (e.g., Norman et al., 2006a), we found some support  
347 for the Initial Floristics Model (IFM) in plots that received 80 or 120 kg P ha<sup>-1</sup>. Specifically,  
348 the number of resprouter species was relatively constant over time for plots receiving 80 and  
349 120 kg P ha<sup>-1</sup> and overall species richness declined, reflecting species losses rather than gains.  
350 In contrast, there was no support for the IFM in the successional trajectory of unfertilised  
351 plots. Support for the model was lacking in key parameters including the increase in  
352 similarity of the restored communities to reference forest plots, and the more than doubling in

353 species richness of resprouter species at zero P over the 20-year experiment. Indeed, the data  
354 suggest a relay floristics model could be appropriate whereby early colonists such as weeds  
355 and ephemerals, and ultimately Acacias and other legumes, are replaced by resprouter  
356 species. Ongoing recruitment of resprouter species may result from extended seed dormancy  
357 in the soil seed bank, or seed dispersal into plots (or both). Rather than resprouter species  
358 failing to arrive at restored sites over time, which has been the assumption under the IFM  
359 (Koch, 2007), our data suggest elevated P may retard species recruitment and hence disrupt,  
360 succession. The mechanisms for these effects could be: (1) P-responsive competitive plants  
361 limiting establishment of these typically slow-growing species, (2) an accumulation of fine  
362 woody debris and leaf litter, which can occur at high P-application rates (Grant et al., 2007),  
363 inhibiting seedling emergence, or (3) direct toxicity from elevated P. Consequently, we  
364 propose that jarrah forest succession is contingent on P fertilisation rather than initial  
365 floristics per se.

366         One year after restoration, applied-P resulted in higher species richness of native  
367 species and a higher plant density. Combined with applied-P increasing plant cover in newly  
368 restored sites (e.g., Daws et al., 2013), this suggests an early assessment of restoration  
369 success would come to the *incorrect* conclusion that applying P was beneficial. However, at  
370 20-years, 80 or 120 kg P ha<sup>-1</sup> resulted in fewer native species, fewer individual plants, a  
371 similar level of understorey cover and lower similarity to target vegetation than zero applied-  
372 P, demonstrating the value of long-term monitoring for determining restoration success.

373

## 374 **5. Conclusion**

375 We propose that P limitation and moderation of fertiliser inputs may be important for re-  
376 establishing long-term species diversity in naturally P-impooverished forests, such as the  
377 jarrah forest. Since there is considerable overlap between areas of high plant species richness,

378 nutrient deficient soils and current and prospective mining areas (Tibbett et al., 2019b), these  
379 findings have potential broad applicability in post mining restoration.

380

381 **Acknowledgements:**

382 MID and AHG are paid employees of Alcoa. We acknowledge earlier studies by John Koch  
383 and Melanie Norman and various Alcoa employees for technical assistance.

384

385 **Funding:** This work was part funded by the Australian Research Council. The completion of  
386 this work was supported by the Building Outstanding Impact Support Programme H&F38:  
387 Restoring biodiversity to phosphorus sensitive forests.

388 **References**

- 389 Banning, N.C., Grant, C.D., Jones, D.L., & Murphy, D.V. (2008) Recovery of soil organic  
390 matter, organic matter turnover and nitrogen cycling in a post-mining forest rehabilitation  
391 chronosequence. *Soil Biology and Biochemistry*, 40, 2021–2031.
- 392 Bates, D., Maechler, M., & Bolker, B. (2011) Lme4: linear mixed-effects models using S4  
393 classes. R package version 0.999375-42. <http://CRAN.Rproject.org/package=lme4>.  
394 Accessed March 2013.
- 395 Bell, D.T. (2001) Ecological response syndromes in the flora of southwestern Western  
396 Australia: fire resprouters versus reseeder. *Botanical Review*, 67, 417–440.
- 397 Bell, I.C. (2001) Establishment of native ecosystems after mining - Australian experience  
398 across diverse biogeographic zones. *Ecological Engineering*, 17, 179–186.
- 399 Bell, D.T., Plummer, J.A., & Taylor, S.K. (1993) Seed germination ecology in southwestern  
400 Western Australia. *Botanical Review*, 59, 24–73.
- 401 Bellairs, S.M., & Bell, D.T. (1990) Canopy-borne seed store in three Western Australian  
402 plant communities. *Australian Journal of Ecology*, 15, 299–305.
- 403 Bizuti, D.T.G., Soares, T.M., Duarte, M.M., Casagrande, J.C., Moreno, V.S., Peinado F.J.M.,  
404 de Medeiros, S.D.S., van Melis, J., Schweizer D., & Brancalion, P.H.S. (2020) Recovery  
405 of soil phosphorus on former bauxite mines through tropical forest restoration. *Restoration*  
406 *Ecology* 28, 1237-1246.
- 407 Bowen, B.J. (1991) Fire response within the family Proteaceae: A comparison of plants  
408 displaying the seeder and resprouter mode of recovery. PhD thesis, University of Western  
409 Australia, Perth, WA, Australia.
- 410 Bowen, B.J., & Pate, J.S. (1993) The significance of root starch in post-fire shoot recovery of  
411 the resprouter *Stirlingia latifolia* R. Br. (Proteaceae). *Annals of Botany*, 72, 7–16.

