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1 **Negative impacts of felling in exotic spruce plantations on moth diversity mitigated by**
2 **remnants of deciduous tree cover**

3 Lucinda Kirkpatrick^{1,2}, Sallie Bailey³, Kirsty J. Park¹

4 Lucinda Kirkpatrick (Corresponding author)

5 ¹Biological and Environmental Sciences
6 University of Stirling,
7 Stirling, Scotland
8 FK9 4LA.

9
10 ²EVECO
11 Universiteit Antwerpen
12 Universiteitsplein 1
13 Wilrijk
14 2610

15
16 ³Forestry Commission Scotland,
17 Edinburgh,
18 United Kingdom
19

20 Email: Lucinda.Kirkpatrick@uantwerpen.be
21 Tel: +32 0495 477620

22

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26 **Abstract:**

27 Moths are a vital ecosystem component and are currently undergoing extensive and severe declines
28 across multiple species, partly attributed to habitat alteration. Although most remaining forest cover
29 in Europe consists of intensively managed plantation woodlands, no studies have examined the
30 influence of management practices on moth communities within plantations. Here, we aimed to
31 determine: (1) how species richness, abundance, diversity of macro and micro moths in commercial
32 conifer plantations respond to management at multiple spatial scales; (2) what the impacts of forest
33 management practices on moth diversity are, and (3) how priority Biodiversity Action Plan (BAP)

34 species respond to management. BAP species were selected as they represent formerly widespread
35 and common species, which have undergone substantial declines in the UK and Europe. We assessed
36 moth communities in three conifer plantations in Northern England and Scotland by light trapping,
37 combining local (e.g. age of planting) and landscape level (e.g. proximity to felled areas)
38 characteristics to evaluate the impacts of forest management on moths. We found no relationship
39 between local factors and moth richness, abundance and diversity but the amount of clear felling in
40 the surrounding landscape had a strongly negative correlation. In contrast, the amount and
41 proximity of broadleaf cover in the surrounding landscape positively influenced macro moth richness
42 and abundance. For six BAP species, abundances were lower close to felled areas but increased with
43 the size of adjacent broadleaf patches. We conclude that clear felling negatively affects moths,
44 probably through alteration of habitats, the loss of larval host plants, and by limiting dispersal. A
45 shift to continuous cover and maintaining broadleaf tree cover within plantations will greatly
46 enhance their value for moth communities.

47 **Keywords: Moth; Lepidoptera; abundance; species richness; plantation management; landscape**
48 **heterogeneity**

49

50 **1. Introduction**

51 Maintaining and restoring biodiversity is a key tenet in sustainable ecosystem management, the
52 paradigm currently guiding habitat management practices across Europe and North America (Ober &
53 Hayes 2010). This is driven by concern about world-wide declines in species and populations across a
54 range of taxa (Dirzo et al. 2014) and recognition that much of this is driven by habitat loss and
55 fragmentation, caused by anthropogenic change (Thomas 2004). In many countries the timber
56 industry has responded to recognition of the importance of biodiversity by shifting focus from purely
57 timber production to one which encourages sustainable practices that promote both wildlife
58 conservation and sustainable timber yields (Macdonald et al. 2009). In Europe this has been driven

59 by policy change initiated as a result of the Convention of Biological Diversity, requiring explicit
60 consideration of environmental, economic and social objectives and a multi-purpose approach to
61 forestry (Watts et al. 2008). However, efforts to assess the impact of forest practices can be
62 challenging as there is often inadequate knowledge of the current distribution and abundance of
63 many taxa in managed forest systems (Ober & Hayes 2010).

64 Plantation forests are generally considered poor for biodiversity as they are primarily
65 composed of non-native tree species, often in monocultures, which are under an intensive
66 management regime (Brockhoff et al. 2008). However, they usually constitute the largest patches
67 of tree cover in many European countries and as such may be valuable for preserving biodiversity if
68 managed sympathetically. One of the few studies carried out at a national scale demonstrated that
69 plantations can support diverse invertebrate communities in the UK, and that invertebrate
70 community composition and abundance is most affected by tree species planted and geographic
71 location (Humphrey et al. 2003). The structure of the plantation was also important for some
72 groups: ground dwelling Carabid diversity decreased with canopy cover whereas overall Coleopteran
73 richness and abundance in the canopy increased (Humphrey et al. 2003). The effect of stand age on
74 invertebrate communities can also vary between taxa. Higher abundance and diversity of Coleoptera
75 has been associated with older *Larix kaempferi* (Larch) and *Picea sitchensis* (Sitka Spruce) plantations
76 in Japan and Northern Ireland due to increased heterogeneity and regeneration of native trees
77 (Ohsawa 2005; Oxbrough et al. 2010). However, the high canopy cover in mature plantations can
78 negatively affect other groups associated with open habitats (e.g. Arachnid diversity; Oxbrough et al.
79 2010).

80 Despite being a speciose taxonomic group and an important component of the invertebrate
81 community, the impacts of plantation forestry on night active Lepidoptera are yet to be explored.
82 Substantial declines of many moth species have occurred in the last few decades; two thirds of
83 common and widespread species in the UK have suffered rapid population decreases (Conrad et al.

84 2006) with similar patterns occurring in Finland (Mattila et al. 2006) and Sweden (Franzén &
85 Johannesson 2007). Rapid economic development, urbanisation, changes to silvicultural
86 management and agricultural expansion have all been implicated in causing these declines (Conrad
87 et al. 2006; Fox et al. 2013). Taken together, these studies provide overwhelming evidence that
88 moths are facing declines on a large geographic scale, across a range of habitats, which mirrors
89 similar effects found in less species rich groups such as butterflies and bumblebees (Warren et al.
90 2001; Goulson et al. 2008). Such losses are likely to have substantial effects at both higher and lower
91 trophic levels. Moths are a key component of terrestrial ecosystems, providing ecosystem services
92 through modification of ecosystem functioning by saproxylic species (Merckx et al. 2012), impacting
93 upon plant growth through larval feeding activity, acting as pollinators and providing food for a
94 range of taxa such as birds, small mammals and bats (Fox et al. 2013).

95 Intensified silvicultural practices have been suggested as one major driver of the decline in
96 moth diversity and abundance (Fox et al. 2013). However, most studies have only focussed on the
97 negative effects that a reduction in traditional deciduous forest management practices has had on
98 lepidopteran species, and have not considered the role that non-native plantations may play.
99 Reductions in deciduous forest management techniques such as coppicing and opening up rides
100 have resulted in lower moth diversity by increasing structural complexity and changing botanical
101 communities (Fox et al. 2013; Merckx et al. 2012; Warren & Bourn 2011). In general, moths
102 associated with deciduous trees have declined throughout Europe, with larval host plant specificity a
103 key factor in extinction likelihood in parts of Scandinavia (Mattila et al. 2006; Franzén & Johannesson
104 2007), whilst species associated with conifer trees have increased (Fox et al. 2013). Our current
105 knowledge of moths in non-native coniferous plantations comes largely from studies which have
106 focused on the management of pest species, and to the best of our knowledge no research has
107 explicitly explored moth community composition and the impacts of forest management in exotic
108 plantations.

109 Whilst little is known about the impacts of timber harvesting on Lepidoptera in non-native
110 plantations, studies in native hardwood forests have suggested that effects are largely negative. In
111 Indiana and Ohio, Summerville and Crist (2002, 2014) demonstrated that clear felling in native
112 hardwood forests disrupted moth communities beyond the stand being felled, limiting the diversity
113 of species able to persist within the landscape. Impacts of timber harvest on Lepidoptera can persist
114 for up to 60 years (Summerville et al. 2009), although Summerville (2013) suggests that less
115 intensive practices such as shelterwood harvest (removal of 15% standing wood) may support a
116 higher richness and abundance of moth communities. In native conifer forests in Oregon, moth
117 dominance and diversity was associated with greater canopy cover whereas richness was only
118 affected by elevation, with higher species richness at lower elevations (Ober & Hayes 2010). These
119 studies from North America demonstrate that managed native forest systems can support diverse
120 lepidopteran communities, but the extent to which this is true in managed non-native plantations
121 has not yet been examined. Specifically, in this study we aim to assess the impact of the following on
122 moth abundance, richness, diversity and dominance in conifer plantations:

- 123 1. Influential, local scale plantation characteristics (e.g. age of planting, ground cover);
- 124 2. Proximity and prevalence of clear felling in the surrounding landscape;
- 125 3. Proximity and prevalence and of broadleaf tree cover within the surrounding landscape.

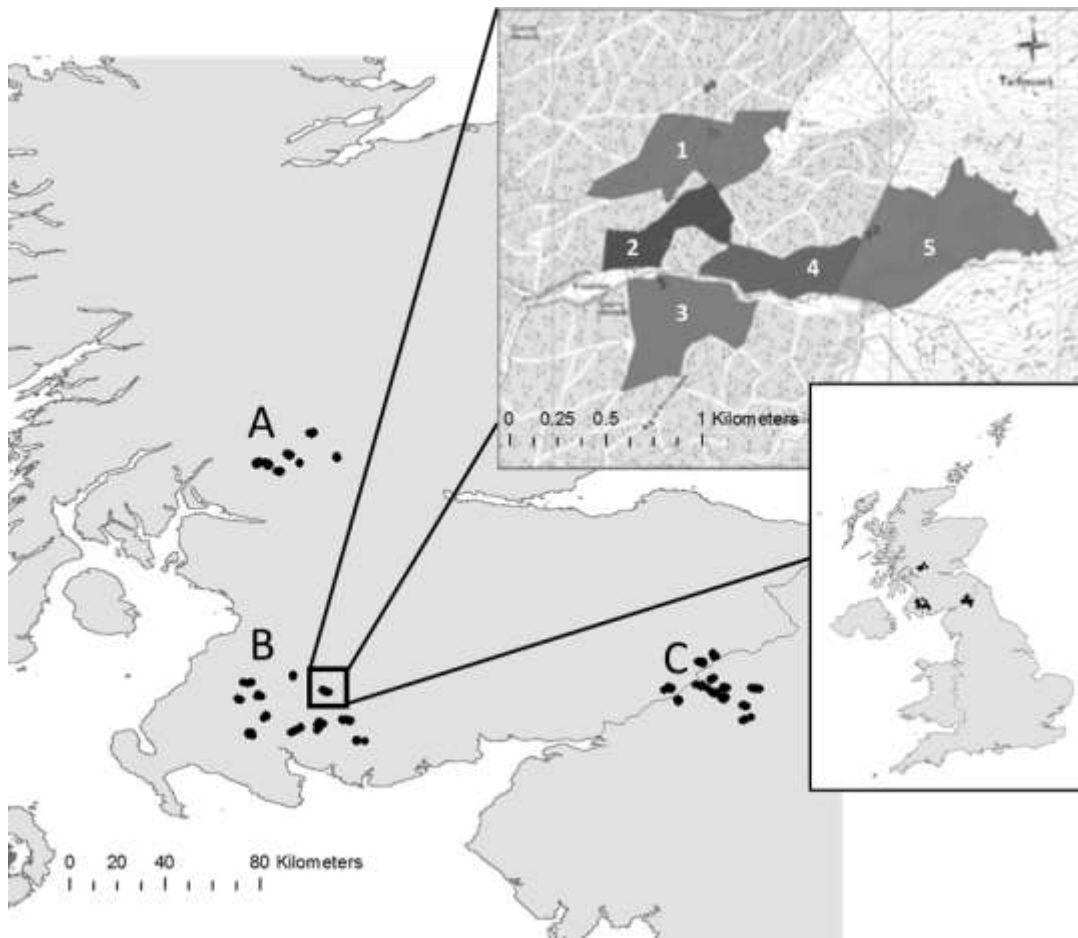
126 Since declining moth species might respond differently to the wider moth community, we examined
127 the impacts of the above characteristics for moth communities as a whole, and separately for
128 priority biodiversity action plan (BAP) species. These are formerly widespread and common species
129 which have undergone population declines of between 70 – 90% in the last few decades, and as such
130 are of particular scientific interest (Fox et al. 2013).