- 412 Brancalion, P.H.S., Campoe, O., Mendes, J.C.T., Noel, C., Moreira, G.G., van Melis, J.,  
413 Stape, J.L. & Guillemot, J. (2019) Intensive silviculture enhances biomass accumulation  
414 and tree diversity recovery in tropical forest restoration. *Ecological Applications* 29,  
415 e01847.
- 416 Burrows, N.D., Wardell-Johnson, G., & Ward, B. (2008) Post-fire juvenile period of plants in  
417 south-west Australia forests and implications for fire management. *Journal of the Royal*  
418 *Society of Western Australia*, 91, 163–174, Fire response database available at  
419 <http://naturemap.dec.wa.gov.au>
- 420 Chan, S.S., Larson, D.J., Maas-Hebner, K.G., Emmingham, W.H., Johnston, S.R. &  
421 Mikowski, D.A. (2006) Overstory and understory development in thinned and  
422 underplanted Oregon Coast Range Douglas-fir stands. *Canadian Journal of Forest*  
423 *Research* 36, 2696-2711.
- 424 Clarke, K.R., & Warwick, R.M. (1994) Change in marine communities: an approach to  
425 statistical analysis and interpretation. Natural Environment Research Council, Swindon,  
426 UK.
- 427 Clarke, P.J., Lawes, M.J., Murphy, B.P., Russell-Smith, J., Nano, C.E.M., Bradstock, R.,  
428 Enright, N.J., Fontaine, J.B., Gosper, C.R., Radford, I., Midgley, J.J., Gunton, R.M. (2015)  
429 A synthesis of postfire recovery traits of woody plants in Australian ecosystems. *Science*  
430 *of the Total Environment* 534, 31-42.
- 431 Daws, M.I., Standish, R.J., Koch, J.M., & Morald, T.K. (2013) Nitrogen and phosphorus  
432 fertiliser regime affect jarrah forest restoration after bauxite mining in Western Australia.  
433 *Applied Vegetation Science*, 16, 610–618.
- 434 Daws, M.I., Standish, R.J., Koch, J.M., Morald, T.K., Tibbett, M., & Hobbs, R.J. (2015)  
435 Phosphorus fertilisation and large legume species affect jarrah forest restoration after  
436 bauxite mining. *Forest Ecology and Management*, 354, 10–17.

- 437 Daws, M.I., Grigg, A.H., Standish, R.J., & Tibbett, M. (2019a) Applied phosphorus has long-  
438 term impacts on vegetation responses in restored jarrah forest, In: A.B. Fourie and M.  
439 Tibbett (eds.) Proceedings of the 13th International Conference on Mine Closure,  
440 Australian Centre for Geomechanics, Perth, pp. 693-704.
- 441 Daws, M.I., Grigg, A.H., Tibbett, M., & Standish, R.J. (2019b) Enduring effects of large  
442 legumes and phosphorus fertiliser on jarrah forest restoration 15 years after bauxite  
443 mining. *Forest Ecology and Management*, 438, 204–214.
- 444 de Campos, M.C.R., Pearse, S.J., Oliveira, R.S. & Lambers, H. (2013) Downregulation of net  
445 phosphorus-uptake capacity is inversely related to leaf phosphorus-resorption proficiency  
446 464 in four species from a phosphorus-impooverished environment. *Annals of Botany* 111,  
447 445–454.
- 448 DFAT (2016) Mine Rehabilitation: Leading Practice Sustainable Development Program for  
449 the Mining Industry. Department of Foreign Affairs and Trade, Commonwealth  
450 Government of Australia, Canberra, Australia.
- 451 Egler, F.E. (1954) Vegetation science concepts. I. Initial floristic composition, a factor in  
452 oldfield vegetation development. *Vegetatio*, 4, 412–417.
- 453 (EPA) Environment Protection Agency. 1995. Rehabilitation and revegetation. Best Practice  
454 Environmental Management in Mining, Commonwealth of Australia, Canberra, Australia
- 455 Grant, C.D., Ward, S.C., & Morley, S.C. (2007) Return of ecosystem function to restored  
456 bauxite mines in Western Australia. *Restoration Ecology*, 15, S94–S103.
- 457 Grigg A.H., & Grant C.D. (2009) Overstorey growth response to thinning, burning and  
458 fertiliser in 10–13-year-old rehabilitated jarrah (*Eucalyptus marginata*) forest after bauxite  
459 mining in south-western Australia. *Australian Forestry* 72, 80–86.