131 **2. Methods**

132 The study was conducted in three plantation forests in Central and Southern Scotland and Northern
133 England (Figure 1). Widespread deforestation had already occurred in this area by the Holocene;

134 prior to the planting of the plantations in 1920 – 1940, the sampling areas would have consisted of
135 open, upland moorland predominantly used for sheep grazing, with small patches of remaining
136 broadleaf. The three forests were chosen for their large size (ranging from 30,000 ha in Cowal and
137 Trossachs to 60,000 ha in Kielder and 114,000 ha in Galloway), high productivity and the
138 predominance of *Picea sitchensis*, the most commonly planted and intensively managed coniferous
139 tree species in the UK, and a common plantation tree species in Europe (Boye & Dietz 2005). Within
140 each forest, multiple sites, a minimum of 4 km from each other, were selected using a Forestry
141 Commission sub-compartment database within a Geographic Information System (GIS) (ArcMap
142 10.1, ESRI) based on stand (a unit of plantation management) age and species composition (Figure
143 1).

144 In total, seven sites were surveyed in Cowal and Trossachs, 12 in Galloway Forest and 12 in
145 Kielder Forest. Where possible a stand of trees at each management stage was selected in each site,
146 which was a maximum of 2km² in size. Not all sites had all stands of each management age resulting
147 in an unbalanced design of between four and six stands per site and a total of 285 stands across 31
148 sites. See supplementary data (4) for a description of the different stand types.



149

150 Figure 1. Location of field sites at three different study areas in (A) Cowal and Trossachs, South West
 151 Scotland, (B) Galloway, South West Scotland and (C) Kielder, Northern England. Stand types were as
 152 follows: Clearfell (1), Young (2), Thicket (3), Thin (4), Mature (5). See Supplementary data 4 for stand
 153 details.

154 *2.1 Invertebrate trapping*

155 Each site was surveyed for one night. Moths were trapped using portable 6W heath light traps using
 156 E7586 9" actinic tube lights, powered with 12V batteries which were activated 15 mins after sunset
 157 and switched off after 4 hours (approximating the duration of the shortest night in the study area).
 158 This ensured that species flying at dusk and during the night were surveyed regardless of night
 159 duration. Species flying at dawn would most likely be missed as traps were often turned off before
 160 dawn. Surveys were only conducted on nights that were above 8°C in temperature and wind speed
 161 of less than Beaufort 4, and were randomised as far as possible during the survey season between

162 the different geographical areas. We recognise that surveying each site only once provides a coarse
163 estimate of local moth assemblages; however, we are primarily interested in comparisons between
164 stand types to identify potentially influential characteristics, which requires a large sample size. This
165 same approach has been used to identify the influence of woodland characteristics on species
166 richness, diversity and abundance of moth populations in both agricultural and urban landscapes
167 (Fuentes-Montemayor et al. 2012; Lintott et al. 2014). In addition, previous studies have suggested
168 that patterns of moth community composition remain consistent despite seasonal turnover
169 (Summerville and Crist 2003).

170 Within each stand a heath trap was placed 15 metres from the edge, at least 200m from the
171 next nearest trap and the location recorded with a GPS. Traps were selectively positioned to ensure
172 that similar light levels were emitted (e.g. avoiding vegetation obscuring the light). In most cases, the
173 traps were not visible from each other, apart from in felled stands. This may introduce a bias in traps
174 at felled sites as the lights were visible from further away, reducing spatial independence (Lacki et al.
175 2007) although the attraction radii of heath light traps is commonly only between 10 – 30m
176 depending on moth family (Truxa & Fiedler 2012). Any moths attached to the outside of the trap at
177 the end of the trapping session were gently removed and released. A cotton wool ball soaked in
178 ethyl acetate was immediately added to the trap and left overnight to kill trapped invertebrates.
179 Macro moths were removed and pinned to boards for later identification and micro moths were
180 separated for identification by an expert at the National Museum of Scotland. Approval for this work
181 was obtained from the Ethical Review Committee within the Department of Biological &
182 Environmental Sciences at the University of Stirling. Species data were shared with local moth
183 recorders and added to the National Moth Monitoring Scheme (Fox et al. 2010).

184 *2.2 Local habitat characteristics*

185 We carried out vegetation surveys in two 0.01 ha plots at each stand type; due to the homogenous
186 nature of stands these plots were considered representative of the stand as a whole. At each plot we

187 recorded the total number of trees with diameter at breast height greater than 7 cm (stand density)
188 and recorded the dominant ground cover (vegetated / non vegetated). Since dead wood is
189 important for saproxylic moths we assessed the amount of dead wood on the forest floor using the
190 following scale: 0 – no coarse woody debris, 1 – small twigs, 2 – large twigs and branches over 7cm
191 in diameter, 3 – both large and small branches. Understory vegetation height was measured at 10
192 evenly spaced points across the radius of the circle and canopy cover was recorded at each point
193 using a sighting tube with an internal crosshair; if the crosshair intersected with any canopy
194 vegetation presence of canopy cover was recorded and converted to a percentage cover score
195 (Lintott et al. 2015).

196 *2.3 Landscape analysis*

197 The GUIDOS toolbox (Soille & Vogt 2009) was used to determine percentage cover of core (more
198 than 20m from the edge), and edge (patches within 20m of the edge) broadleaf tree cover woodland
199 and felled patches within 4km of each moth trap by combining data from the OS Mastermap (EDINA,
200 2014) and a high resolution Forestry Commission database specific to the study areas. Distance to
201 both broadleaf patches and felled areas as well as the size of the nearest broadleaf / felled patch
202 were also recorded. It should be noted that broadleaf cover could be remnants of deciduous
203 woodland cover from before the plantation was planted. Finally, the complexity of the broadleaf
204 patch (a score of the total area of broadleaf / felled divided by the total edge area of broadleaf /
205 felled) was calculated which approximates fragmentation (a highly fragmented area will have a high
206 complexity score, see Appendix 1 for details on landscape variables included in analysis).

207 *2.4 Statistical analysis*

208 All analysis was carried out using R (version 3.4, R core development team) using the following
209 packages: MuMIn, lme4, vegan, ggplot2. We used Margalef diversity to assess species diversity as it
210 is straightforward to interpret and because it can deal with occasions where the number of
211 individuals in a trap is equal to the number of species (Magurran 1988).

212 Many of the local and landscape variables were collinear so we used principle components
213 analysis (PCA) to remove collinearity and reduce the number of predictors. Three separate PCAs
214 were conducted for local characteristics and the felling and broadleaf tree cover metrics (See
215 Supplementary data 1 for an explanation of the variables included in the PCA). For each PCA we
216 retained those axes which explained more variation than random using the “broken stick” approach
217 (Jackson 1993). For the local characteristics (Local PC), the first two axes explained 77% of the
218 variation between stands; Local PC1 mainly described the stands with low canopy cover and high
219 understorey vegetation height (which loaded low on PC1) and stands with low vegetation cover and
220 high canopy cover (which loaded high on PC1), loosely catagorising different stand types
221 (Supplementary data 2, Figure A). Local PC2 was driven largely by differences in altitude, describing
222 the difference between the three different forests, with Galloway sites primarily at low altitudes,
223 Kielder stands predominantly at high altitudes and Cowal and Trossachs falling in between. For
224 felling characteristics (Felling PC), only the first axis explained more variation (63%) than chance;
225 stands with low values of Felling PC1 were closer to patches of clearfell and surrounded by greater
226 areas of felling in a 1km radius and those loading high on Felling PC1 were further from felled areas
227 with less overall felling in a 1km radius (Supplementary data 2, Figure C). For characteristics relating
228 to broadleaf woodland in the landscape (Broadleaf PC), only the first axis explained more variation
229 (67%) than by chance; stands loading high on Broadleaf PC1 tended to be further from smaller
230 patches of broadleaf woodland, with less broadleaf tree cover in the surrounding landscape whereas
231 sites loading low on Broadleaf PC1 were closer to larger broadleaf patches, with more overall
232 broadleaf tree cover in the surrounding habitat (Supplementary data, Figure B).

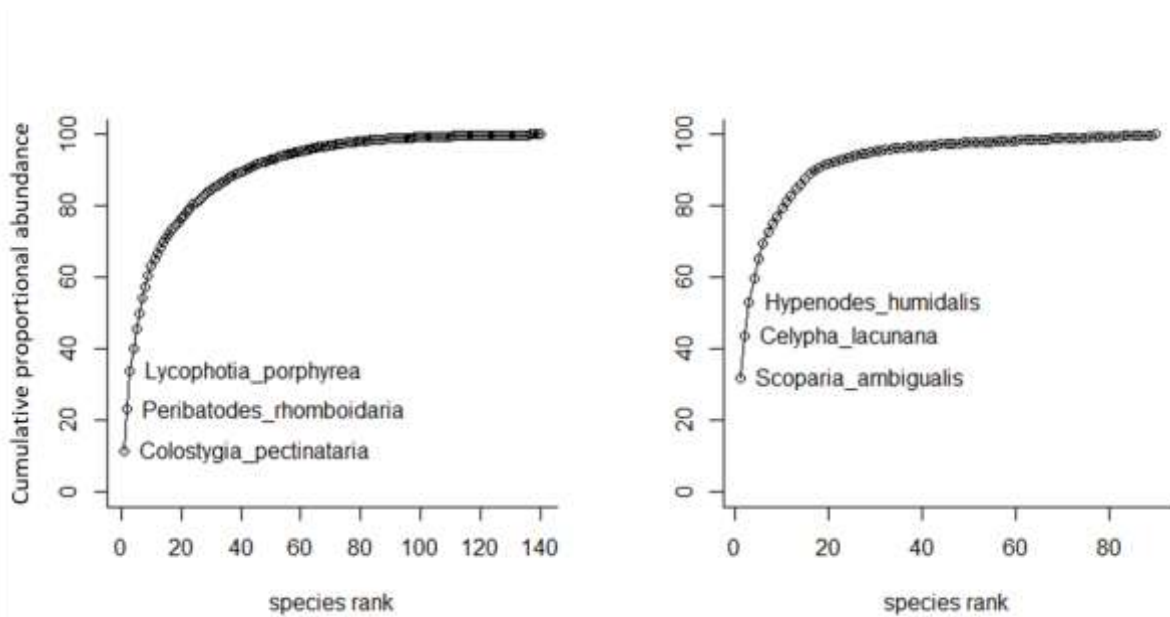
233 Using an information theoretic approach, we assessed the influence of stand and landscape
234 variables on the **abundance** and **species richness** of macro and micro moths separately, using each
235 metric per stand as the unit of replication. We used generalised linear models with a negative
236 binomial error structure to account for overdispersion, and included an interaction between latitude
237 and longitude as a fixed effect in all models to account for spatial autocorrelation. Models were

238 validated by visual assessment of the residuals (Crawley 2007). Continuous variables were
239 standardised and centred around a mean of zero and a standard deviation of 1 to allow direct
240 comparisons of estimates, and model fit was assessed by comparing the change in AIC, retaining the
241 best model (change in AIC greater than 2). McFaddens pseudo R^2 (McFadden 1974) was used to
242 assess the amount of variation explained by each model. Local PC2 was not used, as this mainly
243 described the difference in altitude between the stands and was collinear with date; in all cases
244 simply using date was a better predictor. Models were fitted using either the stand type or the Local
245 PC1, depending on model fit. We assessed the impact of felling and surrounding broadleaf tree cover
246 on each response measure including either Felling PC1 or Broadleaf PC1 separately, then together
247 and as an interaction. The same process was followed for **Margalef diversity** and **dominance** using a
248 Gaussian error distribution. For each response measure, if there was no clear “top” model we
249 averaged the coefficients across the top models in the set which accounted for a change in AIC of
250 less than 2, using full averaged models to reduce the bias from explanatory factors which do not
251 appear in every model (Burnham and Anderson 2002). Explanatory variables were considered to
252 have a “significant” effect on the responses if the standard error of the estimate did not cross zero
253 (Burnham & Anderson 2002). Micro and macro moths were analysed separately. Although the
254 distinction between macro moths and micro moths is not taxonomically supported, micro moths
255 typically have lower dispersal distances apart from some migratory species (Nieminen et al. 1999)

256 In addition to moth community measures outlined above, we modelled the influence of local
257 and landscape characteristics on the occurrence of six of 13 **BAP priority species** recorded in the
258 plantations. The following six species (*Eugnorisma glareosa* (Autumnal Rustic), *Arctia caja* (Garden
259 Tiger), *Celaena haworthii* (Haworths Rustic), *Xestia castanea* (Neglected Rustic), *Ecliptopera silaceata*
260 (Small Phoenix) and *Spilosoma lubricipeda* (White Ermine)) were present at the most sites and
261 represented species which have declined between 70 – 90% over the last ten years (Conrad et al.
262 2006). We had insufficient data to model abundance at stand-level, so presence of these species was
263 modelled using a binomial mixed effects model with species ID as a random intercept and Local PC1

264 as a random slope in order to assess species specific responses to stand level changes. We used the
265 same approach as the previous analyses but here visual inspection of the data and subsequent
266 model checking indicated that species occurrence was strongly and similarly associated with
267 distance to felled areas and the size of broadleaf patches, so these were used in preference to the
268 Felled and Broadleaf PC axis.

269 We graphically present the results for the single best model for each analysis including
270 standardised parameters and standard errors for all explanatory variables. Inferences were made by
271 comparing each parameter's standardised estimate with other predictor variables to assess its
272 relative importance, the upper and lower 95% quantiles of each parameter obtained from N = 2000
273 simulated draws from the estimated distribution (Lintott et al. 2014) and a comparison of selected
274 models using AIC.



275
276 Figure 2. Species rank abundance curves for macro and micro moths considered separately. The
277 three most abundant species are named. Rank abundances are given as cumulative proportions of
278 total abundance.

279 3. Results

280 Composition of moth populations in commercial coniferous plantations

281 We collected a total of 8074 moths comprising 6464 macro moths belonging to 140 species and 10
282 families, and 1762 micro moths, belonging to 90 species and 19 families (Supplementary data 1) over
283 170 trap nights. Of these, 60% were generalist species while only 14% were woodland specialists and
284 26% were associated with open habitats (open specialists). We recorded an average of 38 (± 4.2)
285 macro moth species and 10 (± 1.5) micro moth species per stand. Community composition was
286 dominated by a few, highly abundant species such as the micro moth *Scoparia ambigualis*
287 (Crambidae) and the macro moth *Colostygia pectinataria* (Geometridae), with less than 20% of
288 micro moth species accounting for over 80% of all micro moths collected and 34% of macro moth
289 species accounting for over 80% of all macro moths (Figure 2). We recorded 13 BAP priority species,
290 with an average of 3.2 ± 0.6 per stand.

291 3.1 Influence of local characteristics on moth communities

292 After accounting for date and temperature, there was relatively little correlation between local
293 characteristics and moth communities (Table 1), with correlations between Local PC1 and macro
294 moth abundance only. Abundance was highest in stands with a low Local PC1 score (low canopy
295 cover and high understorey vegetation height), falling 53% in older stands with a closed canopy and
296 lower understorey vegetation height. Fewer moths of both groups were collected later in the season,
297 with a similar pattern for species richness and diversity, but not dominance. Finally, the interaction
298 between latitude and longitude influenced richness, abundance and diversity for micro moths but
299 not macro moths (Table 1) revealing regional differences in species richness and abundance, with
300 the highest abundance in Galloway plantation (27.0 ± 3) and lower in Kielder (13.0 ± 1.6) and Cowal
301 and Trossachs (8.5 ± 1.3).

302 Table 1. Best approximating GLM's assessing influence of local, felling and broadleaf parameters on moth richness, abundance, diversity and dominance,
 303 conducted using an information theoretic approach with model averaging to assess importance of parameters. NA's indicate parameters not included in the
 304 top model sets. Dominant ground cover, coarse woody debris and the interaction between Felling PC1 and Broadleaf PC1 was never included in any top
 305 models and are not presented here. Parameters in bold are those which have a significant effect on response values, determined by whether the standard
 306 error of the estimate crosses zero (Burnham & Anderson 2002). Akaike's weight is the total weight explained by all models. Averaged estimates are
 307 presented \pm the standard error.

		No. models averaged across	Intercept	Local PC1	Felling PC1	Broadleaf PC1	Date	Temp	Lat:Long	Akaike's weight
Macro moths										
	Sp. Richness	7	264.9 \pm 177.6	-0.70 \pm 0.20	0.27 \pm 0.08	-0.10 \pm 0.00	-0.35 \pm 0.11	0.01 \pm 0.05	0.27 \pm 0.34	0.62
	Abundance	5	3.21 \pm 0.17	-0.26 \pm 0.11	0.49 \pm 0.12	-0.38 \pm 0.17	-0.61 \pm 0.19	0.16 \pm 0.16	0.21 \pm 0.39	0.63
	Marg. Diversity	6	2.01 \pm 0.18	-0.11 \pm 0.12	0.34 \pm 0.11	-0.33 \pm 0.14	-0.39 \pm 0.15	0.02 \pm 0.07	0.30 \pm 0.42	0.64
	Simp. diversity	9	1.22 \pm 0.19	-0.04 \pm 0.12	-0.03 \pm 0.10	NA	0.08 \pm 0.21	0.04 \pm 0.11	NA	0.27
Micro moths										
	Sp. Richness	4	1.23 \pm 0.15	-0.02 \pm 0.06	0.24 \pm 0.09	-0.28 \pm 0.11	-0.10 \pm 0.13	0.33 \pm 0.10	1.54 \pm 0.40	0.77
	Abundance	2	2.28 \pm 0.20	NA	0.47 \pm 0.12	-0.45 \pm 0.17	-0.32 \pm 0.25	0.42 \pm 0.15	1.93 \pm 0.50	0.73
	Marg. Diversity	4	1.07 \pm 0.10	-0.01 \pm 0.03	0.13 \pm 0.06	-0.14 \pm 0.10	-0.13 \pm 0.10	0.30 \pm 0.07	0.86 \pm 0.24	0.65
	Simp. diversity	13	0.75 \pm 0.17	-0.04 \pm 0.11	NA	0.01 \pm 0.05	-0.03 \pm 0.11	0.10 \pm 0.16	NA	0.43

308

309

310 Table 2: Best approximating GLM's assessing influence of local, felling and broadleaf parameters on BAP moth species probability of being detected. These
 311 were conducted using an information theoretic approach with model averaging to assess importance of parameters. NA's indicate parameters which were
 312 not included in the model. Dominant ground cover, coarse woody debris and the interaction between Felling PC1 and Broadleaf PC1 was never included in
 313 any top models and is not presented here. Parameters in bold are those which have a significant effect on response values, determined by whether the
 314 standard error of the estimate crosses zero (Burnham and Anderson 2002). Akaike's weight is the total weight explained by all models. Estimates for the full
 315 averaged model are presented \pm the standard error. Estimates provided for the top 7 models, with a change in AIC of less than 2. The same variables as for
 316 the overall moth communities were originally used but inspection of the broadleaf and felling PC output showed that the main relationships were with
 317 specific components of the principle components.

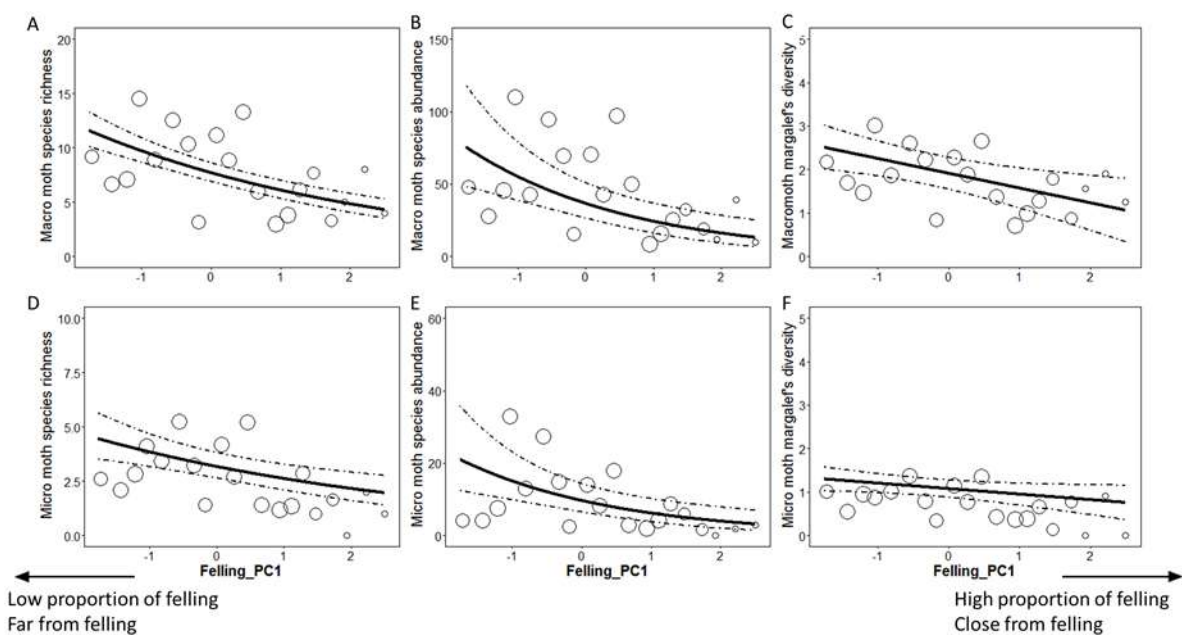
	Intercept	Size of nearest broadleaved patch	Altitude	Distance to felled stand	Lat:Long	Local_PC1	AICc	Akaike's weight
Averaged Model	-2.88 \pm 0.25	0.22 \pm 0.09	-0.04 \pm 0.10	0.16 \pm 0.14	0.02 \pm 0.09	-0.44 \pm 0.35		0.50
1	-2.95	0.22	NA	0.22	NA	-0.59	479.90	0.13
2	-2.92	0.22	NA	NA	NA	-0.53	480.92	0.08
3	-2.97	0.23	-0.13	0.24	NA	-0.59	480.96	0.07
4	-2.67	0.21	NA	0.21	NA	NA	481.17	0.07
5	-2.96	0.23	NA	0.20	0.09	-0.59	481.53	0.06
6	-2.98	0.27	-0.22	0.21	0.18	-0.59	481.56	0.05
7	-2.67	0.21	NA	NA	NA	NA	481.82	0.05

318

319

320 3.2 Influence of felling on moth communities

321 There appeared to be a large, negative impact of clear felling on species richness, abundance and
322 diversity for both macro- and micro moths (Figure 3, Table 1). Macro moth species richness declined
323 from 13.4 (9.3 – 19.4) in sites furthest from clear felled areas and with less felling within 1km to 4.0
324 (2.5 – 6.6) in sites nearest to felled areas or surrounded by more felling in 1km. Similarly, micro moth
325 species richness fell from 4.2 (2.9 – 6.2) to 1.5 (0.9 – 2.5) in sites close to felling or with a greater
326 proportion of felling in the surrounding landscape (Figure 3 A, D).