- 460 Holmes, P.M. (2001) Shrubland restoration following woody alien invasion and mining:  
461 effects of topsoil depth, seed source, and fertilizer addition. *Restoration Ecology*, 9, 71–  
462 84.
- 463 Hopper, S.D. (2009) OCBIL theory: towards an integrated understanding of the evolution,  
464 ecology and conservation of biodiversity on old, climatically buffered, infertile  
465 landscapes. *Plant and Soil* 322, 49-86.
- 466 Hopper, S.D., & Gioia, P. (2004) The Southwest Australian Floristic region: evolution and  
467 conservation of a global hot spot of biodiversity. *Annual Review of Ecology, Evolution*  
468 *and Systematics* 35, 623-650.
- 469 Isbell, F., Reich, P.B., Tilman, D., Hobbie, S.E., Polasky, S., & Binder, S. (2013). Nutrient  
470 enrichment, biodiversity loss, and consequent declines in ecosystem productivity.  
471 *Proceedings of the National Academy of Sciences*, 110, 11911-11916.
- 472 Koch, J.M. (2007) Restoring a Jarrah forest understorey vegetation after bauxite mining in  
473 Western Australia. *Restoration Ecology*, 15, S26–S39.
- 474 Lambers, H., Raven, J.A., Shaver, G.R. & Smith, S.E. (2008) Plant nutrient-acquisition  
475 strategies change with soil age. *Trends in Ecology and Evolution*, 23, 95–103.
- 476 Lockley, I.R., & Koch, J.M. (1996) Response of two eucalypt species to fertilizer application  
477 on rehabilitated bauxite mines in Western Australia. Alcoa of Australia Ltd., Research  
478 Bulletin No.27.
- 479 Minerals Council of South Africa, (2007) Guidelines for the rehabilitation of mine land.  
480 Chamber of Mines of South Africa / Coaltech.
- 481 Norman, M.A., Koch, J.M., Grant, C.D., Morald, T.K., & Ward, S.C. (2006a) Vegetation  
482 succession after bauxite mining in Western Australia. *Restoration Ecology*, 14, 278–288.

- 483 Norman, M.A., Plummer, J.A., Koch, J.M., & Mullins, G.R. (2006b) Optimising smoke  
484 treatments for jarrah (*Eucalyptus marginata*) forest rehabilitation. Australian Journal of  
485 Botany, 54, 571–581.
- 486 Pate, J.S., Froend, R.H., Bowen, B.J., Hansen, A., & Kuo, J. (1990) Seedling growth and  
487 storage characteristics of seeder and resprouter species of Mediterranean-type ecosystems  
488 of S.W. Australia. Annals of Botany, 65, 585–601.
- 489 Prematuri, R., Turjaman, M., Sato, T., & Tawaraya, K. (2020) Post bauxite mining land soil  
490 characteristics and its effects on the growth of *Falcataria moluccana* (Miq.) Barneby & J.  
491 W. Grimes and *Albizia saman* (Jacq.) Merr. Applied and Environmental Soil Science  
492 article ID 6764380.
- 493 Prober, S.M., & Wiehl, G. (2012) Relationships among soil fertility, native plant diversity  
494 and exotic plant abundance inform restoration of forb-rich eucalypt woodlands. Diversity  
495 and Distributions, 18, 795–807.
- 496 R Development Core Team (2009) R 2.9.0. The R Foundation for Statistical Computing,  
497 Vienna.
- 498 Rayment, G.E., & Higginson, F.R. (1992) Australian Laboratory Handbook of Soil and  
499 Water Chemical Methods. Inkata Press, Melbourne.
- 500 Smith, M.A., Grant, C.D., Loneragan, W.A., & Koch, J.M. (2004) Fire management  
501 implications of fuel loads and vegetation structure in jarrah forest restoration on bauxite  
502 mines in Western Australia. Forest Ecology and Management, 187, 247–266.
- 503 Smith, M.A., Loneragan, W.A., Grant, C.D., & Koch, J.M. (2000) Effect of fire on the topsoil  
504 seed banks of rehabilitated bauxite mine sites in the jarrah forest of Western Australia.  
505 Ecological Management and Restoration, 1, 50–58.
- 506 Spain, A.V., Tibbett, M., Ridd, M., & McLaren, T.I. (2018) Phosphorus dynamics in a  
507 tropical forest soil restored after strip mining. Plant and Soil, 427, 105–123.

- 508 Standish, R.J., Cramer, V.A., Hobbs, R.J., & Kobryn, H.T. (2006) Legacy of land-use evident  
509 in soils of Western Australia's wheatbelt. *Plant and Soil*, 280, 189–207.
- 510 Standish, R.J., Morald, T.K., Koch, J.M., Hobbs, R.J., & Tibbett, M. (2008) Restoring jarrah  
511 forest after bauxite mining in Western Australia: the effect of fertiliser on floristic  
512 diversity and composition. In: Fourie, A.B., Tibbett, M., Weiersbye, I.M., Dye, P.J. (Eds.),  
513 Mine Closure 2008. Australian Centre for Geomechanics, Perth, Western Australia, pp.  
514 717–725.
- 515 Standish, R.J., Daws, M.I., Gove, A.D., Didham, R.K., Grigg, A.H., Koch, J.M., & Hobbs,  
516 R.J. (2015) Long-term data suggest jarrah-forest establishment at restored mine sites is  
517 resistant to climate variability. *Journal of Ecology*, 103, 78–89.
- 518 Stock, W.D., & Allsopp, N. (1992) Functional perspective of ecosystems. Pages 241–259 in  
519 R. M. Cowling, editor. *The ecology of fynbos: nutrients, fire and diversity*. Oxford  
520 University Press, Cape Town.
- 521 Tibbett, M. (2010) Large-scale Mine Site Restoration of Australian Eucalypt Forests After  
522 Bauxite Mining: Soil Management and Ecosystem Development. In: *Ecology of Industrial  
523 Pollution*, L.C. Batty & K. Hallberg, (Eds), pp. 309-326. Cambridge University Press, UK.
- 524 Tibbett, M., Gil-Martínez, M., Fraser, T., Green, I.D., Duddigan, S., De Oliveira V., ... Diaz,  
525 A. (2019a) Long-term acidification of pH neutral grasslands affects soil biodiversity,  
526 fertility and function in a heathland restoration. *Catena*, 180, 401–415.
- 527 Tibbett, M., O'Connor, R & Daws, M.I. (2019b) Too much of a good thing: phosphorus over-  
528 fertilisation in rehabilitated landscapes of high biodiversity value. In: A.B. Fourie and M.  
529 Tibbett (eds.) *Proceedings of the 13th International Conference on Mine Closure*,  
530 Australian Centre for Geomechanics, Perth, pp. 651-665.

- 531 Tibbett, M., Daws, M.I., George, S.J., & Ryan, M.H. (2020) The where, when and what of  
532 phosphorus fertilisation for seedling establishment in a biodiverse jarrah forest restoration  
533 after bauxite mining in Western Australia. *Ecological Engineering*, 153, 105907.
- 534 Ward, S.C. (2000) Soil development on rehabilitated bauxite mines in south-west Australia.  
535 *Australian Journal of Soil Research*, 38, 453–464.
- 536 Ward, S.C., Koch, J.M., & Grant, C.D. (1997) Ecological aspects of soil seed-banks in  
537 relation to bauxite mining. I. Unmined jarrah forest. *Australian Journal of Ecology*, 22,  
538 169–176.
- 539 Ward, S.C., Koch, J.M., & Nichols, O.G. (1990) Bauxite mine rehabilitation in the Darling  
540 Range, Western Australia. *Proceedings of the Ecology Society of Australia*, 16, 557–565.
- 541 Wassen, M.J., Venterink, H.O., Lapshina, E.D., & Tanneberger, F. (2005) Endangered plants  
542 persist under phosphorus limitation. *Nature*, 437, 547–550.
- 543 Western Australian Herbarium (2012) FloraBase – the Western Australian Flora. Department  
544 of Environment and Conservation. <<http://florabase.dec.wa.gov.au/>>.
- 545 Wheeler, B.D., & Shaw, S.C. (1991) Aboveground crop mass and species richness of the  
546 principal types of herbaceous rich-fen vegetation of lowland England and Wales. *Journal*  
547 *of Ecology*, 79, 285–301.
- 548 Williams, A., George, S., Birt, H.W.G., Daws, M.I., & Tibbett, M. (2019) Sensitivity of  
549 seedling growth to phosphorus supply in six tree species of the Australian Great Western  
550 Woodlands. *Australian Journal of Botany*, 67, 390-396.
- 551
- 552

553

554 Table 1: Mean ( $\pm 1$ SE) soil  $\text{NH}_4^+$ ,  $\text{NO}_3^-$  Colwell(available)-P and soil pH, at 0-10 cm depth,

555 across the phosphorus fertiliser treatments in 20-year-old restored jarrah forest.

556

Fertiliser treatment	Application rate (kg ha <sup>-1</sup> )	Colwell P (mg kg <sup>-1</sup> )	$\text{NH}_4^+$ (mg kg <sup>-1</sup> )	$\text{NO}_3^-$ (mg kg <sup>-1</sup> )	Soil pH
	0	1.22 $\pm$ 0.11 <sup>a</sup>	4.73 $\pm$ 0.56 <sup>a</sup>	< 1	4.97 $\pm$ 0.06 <sup>a</sup>
Phosphorus	80	4.58 $\pm$ 0.51 <sup>b</sup>	4.72 $\pm$ 0.49 <sup>a</sup>	< 1	4.70 $\pm$ 0.07 <sup>b</sup>
	120	6.98 $\pm$ 0.81 <sup>c</sup>	4.77 $\pm$ 0.36 <sup>a</sup>	< 1	4.63 $\pm$ 0.06 <sup>b</sup>

557 Superscripts indicate a significant difference (One-way ANOVA,  $P < 0.05$ ) between values

558 within each column.

559

560

561 **Table 2:** Summary of results from the general linear model analyses of the effects of P-  
 562 application rate and time since initiation of restoration on responses of the species groupings.