327

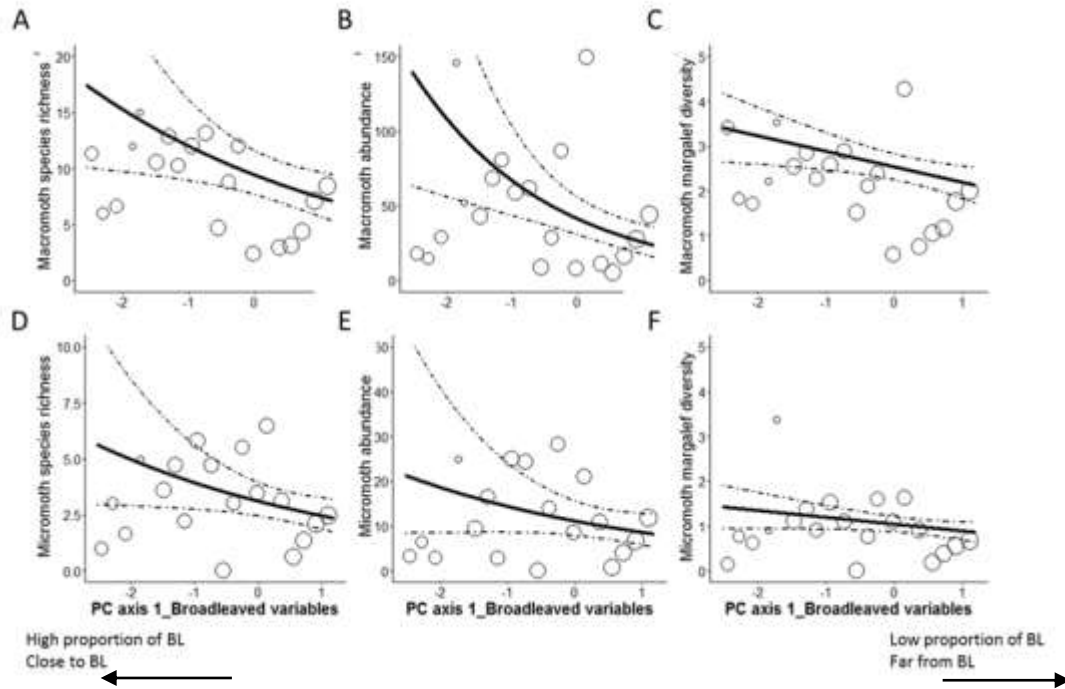
328

329 Figure 3. Impacts of felling on (A – C) Macro moth species richness, abundance and diversity and (D
330 – F) Micro moth species richness, abundance and diversity per site. Different scales are used for
331 abundance and richness due to higher richness and abundance in macro moths compared to micro
332 moths. Original data on richness, abundance and diversity are superimposed as grey circles with
333 diameter proportional to the number of sampling points where mean values occurred. Dashed lines
334 represent 95% confidence intervals around the predictions (solid line).

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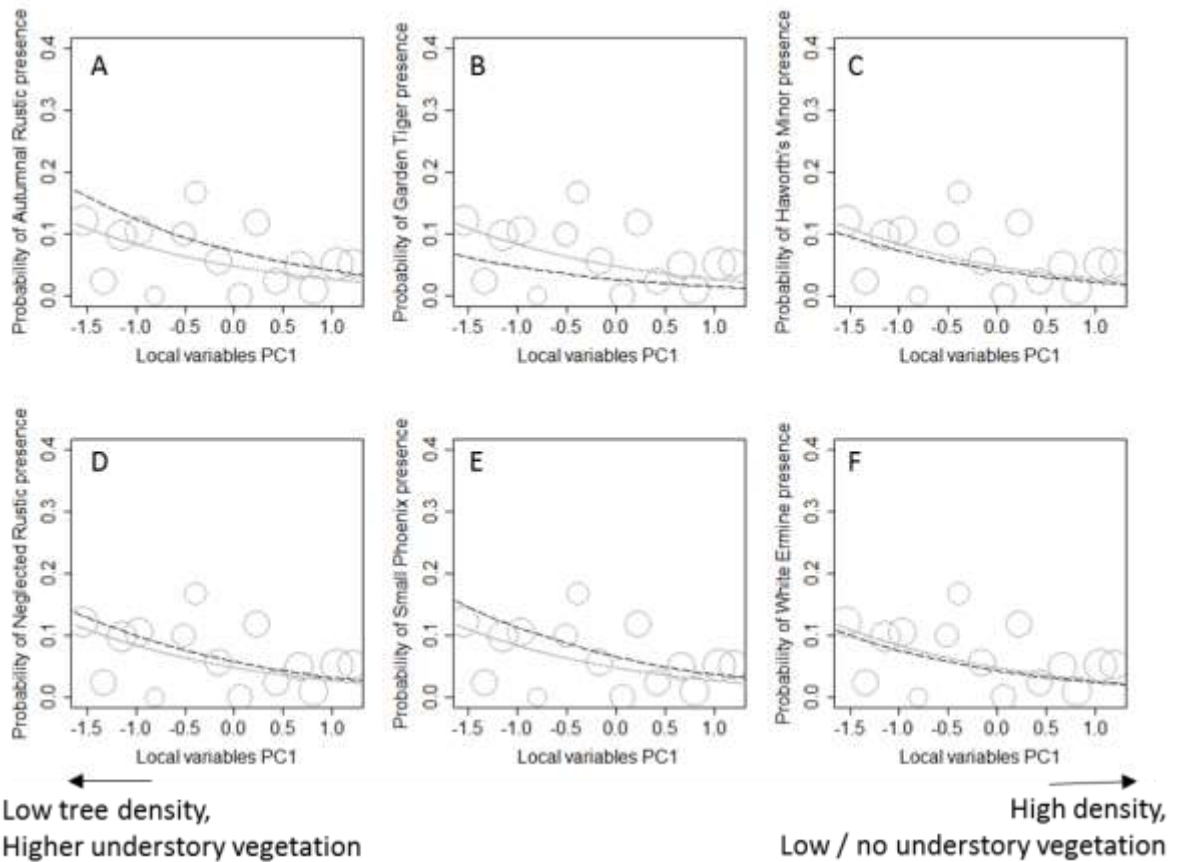
339 Fig. 4. Impacts of the amount and proximity of broadleaf woodland (BL) on (A – C) Macro moth
340 species richness, abundance and diversity, and (D – F) Micro moth species richness, abundance and
341 diversity per stand. Different scales are used for abundance and richness due to higher richness and
342 abundance in macro moths compared to micro moths. Original data on richness, abundance and
343 diversity are superimposed as grey circles with diameter proportional to the number of stands
344 where mean values occurred. Dashed lines represent 95% confidence intervals around the
345 predictions.

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352 Figure 5. Probability of recording priority BAP species by Local PC1 scores (associated with a shift
 353 from stands with low canopy cover and taller vegetation height to stands with high canopy cover
 354 and low vegetation height). Sites low on PC1 are predominantly clear fell and young, moving to
 355 thinned and mature aged stands loading high on PC1. Dotted lines are species specific, whilst the
 356 grey line shows the trend across all six BAP species. Original data on richness, abundance and
 357 diversity are superimposed as grey circles with diameter proportional to the number of stands
 358 where mean values occurred.

359 Both micro and macro moth abundance responded strongly to Felling PC1; macro moth abundance
 360 decreased from 68.0 (40.0 – 114.0) moths in sites far from felling or with a low proportion of felling
 361 in the surrounding landscape to 10.0 (5. 4 – 18.5) in sites closer to felling or with more felling in the
 362 surrounding landscape, and micro moth abundance decreased from 25.0 (14.0 – 42.0) individuals to

363 2.5 (1.3 – 4.9) individuals (Figure 3 B, E). There was little response of diversity of either group to the
364 proximity or prevalence of felling (Figure 3 C, F).

365 *3.3 Effects of the presence of broadleaf tree cover on moth communities*

366 In general, the proximity and amount of broadleaf tree cover within 4km of sampling sites appeared
367 to be positively associated with species richness, abundance and diversity for both macro and micro
368 moths, although the effect was smaller than the impact of felling (Table 1, Figure 4). The effect is
369 clearest for species richness, with richness of macro moths in stands nearest to the largest patches
370 of broadleaf tree cover double that of stands furthest from smaller patches of broadleaf, increasing
371 from 7.0 (5.0 – 9.0) species to 15.0 (8.0 – 29.0) species per stand (Figure 4 A). Similarly, micro moth
372 richness increased from 2.5 (1.3 – 3.1) species in stands far from broadleaf tree cover and with a low
373 proportion of broadleaf in the surrounding area to 5.0 (3.0 – 10.0) species richness in stands closest
374 to broadleaf patches or with a high proportion of broadleaf tree cover in the surrounding landscape
375 (Figure D). Whilst the influence of broadleaf woodland on abundance of both groups is similar, the
376 relationship appears to be weaker than for species richness (Figures 4B, E), and for macro moths
377 appears to be driven by high abundance at one site (Figure 4 B). Neither local variables, felling nor
378 broadleaf characteristics had any correlation with macro or micro moth dominance.

379 *3.4 Influence of local characteristics, felling and broadleaved woodland on BAP priority species*

380 The likelihood of catching a BAP species increased further from felled areas, and as the size of the
381 nearest broadleaf patch increased, with all six species having very similar response to both variables.
382 However, the correlation of Local PC1 with micro moth presence differed between the BAP priority
383 species. *Eugnorisma glareosa* (Autumnal Rustic; Figure 5 A) and *Ecliptopera silaceata* (Small Phoenix;
384 Figure 5 E) responded relatively strongly to Local PC1, and were more likely to be recorded in open
385 stands with taller vegetation whereas there was relatively little change in the probability of capture
386 for *Arctia caja* (Garden Tiger; Figure 5 B).

387 **4. Discussion:**

388 Here, we demonstrate that plantations can support large communities of moths, including several
389 BAP priority species. Lepidoptera are one of the most abundant and diverse insect orders, but are
390 currently undergoing widespread declines across Europe (Fox et al. 2013). Loss of habitat and
391 changes to silvicultural practices in native woodlands have been cited as drivers of these losses, but
392 to date the value of coniferous plantations for moths has been ignored due to their perception as
393 being a poor habitat for biodiversity.