563

<b>Species richness</b>					
Factors	All native species	Weeds	Ephemerals	Seeder species	Resprouter species
P-application rate	*	*	n.sig.	*	*
Time	***	***	***	***	***
P × time	***	*	*	n.sig.	*
<b>Stem density</b>					
P-application rate	n.sig.	n.sig.	n.sig.	*	*
Time	**	n.sig.	**	***	***
P × time	*	n.sig.	*	n.sig.	*
<b>Plant cover</b>					
P-application rate	n.sig.	n.sig.	n.sig.	n.sig.	*
Time	***	n.sig.	n.sig.	***	***
P × time	**	n.sig.	n.sig.	n.sig.	*

564 n.sig.  $P > 0.05$ ; \* $P < 0.05$ ; \*\* $P < 0.01$ ; \*\*\* $P < 0.001$

565 **Table 3:** Species associated with unfertilised and fertilised plots in restored sites at 20 years.

566 Species denoted with an asterisk are exotic, those in bold are resprouters.

	0P plots		80P and 120P plots
Anthericaceae	<i>Thysanotus fastigiatus</i>	Asteraceae	<i>Hypochaeris glabra*</i>
Apiaceae	<b><i>Pentapeltis peltigera</i></b>	Colchicaceae	<b><i>Burchardia congesta</i></b>
	<i>Xanthosia candida</i>	Euphorbiaceae	<i>Phyllanthus calycinus</i>
Campanulaceae	<i>Wahlenbergia preisii</i>	Fabaceae	<i>Acacia extensa</i>
Cyperaceae	<b><i>Tetraria capillaris</i></b> †		<i>Paraserianthes lophantha</i>
Dasyogonaceae	<b><i>Lomandra caespitosa</i></b> †	Orchidaceae	<b><i>Caladenia latifolia</i></b>
	<b><i>L. hermaphrodita</i></b> †	Pittosporaceae	<i>Billardiera heterophylla</i>
	<b><i>L. sonderi</i></b> †	Poaceae	<i>Aira caryophyllea*</i>
Epacridaceae	<i>Andersonia lehmanniana</i>		<i>Pentaschistis airoides*</i>
Dilleniaceae	<b><i>Hibbertia acerosa</i></b>		
	<b><i>H. amplexicaulis</i></b> †		
Fabaceae	<i>Acacia drummondii</i>		
Goodeniaceae †	<b><i>Lechenaultia biloba</i></b>		
	<b><i>Scaevola calliptera</i></b> †		
Haemodoraceae	<b><i>Conostylis setosa</i></b>		
Orchidaceae	<b><i>Microtis media</i></b>		
Proteaceae	<b><i>Hakea undulata</i></b>		
Rutaceae	<b><i>Boronia fastigiatus</i></b>		
Tremandraceae	<b><i>Tetratheca hirsuta</i></b>		

567 †Currently propagated by Alcoa and planted into newly restored sites.

568 **Figure legends:**

569 Figure 1. Photograph of one of the 20-year-old experimental plots.

570 Figure 2. Effects of P-fertiliser application rate on: (A) species richness, (B) stem density,  
571 and (C) total native species cover, over time. Error bars  $\pm 1$ SE of the mean.

572 Figure 3. Effects of P-fertiliser application rate on the species richness of, (A) weeds, (B)  
573 ephemerals, (C) seeders, and (D) resprouters, over time. Error bars  $\pm 1$ SE of the mean.

574 Figure 4. Effects of P-fertiliser application rate on density of, (A) weeds, (B) ephemerals, (C)  
575 seeders, and (D) resprouters, over time. Error bars  $\pm 1$ SE of the mean although, for  
576 clarity, error bars for the weeds are  $+1$ SE of the mean. High weed density at  $120 \text{ kg P ha}^{-1}$   
577 at 13 years was driven by an extremely high density of *Aira caryophyllea* in a single  
578 plot.

579 Figure 5. Effects of P-fertiliser application rate on the percentage cover of, (A) weeds, (B)  
580 ephemerals, (C) seeders, and (D) resprouters, over time. Error bars  $\pm 1$ SE of the mean.  
581 High weed cover at  $120 \text{ kg P ha}^{-1}$  at 13 years was driven by an extremely high abundance  
582 of *Aira caryophyllea* in a single plot.

583 Figure 6. Effects of P-fertiliser application rate on the similarity of species composition  
584 between the fertiliser treatments and adjacent reference forest plots over time. Similarity  
585 values for the treatment plots are expressed as a percentage of the average value of forest  
586 plot: forest plot similarity. Error bars  $\pm 1$ SE of the mean.