394 Moth abundance was dominated by generalist species which are preferentially found in heath or
395 bog habitats, or by a small number of conifer specialist species. Macro moth abundance was highest
396 in relatively low density stands with vegetation cover, which are more likely to support appropriate
397 larval host plants, compared to dense stands with predominantly bare or moss as dominant ground
398 cover. In addition, sites loading low on Local PC1 were often recently felled and young stands with
399 large amounts of dead wood remaining which would benefit saproxylic species (Thorn et al. 2015).
400 However, we saw no effect of stand characteristics on species richness or diversity in macro moths
401 or for any micro moth response metric, possibly because we captured a high proportion of generalist
402 moth species which have less strict habitat associations. We have no data on the species
403 composition of moth communities prior to afforestation but it is likely to include species which
404 specialise on low nitrogen, open habitats.

405 Felling was strongly and negatively correlated with both macro- and micro moth species
406 richness, abundance and diversity. Macro and micro moth species richness was three times higher in
407 sites furthest from felling, and with fewer felled patches in the immediate landscape, whereas
408 abundance for macro and micro moths was between 7 and 10 times higher in sites further from
409 felling and with less felling in the surrounding landscape. This reflects patterns reported from
410 managed native broadleaf forests in Indiana, which found that clear felling significantly reduced
411 moth species richness compared to either no management or selective felling (Summerville & Crist

412 2002). Clear felling causes substantial changes in the floristic composition of the forest habitat and
413 through substantial changes in microclimate, to herbaceous ground cover and host plant availability
414 (Summerville 2011). Summerville (2011, 2013) similarly found that species richness of moths was
415 40% lower after timber removal, with the impacts of felling persisting up to 200m from the cleared
416 site itself.

417 The nature of the landscape matrix stands are embedded in may impede or facilitate
418 dispersal between habitat types (Tscharntke et al. 2012); if there is too much felling in the
419 surrounding landscape it may impede moth movements. Felled stands themselves may still be
420 attractive to particular moth species due to intermediate levels of disturbance allowing pioneer and
421 to some degree specialist species to coexist (Hamer et al. 2003). Indeed, in simplified landscapes,
422 characterised by high disturbance, dynamics in habitat patches are likely to be determined by the
423 availability of landscape wide remnant communities, particularly for species able to disperse over
424 wide distances (Tscharntke et al. 2012).

425 Disturbed habitats are often characterised by a high abundance of a few generalist species,
426 with the same subset of taxa dominating local stands and at the regional level. The majority of the
427 moths we trapped were generalist species (Supplementary data 1), this may reflect the fact that
428 moths using the plantations are those which can persist in a disturbed environment, as generalist
429 species are more resilient to disturbance (Franzén & Johannesson 2007). For example, although 14%
430 of all the moths we recorded are deciduous specialist feeders, the tree species they specialise on are
431 often planted as deciduous tree cover in plantations (Tallamy & Shropshire 2009). It is not possible
432 to tell from our study whether moth populations in plantations differ significantly from those in
433 native broadleaf woodlands. However, due to the levels of disturbance caused by felling and the
434 potential lack of host plants, as well as the predominance of generalist species we found in our
435 plantation sites, we would expect plantation woodlands to support a less diverse moth population
436 than broadleaf woodlands do. Macro moth species richness in the plantations was similar to that

437 found in broadleaf woodlands within an agricultural matrix, although abundance was lower, while
438 micro moth richness was 25% higher than in agricultural woodlands (Fuentes-Montemayor et al.
439 2012). Micro moth richness was similar to that reported from urban woodlands, but macro moth
440 richness was 40% higher in plantations (Lintott et al. 2014). It is surprising that similar or lower
441 species richness and abundance was found in urban (Lintott et al. 2014) and agricultural woodlands
442 (Fuentes-Montemayor et al. 2012). It would be interesting to determine whether this is due to
443 geographical differences (sites surveyed by Fuentes-Montemayor et al. 2012, and Lintott et al. 2014
444 were in Scotland but further north than the majority of sites surveyed for this study) or whether
445 woodlands surrounded by agricultural and urban land are similarly disturbed habitats due to a more
446 hostile matrix (Tschardt et al. 2012), although the drivers of disturbance may differ.

447 Continuous cover forestry, which involves the continuous and uninterrupted maintenance of
448 forest cover and avoids clear felling (Pommerening & Murphy 2004), has been advocated as an
449 alternative forest management system. The UK forest standard requires managers to identify areas
450 “which can be managed under a continuous cover forestry system and build them into forest design”
451 (Mason et al. 1999). Despite not being appropriate for widespread use in all plantation forests due
452 to the potential risk of wind damage to stands, there is evidence to suggest that multi aged systems
453 may be more resilient to impacts of wind (O’Hara & Ramage 2013) and the potential forest health
454 and yield benefits are increasingly recognised, with over 10% of Forestry Commission woodlands
455 now under continuous cover management (Macdonald et al. 2009; O’Hara & Ramage 2013).
456 Switching to continuous cover forestry may benefit moth communities; in Indiana (USA) Summerville
457 et al (2009) found that shelterwood harvesting (removal of 15% biomass and similar in concept to
458 continuous cover forestry) did not reduce functional and compositional resilience of lepidopteran
459 communities compared to group selection harvesting (80% of tree biomass removed) and clear
460 felling which had a significant negative impact. Additionally, moth communities showed signs of
461 recovery within three years compared to other studies showing impoverished moth communities up
462 to 60 years after clear felling (Summerville 2013; Summerville et al. 2009)

463 We found that the amount and proximity of broadleaf tree cover positively influenced moth species
464 richness, and to some extent abundance. Many native tree species such as *Betula*, *Quercus* and *Salix*
465 have large numbers of moth species associated with them (Tallamy & Shropshire 2009) and are
466 commonly planted in conifer plantations as broadleaf regeneration trees. Fuentes-Montemayor et al
467 (2012) found that species richness was highest in woodland with no conifers, so increasing landscape
468 heterogeneity by planting patches of broadleaf tree cover within the plantation landscape may be
469 invaluable islands allowing moth species to persist within the plantation matrix despite felling
470 disturbance.

471 We recorded 13 BAP priority species using plantation woodlands. BAP priority species are so
472 designated due to their rapidly declining populations across the United Kingdom and the need for
473 further scientific study in order to assess and understand their population declines (Conrad et al.
474 2006). Of these, seven were present in fewer than 10 sites and were removed from further
475 modelling. Of the six remaining species, all are habitat generalists or conifer and moorland habitat
476 specialists. These species responded to stand type characteristics (separated by local PC1)
477 differently. The Autumnal Rustic (*Eugnorisma glareosa*) and the Small Phoenix (*Ecliptopera silaceata*)
478 were most likely to be detected in open stands with low canopy cover and stand density; the
479 Autumnal Rustic is a generalist species often associated with moorland habitats which constitute a
480 large proportion of the surrounding landscape and the Small Phoenix is a conifer specialist, and
481 therefore likely to thrive in conifer plantations. All BAP species were significantly less likely to be
482 recorded in stands closer to felled areas regardless of the size of the felled area or the proportion of
483 felling in the surrounding area which, considering the two species' preference for open stands is
484 somewhat surprising. All BAP species also responded equally positively to the size of the nearest
485 patch of broadleaf tree cover. Broadleaf patches within plantations are not part of active harvesting
486 programs, and are maintained or increased to meet biodiversity and restructuring guidelines (Watts
487 et al. 2008), so may provide a potential source from which moth species can disperse.

488 *4. 1 Management recommendations:*

489 Worldwide, forest managers increasingly recognise the importance of sustainable forest
490 management to improve biodiversity, but exotic pine plantations have received relatively little
491 attention for their potential contribution to moth communities above and beyond the impacts of
492 pest moth species. However, we found similar or higher levels of abundance and diversity compared
493 to fragmented urban and agricultural woodlands in nearby regions (Lintott et al. 2014; Fuentes-
494 Montemayor et al. 2012), and more BAP priority species in conifer plantations than urban
495 woodlands (Lintott et al. 2014). We found that moth richness, abundance and diversity were
496 influenced by plantation management and consider that the following should be taken into account
497 when considering how plantation management may affect moth communities:

498 **1. Switching to continuous cover forestry:**

499 Similar to other studies in native woodlands under felling pressure (Summerville 2014;
500 Summerville 2011; Summerville & Crist 2002; Summerville 2013; Summerville et al. 2009), felling
501 significantly affected moth populations in our study sites, reducing species richness and
502 abundance. Since clear felling was the only timber extraction technique used at our sites we
503 were not able to compare with other lower-intensity methods. Switching to continuous cover
504 forestry where appropriate will benefit moth communities and in turn the small mammal, bird
505 and bat species which rely on them as a prey source while not negatively impacting forest
506 productivity (Macdonald et al. 2009).

507 **2. Maintaining broadleaf woodland:**

508 Moth abundance and richness was far higher close to broadleaf tree cover; continued replanting
509 of broadleaf trees and reduced intensity of management where possible near broadleaf stands
510 should benefit both micro and macro moth richness and abundance. Many moth species can
511 only disperse over relatively short distances (Merckx et al. 2012), therefore increasing the
512 amount and connectivity of broadleaf woodland may allow moth species to persist within and

513 disperse throughout plantations. All BAP priority species responded strongly to the size of the
514 nearest patch of broadleaf tree cover, so reducing forestry operations near large patches of
515 broadleaf trees is likely to benefit moth communities in general and BAP species in particular.

516 **3. Monitoring BAP priority species in plantations:**

517 Of all the BAP priority species, the Garden Tiger (*Arctia caja*) moth was of particular interest as it
518 is a conspicuous species that has declined widely across the UK, possibly due to climatic changes
519 such as warmer wetter winters (Conrad 2002). More northerly habitats may be essential for the
520 persistence of this species, and low density plantation stands may be an important refuge for
521 this species in the face of future climate change. In addition, the Autumnal Rustic (*Eugnorisma*
522 *glareosa*) which was abundant in plantation sites, has undergone substantial declines
523 throughout the UK, thought to be related to pesticide use. Plantation sites should be included in
524 long term monitoring programs to understand further how BAP priority species are using
525 plantation woodlands.

526 Moth populations in Sitka spruce plantations appear to be predominantly generalist species, which
527 may imply a disturbed community (Summerville et al. 2009). However, the presence of some BAP
528 species demonstrates the importance of surveying sites that may historically be perceived as poor
529 for biodiversity. With sympathetic management, plantation forests may have a role to play in
530 preserving and supporting moth populations, particularly as climate change may result in changing
531 species distributions.

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540 **References:**

- 541 Boye, P. & Dietz, M., 2005. Development of good practice guidelines for woodland
542 management for bats. *English Nature Research Reports*, 661, pp.1–89.
- 543 Brockerhoff, E.G. et al., 2008. Plantation forests and biodiversity: Oxymoron or opportunity?
544 *Biodiversity and Conservation*, 17(5), pp.925–951.
- 545 Burnham, K. & Anderson, D., 2002. *Model selection and multimodel inference: a practical*
546 *information theoretic approach*, Springer Verlag, New York.
- 547 Conrad, K.F. et al., 2006. Rapid declines of common, widespread British moths provide
548 evidence of an insect biodiversity crisis. *Biological Conservation*, 132(3), pp.279–291.
- 549 Crawley, M., 2007. *The R Book*, West Sussex: John Wiley & Sons.
- 550 Dirzo, R. et al., 2014. Defaunation in the Anthropocene. *Science*, 345(6195), pp.401–406.
551 Available at:
552 <http://www.sciencemag.org/cgi/doi/10.1126/science.1251817>
553 <http://www.sciencemag.org/content/345/6195/401.short>.
- 554 Fox, R. et al., 2010. Moths count : recording moths for conservation in the UK. *Journal of*
555 *Insect Conservation*, 15, pp.55–68.
- 556 Fox, R., Parsons, M. & Chapman, J., 2013. The State of Britain’s Larger Moths 2013. *Butterfly*
557 *Conservation and Rothamsted Research, Wareham, Dorset, UK*, pp.1–32. Available at:
558 [http://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britai](http://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britain+s+Larger+Moths+2013#1%5Cnhttp://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britain+s+Larger+Moths+2013#0)
559 [n+’+s+Larger+Moths+2013#1%5Cnhttp://scholar.google.com/scholar?hl=en&btnG=Se](http://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britain+s+Larger+Moths+2013#1%5Cnhttp://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britain+s+Larger+Moths+2013#0)
560 [rch&q=intitle:The+State+of+Britain’s+Larger+Moths+2013#0](http://scholar.google.com/scholar?hl=en&btnG=Search&q=intitle:The+State+of+Britain+s+Larger+Moths+2013#0).
- 561 Franzén, M. & Johannesson, M., 2007. Predicting extinction risk of butterflies and moths
562 (Macrolepidoptera) from distribution patterns and species characteristics. *Journal of*
563 *Insect Conservation*, 11(4), pp.367–390.
- 564 Fuentes-Montemayor, E. et al., 2012. Factors influencing moth assemblages in woodland
565 fragments on farmland: Implications for woodland management and creation schemes.
566 *Biological Conservation*, 153, pp.265–275. Available at:
567 <http://dx.doi.org/10.1016/j.biocon.2012.04.019>.
- 568 Goulson, D., Lye, G.C. & Darvill, B., 2008. Decline and conservation of bumble bees. *Annual*
569 *review of entomology*, 53, pp.191–208. Available at:
570 <http://www.ncbi.nlm.nih.gov/pubmed/17803456>.
- 571 Hamer, K.C. et al., 2003. Ecology of butterflies in natural and selectively logged forests of
572 northern Borneo: The importance of habitat heterogeneity. *Journal of Applied Ecology*,

573 40(1), pp.150–162.

574 Humphrey, J.W., Ferris, R. & Quine, C., 2003. Biodiversity in Britain’s planted forests: Results
575 from the Forestry Commission’s Biodiversity Assessment Project. *Forestry Commission*,
576 *Edinburgh*, pp.51–62.

577 Jackson, D.A., 1993. Stopping Rules in Principal Components Analysis: A Comparison of
578 Heuristical and Statistical Approaches. *Ecology*, 74(8), pp.2204–2214.

579 Lacki, M.J. et al., 2007. Prey consumption of insectivorous bats in coniferous forests of
580 north-central Idaho. *Northwest Science*, 81(3), pp.199–205.

581 Lintott, P.R. et al., 2015. Differential responses to woodland character and landscape
582 context by cryptic bats in urban environments. *Plos One*, 10(5), p.e0126850. Available
583 at: <http://dx.plos.org/10.1371/journal.pone.0126850>.

584 Lintott, P.R. et al., 2014. Moth species richness, abundance and diversity in fragmented
585 urban woodlands: implications for conservation and management strategies.
586 *Biodiversity and Conservation*, pp.2875–2901.

587 Macdonald, E., Gardiner, B. & Mason, W., 2009. The effects of transformation of even-aged
588 stands to continuous cover forestry on conifer log quality and wood properties in the
589 UK. *Forestry*, 83(1), pp.1–16. Available at:
590 <http://forestry.oxfordjournals.org/cgi/content/long/83/1/1>.

591 Magurran, A.E., 1988. *Ecological diversity and its measurement.*, Princeton: Princeton
592 University Press.

593 Mason, W., Kerr, G. & Simpson, J., 1999. *What is continuous cover forestry?*, Information
594 Note.

595 Mattila, N. et al., 2006. Ecological determinants of distribution decline and risk of extinction
596 in moths. *Conservation Biology*, 20(4), pp.1161–1168.

597 McFadden, D., 1974. Conditional logit analysis of qualitative choice behaviour. In P.
598 Zarembka, ed. *Frontiers in Econometrics*. New York: Academic Press, pp. 105–142.

599 Merckx, T. et al., 2012. Conserving threatened Lepidoptera: Towards an effective woodland
600 management policy in landscapes under intense human land-use. *Biological*
601 *Conservation*, 149(1), pp.32–39. Available at:
602 <http://dx.doi.org/10.1016/j.biocon.2012.02.005>.

603 Nieminen, M., Rita, H. & Uuvana, P., 1999. Body size and migration in moths. *Ecography*,
604 22(6), pp.697–707.

605 O’Hara, K.L. & Ramage, B.S., 2013. Silviculture in an uncertain world: Utilizing multi-aged
606 management systems to integrate disturbance. *Forestry*, 86(4), pp.401–410.

607 Ober, H.K. & Hayes, J.P., 2010. Determinants of nocturnal Lepidopteran diversity and
608 community structure in a conifer-dominated forest. *Biodiversity and Conservation*,
609 19(3), pp.761–774.

610 Ohsawa, M., 2005. Species richness and composition of Curculionidae (Coleoptera) in a
611 conifer plantation, secondary forest, and old-growth forest in the central mountainous
612 region of Japan. *Ecological Research*, 20(6), pp.632–645.

613 Oxbrough, A. et al., 2010. Ground-dwelling invertebrates in reforested conifer plantations.
614 *Forest Ecology and Management*, 259(10), pp.2111–2121.

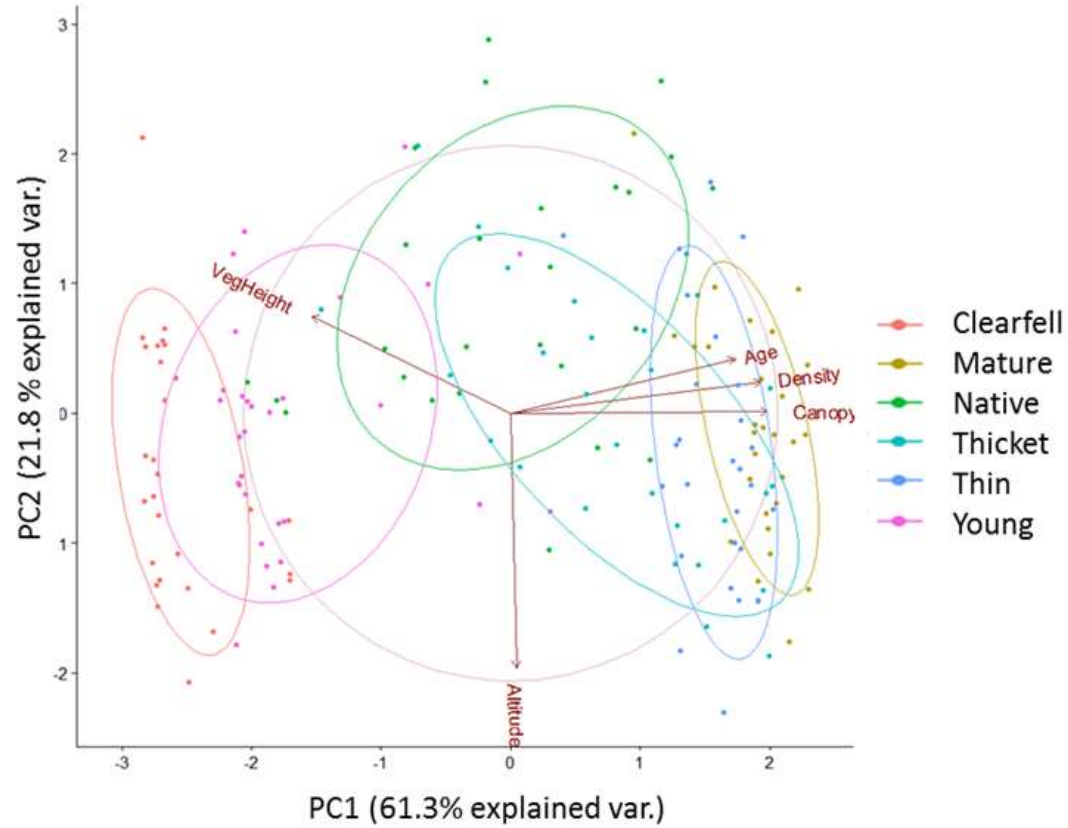
- 615 Pommerening, A. & Murphy, S.T., 2004. A review of the history , definitions and methods of
616 continuous cover forestry with special attention to afforestation and restocking. , 77(1).
- 617 Soille, P. & Vogt, P., 2009. Morphological segmentation of binary patterns. *Pattern*
618 *Recognition Letters*, 30, pp.456–459.
- 619 Summerville, K.S., 2014. Do seasonal temperatures, species traits and nearby timber harvest
620 predict variation in moth species richness and abundance in unlogged deciduous
621 forests? *Agricultural and Forest Entomology*, 16(1), pp.80–86.
- 622 Summerville, K.S., 2013. Forest lepidopteran communities are more resilient to shelterwood
623 harvests compared to more intensive logging regimes. *Ecological Applications*, 23(5),
624 pp.1101–1112.
- 625 Summerville, K.S., 2011. Managing the forest for more than the trees: Effects of
626 experimental timber harvest on forest Lepidoptera. *Ecological Applications*, 21(3),
627 pp.806–816.
- 628 Summerville, K.S., Courard-Hauri, D. & Dupont, M.M., 2009. The legacy of timber harvest:
629 Do patterns of species dominance suggest recovery of lepidopteran communities in
630 managed hardwood stands? *Forest Ecology and Management*, 259(1), pp.8–13.
- 631 Summerville, K.S. & Crist, T.O., 2002. Effects of timber harvest on forest Lepidoptera:
632 community, guild, and species responses. *Ecological Applications*, 12(3), pp.820–835.
- 633 Tallamy, D.W. & Shropshire, K.J., 2009. Ranking lepidopteran use of native versus
634 introduced plants. *Conservation Biology*, 23(4), pp.941–947.
- 635 Thomas, J.A., 2004. Comparative Losses of British Butterflies , Birds , and Plants and. ,
636 1879(2004).
- 637 Thorn, S. et al., 2015. Forest Ecology and Management Guild-specific responses of forest
638 Lepidoptera highlight conservation-oriented forest management – Implications from
639 conifer-dominated forests. *Forest Ecology and Management*, 337, pp.41–47. Available
640 at: <http://dx.doi.org/10.1016/j.foreco.2014.10.031>.
- 641 Truxa, C. & Fiedler, K., 2012. Attraction to light-from how far do moths (Lepidoptera) return
642 to weak artificial sources of light? *European Journal of Entomology*, 109(1), pp.77–84.
- 643 Tschardtke, T. et al., 2012. Landscape moderation of biodiversity patterns and processes -
644 eight hypotheses. *Biological Reviews*, 87(3), pp.661–685.
- 645 Warren, M.S. et al., 2001. Rapid responses of British butterflies to opposing forces of climate
646 and habitat change. , pp.65–70.
- 647 Warren, M.S. & Bourn, N. a D., 2011. Ten challenges for 2010 and beyond to conserve
648 Lepidoptera in Europe. *Journal of Insect Conservation*, 15(1), pp.321–326.
- 649 Watts, K. et al., 2008. Conserving forest biodiversity: recent approaches in UK forest
650 planning and management. *Patterns and Processes in Forest Landscapes*, pp.373–398.