587

588

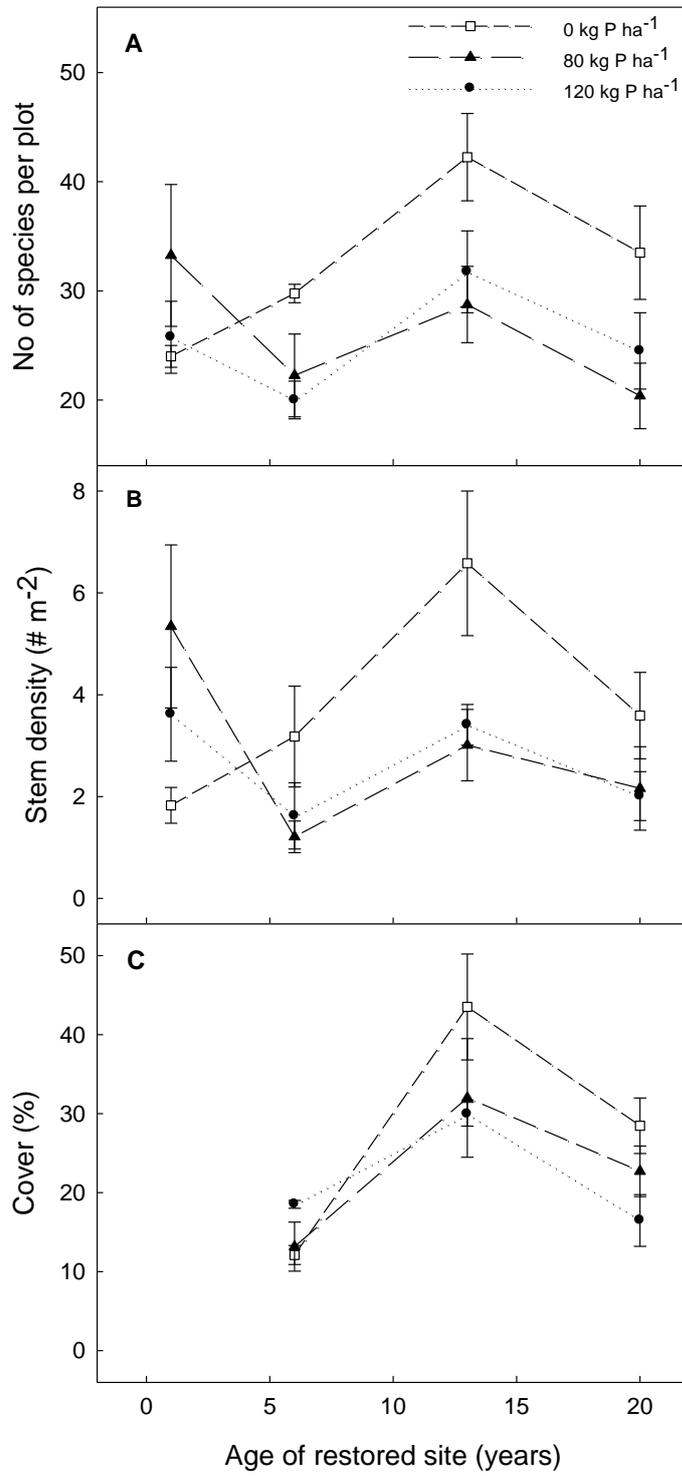
589 Figure 1:



590

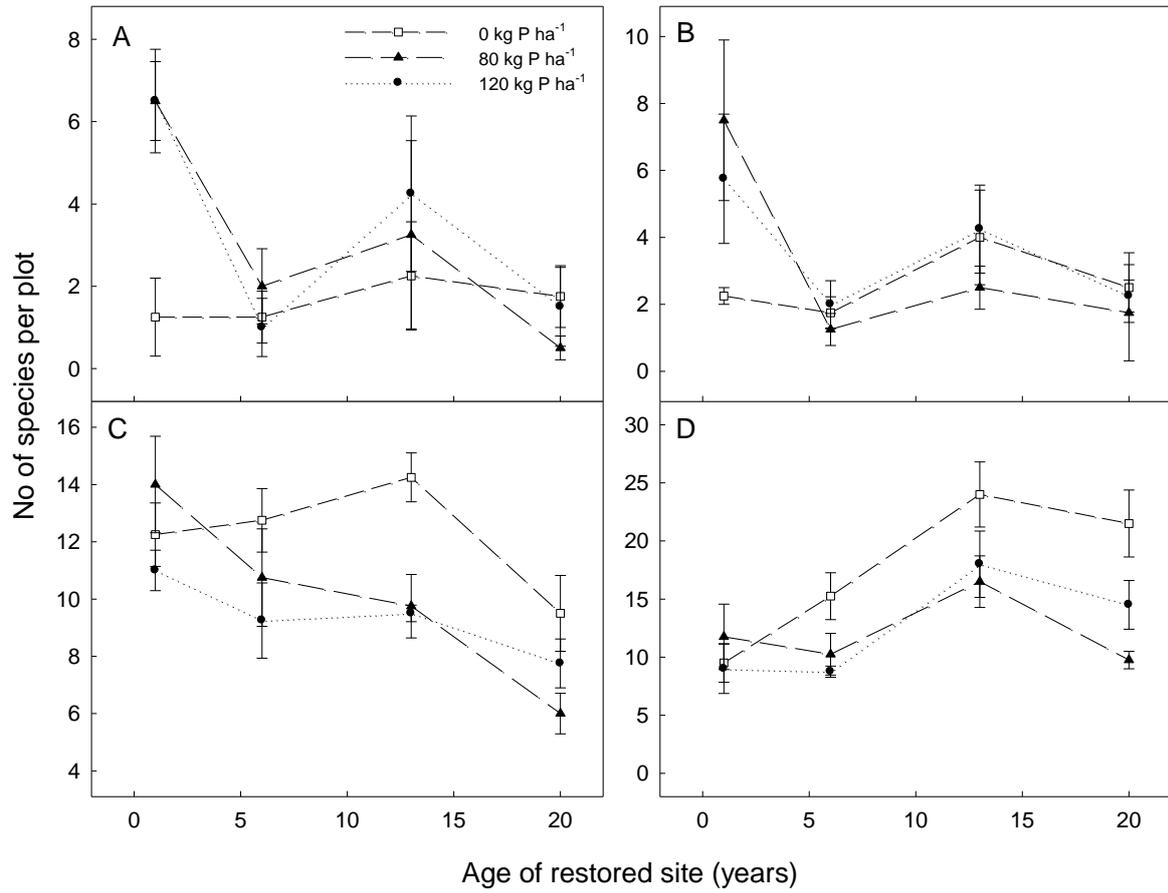
591 Figure 2:

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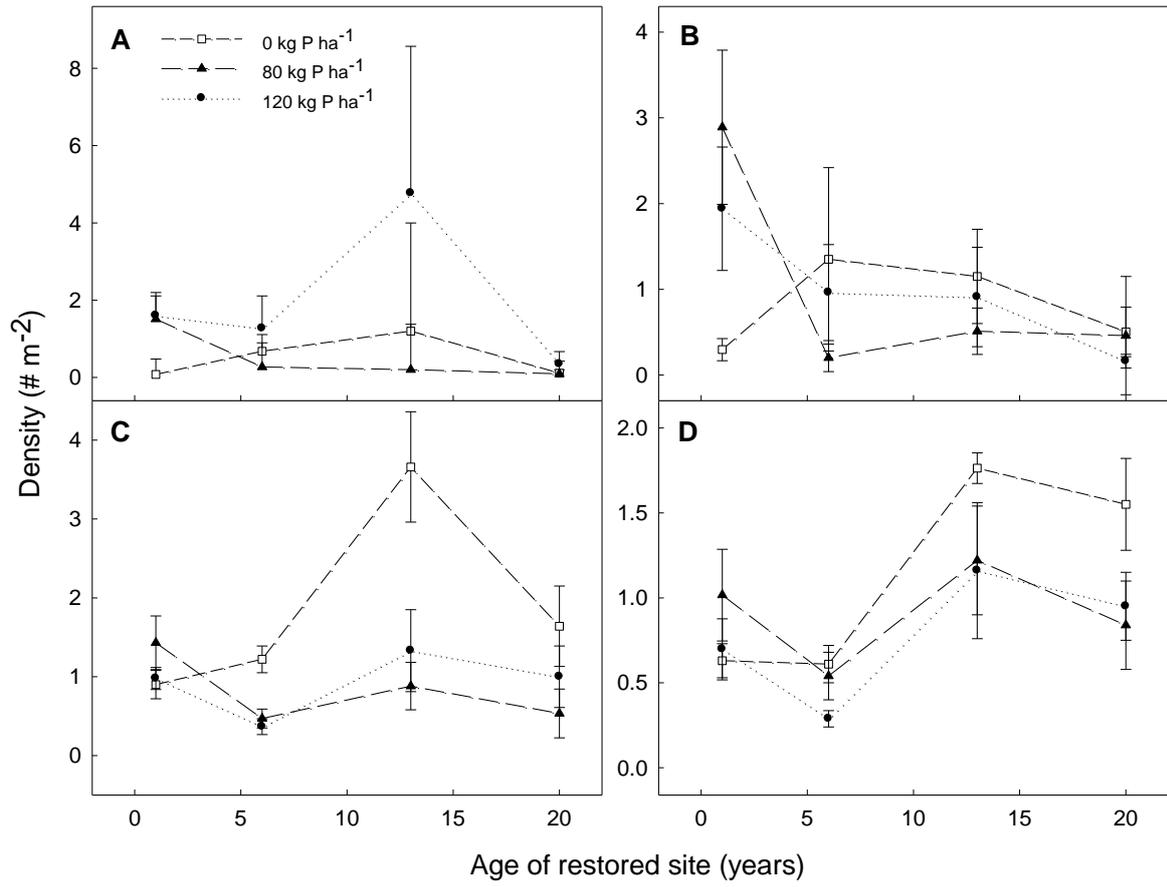
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594 Figure 3:



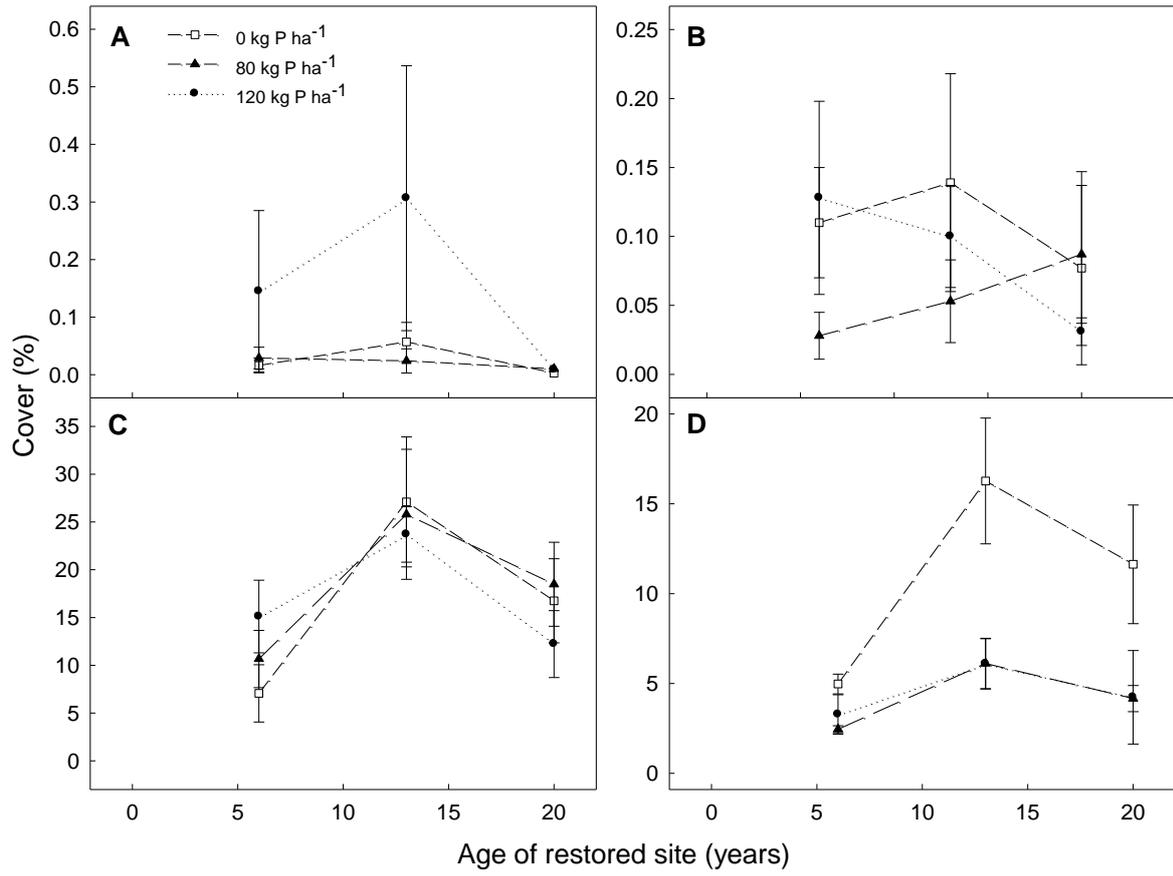
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596 Figure 4:



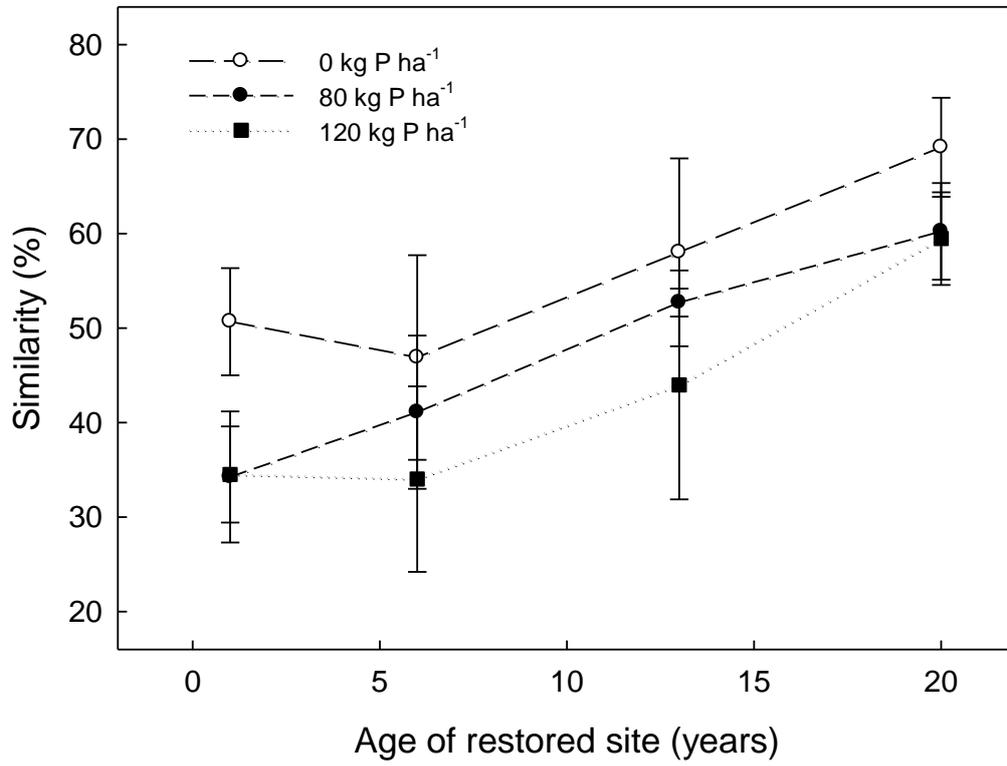
597

598 Figure 5:



599

600 Figure 6:



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