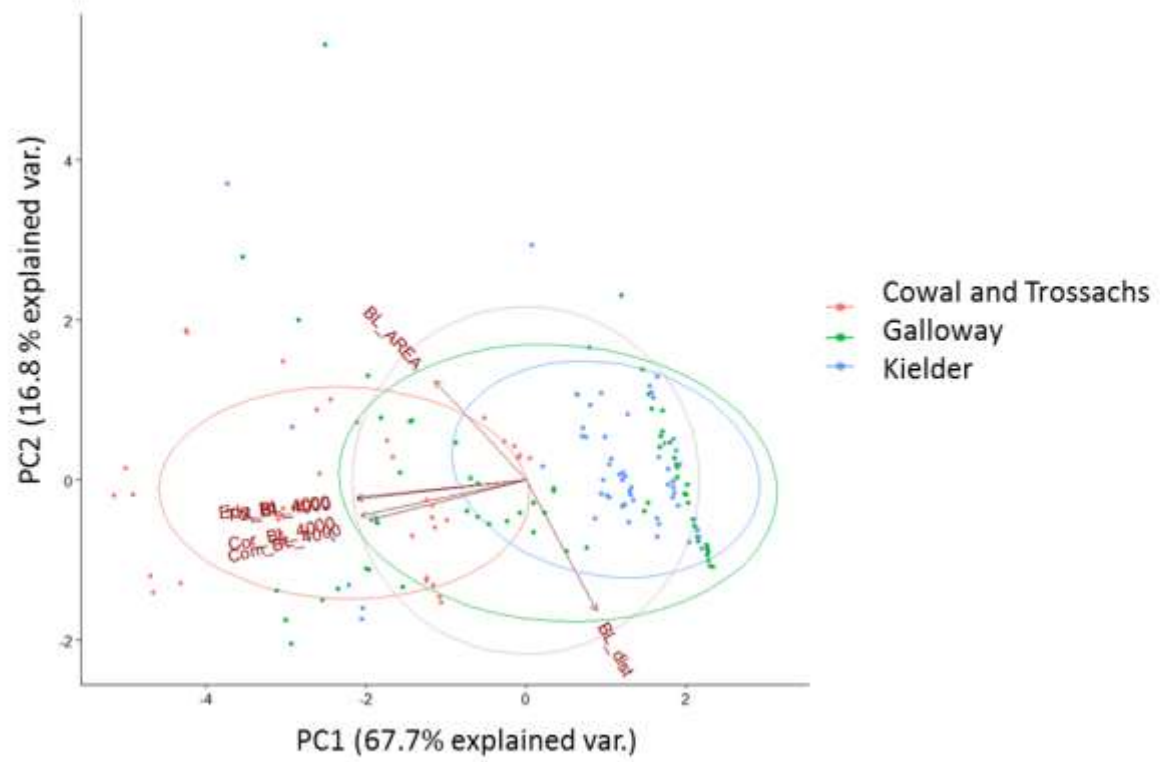
Supplementary data 1:

Table 1: Variables included in Principle Components Analysis.

PC axis	Measure	Unit	Minimum	Maximum	Median	Description
Local PC1	Altitude	m	83.8	466	230.7	Height above sea level
Local PC1	Density	trees per ha	0	3000	600	Number of trees per hectare
Local PC1	Veg height	mm	0	1744.1	156.6	Height of vegetation measured at 10 points across plot
Local PC1	Canopy cover	%	0	1	0.67	Total canopy cover as a percent
Local PC1	Stand Age	years	0	133	14	Stand age calculated from year of planting
Broadleaf PC1	BL_distance	m	0	3934	682	Distance in metres to nearest patch of mature broadleaf
Broadleaf PC1	BL_area	m ²	0.1	163.2	1.3	Size of nearest mature broadleaf patch
Broadleaf PC1	Tot_BL_4000	%	0	11.3	0.8	Total broadleaf cover as a % of a 4km ² circle
Broadleaf PC1	Edge_BL_4000	%	0	2.9	0.2	Edge broadleaf cover as % of a 4km ² circle
Broadleaf PC1	Core_BL_4000	%	0	4.9	0.05	Core broadleaf (at least 10m from an edge) as a % of a 4km ² circle
Broadleaf PC1	Com_BL_4000	%	0	2.1	0.3	Total area / Edge area - complexity of cover within the landscape
Felled PC1	FE_distance	m	0	2670	527	Distance in metres to nearest felled stand
Felled PC1	FE_area	m ²	0.04	92	13.9	Size of nearest felled stand
Felled PC1	Tot_FE_4000	%	0	35	5.1	Felled cover as a % of a 4km ² circle
Felled PC1	Edge_FE_4000	%	0	8	1.9	Edge felled cover as % of a 4km ² circle
Felled PC1	Core_FE_4000	%	0	26.5	2.4	Core felled (at least 10m from an edge) as a % of a 4km ² circle
Felled PC1	Com_FE_4000	%	0.8	2.1	1.5	Total area / Edge area - complexity of cover within the landscape

Supplementary data 2. Output from principle components analysis:





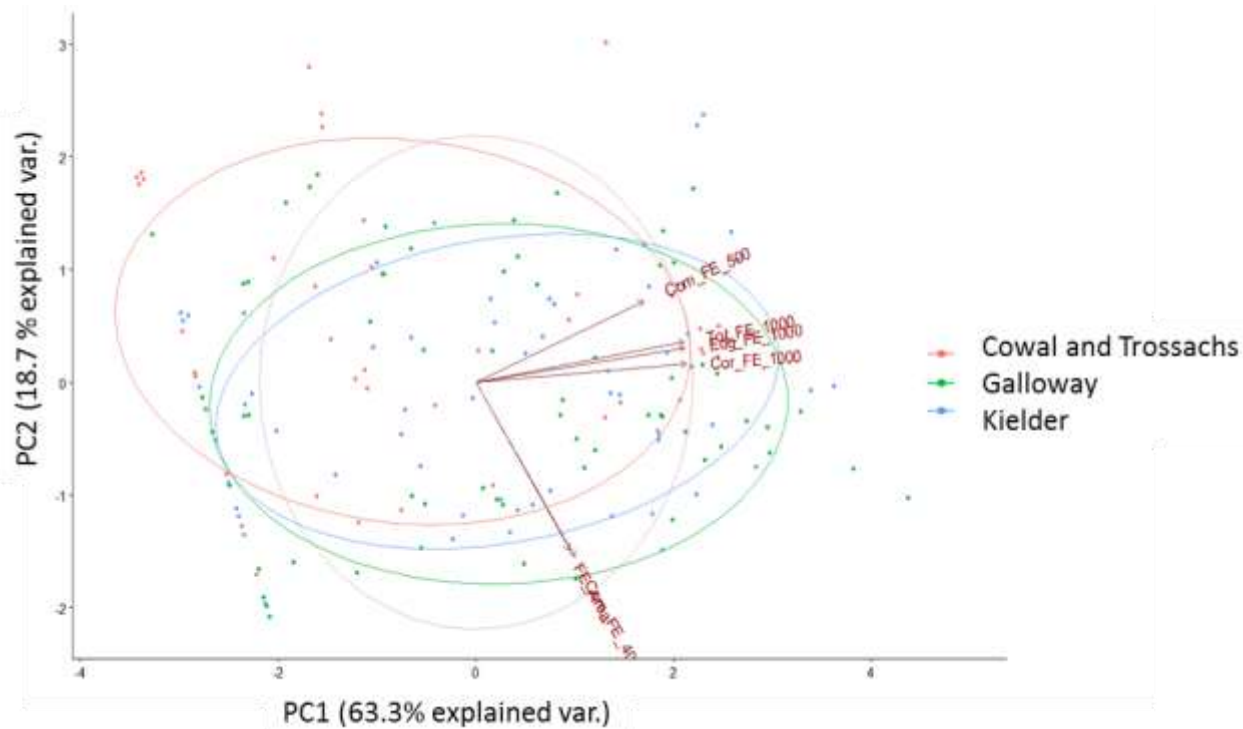


Figure 1. Principle components loadings for A) Local variables, B) Broadleaf variables and C) Felling variables. (see Supplementary data 1 for a description of the variables included in each PCA). Sites are coloured by stand type (Local PC) and by plantation (Broadleaf PC and Felling PC). Coloured ellipses delineate sites within each plantation that are similar to each other based on a normal probability distribution of 0.68. Dark red ellipsoid encompasses sites across all three plantations within a normal probability distribution of 0.68. Arrows indicate direction and magnitude of relationship, variables that are close together or directly opposite are highly correlated.

Supplementary data 3: Full list of moth species recorded as part of study

Table 1: Macro moth species:

Common name (Family)	Latin Name	Abundance per trap (\pm SE)	Habitat preference
Antler Moth (Noctuidae)	<i>Cerapteryx graminis</i>	0.24 \pm 0.08	Grassland
Autumnal Rustic (Noctuidae) ^a	<i>Eugnorisma glareosa</i>	0.47 \pm 0.17	Generalist
Barred Chestnut (Noctuidae)	<i>Diarsia dahlia</i>	0.01 \pm 0.01	Deciduous
Barred Red (Geometridae)	<i>Hylaea fasciaria</i>	1.21 \pm 0.33	Conifer
Barred Straw (Geometridae)	<i>Gandaritis pyraliata</i>	0.34 \pm 0.13	Generalist
Barred Umber (Geometridae)	<i>Plagodis pulveraria</i>	0.01 \pm 0.01	Deciduous
Beautiful Carpet Moth (Geometridae)	<i>Mesoleuca albicillata</i>	0.04 \pm 0.02	Deciduous
Beautiful Golden Y (Noctuidae)	<i>Autographa pulchrina</i>	0.39 \pm 0.10	Generalist
Bordered Beauty (Geometridae)	<i>Epione repandaria</i>	0.01 \pm 0.01	Deciduous
Bordered Gothic (Noctuidae)	<i>Sideridis reticulata</i>	0.01 \pm 0.01	Open ground
Bordered Pug (Geometridae)	<i>Eupithecia succenturiata</i>	0.01 \pm 0.01	Generalist
Bordered Sallow (Noctuidae)	<i>Pyrrhia umbra</i>	0.02 \pm 0.01	Grassland
Bordered White (Noctuidae)	<i>Bupalus piniaria</i>	0.07 \pm 0.03	Conifer
Bright Line Brown Eye (Geometridae)	<i>Lacanobia oleracea</i>	0.02 \pm 0.01	Generalist
Brimstone Moth (Noctuidae)	<i>Opisthograptis luteolata</i>	0.04 \pm 0.03	Generalist
Broom Moth (Crambidae)	<i>Ceramica pisi</i>	0.12 \pm 0.05	Moorland
Brown Rustic (Arctiidae)	<i>Elophila nymphaeata</i>	0.17 \pm 0.11	Deciduous
Buff Ermine (Erebidae)	<i>Spilosoma lutea</i>	0.08 \pm 0.03	Generalist
Buff Footman (Notodontidae)	<i>Eilema depressa</i>	0.19 \pm 0.13	Wood generalist
Buff Tip (Noctuidae)	<i>Phalera bucephala</i>	0.01 \pm 0.01	Deciduous
Burnished Brass (Geometridae)	<i>Diachrysia chrysitis</i>	0.12 \pm 0.04	Open ground
Chevron (Arctiidae)	<i>Eulithis testata</i>	0.01 \pm 0.01	Open ground
Clouded Border (Noctuidae)	<i>Tyria jacobaeae</i>	0.11 \pm 0.05	Deciduous
Clouded Bordered Brindle (Erebidae)	<i>Apamea crenata</i>	0.06 \pm 0.03	Grassland
Clouded Buff (Geometridae)	<i>Diacrisia sannio</i>	0.02 \pm 0.01	Moorland
Clouded Magpie (Geometridae)	<i>Abraxas sylvata</i>	0.04 \pm 0.03	Grassland
Common Carpet (Erebidae)	<i>Epirrhoe alternata</i>	0.23 \pm 0.06	Generalist
Common Footman (Drepanidae)	<i>Eilema lurideola</i>	0.01 \pm 0.01	Generalist
Common Lute String (Geometridae)	<i>Ochropacha duplaris</i>	0.01 \pm 0.01	Deciduous
Common Marbled Carpet (Noctuidae)	<i>Dysstroma truncata</i>	0.01 \pm 0.01	Wood generalist
Common Rustic (Hepialidae)	<i>Mesapamea secalis</i>	0.11 \pm 0.07	Generalist
Common Wainscot (Geometridae)	<i>Korscheltellus lupulina</i>	0.42 \pm 0.22	Grassland
Common Wave (Geometridae)	<i>Cabera exanthemata</i>	0.8 \pm 0.22	Deciduous
Coxcomb Prominent (Noctuidae)	<i>Cabera pusaria</i>	0.07 \pm 0.03	Deciduous
Dark Arches (Geometridae)	<i>Apamea monoglypha</i>	0.27 \pm 0.09	Generalist
Dark Brocade (Geometridae) ^a	<i>Xanthorhoe ferrugata</i>	0.17 \pm 0.10	Generalist
Dark Marbled Carpet (Noctuidae)	<i>Dysstroma citrata</i>	0.54 \pm 0.14	Generalist
Dark Tussock (Noctuidae)	<i>Abrostola triplasia</i>	0.01 \pm 0.01	Open ground
Dotted Carpet (Noctuidae)	<i>Aporophyla lutulenta</i>	0.01 \pm 0.01	Wood generalist

Dotted Clay (Noctuidae)	<i>Xestia baja</i>	0.22 ± 0.10	Generalist
Double Dart (Noctuidae) ^a	<i>Graphiphora augur</i>	0.02 ± 0.01	Wood generalist
Double Square Spot (Geometridae)	<i>Xestia triangulum</i>	0.27 ± 0.13	Deciduous
Double Striped Pug (Lasiocampidae)	<i>Gymnoscelis rufifasciata</i>	0.04 ± 0.02	Generalist
Drinker Moth (Noctuidae)	<i>Euthrix potatoria</i>	0.36 ± 0.09	Generalist
Dusky Brocade (Noctuidae) ^a	<i>Apamea remissa</i>	0.02 ± 0.02	Generalist
Dwarf Pug (Geometridae)	<i>Eupithecia tantillaria</i>	0.02 ± 0.01	Conifer
Ear Moth (Geometridae) ^a	<i>Amphipoea oculea</i>	0.08 ± 0.04	Generalist
Flame Carpet (Noctuidae)	<i>Selenia dentaria</i>	0.53 ± 0.14	Generalist
Flame Shoulder (Noctuidae)	<i>Ochropleura plecta</i>	0.58 ± 0.14	Generalist
Four Dotted Footman (Geometridae)	<i>Luperina testacea</i>	0.1 ± 0.05	Generalist
Foxglove Pug (Geometridae)	<i>Eupithecia pulchellata</i>	0.01 ± 0.01	Generalist
Frosted Orange (Noctuidae)	<i>Gortyna flavago</i>	0.01 ± 0.01	Generalist
Garden Carpet (Geometridae)	<i>Xanthorhoe fluctata</i>	0.13 ± 0.10	Generalist
Garden Tiger (Erebidae) ^a	<i>Arctia caja</i>	0.33 ± 0.12	Generalist
Gold Spangle (Noctuidae)	<i>Autographa bractea</i>	0.01 ± 0.01	Generalist
Gold Swift (Hepialidae)	<i>Phymatopus hecta</i>	0.02 ± 0.02	Generalist
Golden Rod Pug (Geometridae)	<i>Eupitheca virgaureata</i>	0.02 ± 0.01	Generalist
Golden Y (Noctuidae)	<i>Autographa jota</i>	0.09 ± 0.04	Generalist
Gothic (Noctuidae)	<i>Naenia typica</i>	0.01 ± 0.01	Deciduous
Green Arches (Noctuidae)	<i>Anaplectoides prasina</i>	0.01 ± 0.01	Generalist
Green Carpet (Geometridae)	<i>Colostygia pectinataria</i>	4.44 ± 0.86	Deciduous
Green Pug (Geometridae)	<i>Pasiphila rectangulata</i>	0.01 ± 0.01	Deciduous
Grey Arches (Noctuidae)	<i>Polia nebulosa</i>	0.01 ± 0.01	Generalist
Grey Dagger (Noctuidae) ^a	<i>Acronicta psi</i>	0.01 ± 0.01	Deciduous
Grey Mountain Carpet (Geometridae) ^a	<i>Entephria caesiata</i>	0.13 ± 0.05	Generalist
Grey Pine (Geometridae)	<i>Thera obeliscata</i>	0.03 ± 0.03	Moorland
Haworths Minor (Noctuidae) ^a	<i>Celaena haworthii</i>	0.18 ± 0.08	Conifer
Heath Rustic (Noctuidae) ^a	<i>Xestia agathina</i>	0.15 ± 0.13	Moorland
Ingrailed Clay (Noctuidae)	<i>Diarsia mendica</i>	2.36 ± 0.50	Open ground
July Highflyer (Geometridae)	<i>Hydriomena furcata</i>	1.54 ± 0.44	Generalist
Knotgrass (Noctuidae) ^a	<i>Acronicta rumicis</i>	0.03 ± 0.01	Wood generalist
Larch Pug (Geometridae)	<i>Eupithecia lariciata</i>	0.05 ± 0.03	Conifer
Large Emerald (Geometridae)	<i>Geometra papilionaria</i>	0.09 ± 0.04	Generalist
Large Yellow Underwing (Noctuidae)	<i>Noctua pronuba</i>	1.66 ± 1.01	Generalist
Latticed Heath (Geometridae) ^a	<i>Chiasmia clathrata</i>	0.01 ± 0.01	Generalist
Lempkes Gold Spot (Noctuidae)	<i>Plusia putnami</i>	0.14 ± 0.05	Generalist
Lesser Swallow Prominent (Notodontidae)	<i>Pheosia gnoma</i>	0.01 ± 0.01	Open ground
Lesser Yellow Underwing (Noctuidae)	<i>Noctua comes</i>	0.36 ± 0.15	Generalist
Light Emerald (Geometridae)	<i>Campaea margaritaria</i>	0.15 ± 0.07	Generalist
Map Winged Swift (Hepialidae)	<i>Korscheltellus fusconebulosa</i>	2.09 ± 0.39	Wood generalist
Marbled Minor (Noctuidae)	<i>Oligia strigilis</i>	0.11 ± 0.06	Generalist
Middle Barred Minor (Noctuidae)	<i>Oligia fasciuncula</i>	0.39 ± 0.14	Generalist
Mouse Moth (Noctuidae) ^a	<i>Amphipyra tragopoginis</i>	0.01 ± 0.01	Generalist

Muslin Footman (Arctiidae)	<i>Nudaria mundana</i>	0.09 ± 0.03	generalist
Narrow Winged Pug (Geometridae)	<i>Eupithecia nanata</i>	0.21 ± 0.09	Generalist
Neglected Rustic (Noctuidae) ^a	<i>Xestia castanea</i>	0.04 ± 0.02	Open ground
Northern Arches (Noctuidae)	<i>Apamea exulis</i>	0.91 ± 0.31	Open ground
Northern Spinach (Geometridae)	<i>Eulithis populata</i>	0.01 ± 0.01	Open ground
Pale Eggar (Lasiocampidae) ^a	<i>Trichiura crataegi</i>	0.02 ± 0.01	Generalist
Peach Blossom (Drepanidae)	<i>Thyatira batis</i>	0.04 ± 0.02	Generalist
Pebble Prominent (Notodontidae)	<i>Notodonta ziczac</i>	0.05 ± 0.02	Deciduous
Pine Carpet (Geometridae)	<i>Pennithera firmata</i>	0.01 ± 0.01	Deciduous
Pink Barred Sallow (Noctuidae)	<i>Xanthia togata</i>	0.04 ± 0.02	Conifer
Poplar Grey (Noctuidae)	<i>Subacronicta megacephala</i>	0.02 ± 0.01	Generalist
Poplar Hawk Moth (Sphingidae)	<i>Laothoe populi</i>	0.01 ± 0.01	Deciduous
Pretty Pinion (Geometridae)	<i>Perizoma blandiata</i>	0.29 ± 0.07	Generalist
Purple Bar (Geometridae)	<i>Cosmorhoe ocellata</i>	0.56 ± 0.15	Moorland
Purple Clay (Noctuidae)	<i>Diarsia brunnea</i>	0.09 ± 0.03	Open ground
Red Carpet (Geometridae) ^a	<i>Xanthorhoe decoloraria</i>	0.01 ± 0.01	Generalist
Red Twin Spot Carpet (Geometridae)	<i>Xanthorhoe spadicearia</i>	0.01 ± 0.01	Generalist
Riband Wave (Geometridae)	<i>Idaea aversata</i>	0.01 ± 0.01	Moorland
Rosy Minor (Noctuidae)	<i>Litoligia literosa</i>	0.01 ± 0.01	Generalist
Rustic (Noctuidae) ^a	<i>Hoplodrina blanda</i>	0.01 ± 0.01	Grassland
Sallow (Noctuidae) ^a	<i>Cirrhia icteritia</i>	0.01 ± 0.01	Generalist
Satyr Pug (Geometridae)	<i>Eupithecia satyrata</i>	0.01 ± 0.01	Moorland
Saxon (Noctuidae)	<i>Hyppa rectilinea</i>	0.28 ± 0.14	Generalist
Scalloped Hazel (Geometridae)	<i>Odontopera bidentata</i>	0.01 ± 0.01	Generalist
Scalloped Hooktip (Drepanidae)	<i>Falcaria lacertinaria</i>	0.04 ± 0.02	Wood generalist
Scalloped Oak (Geometridae)	<i>Crocallis elinguaris</i>	0.01 ± 0.01	Generalist
Scalloped Shell (Geometridae)	<i>Hydria undulata</i>	0.08 ± 0.04	Wood generalist
Scarce Silver Y (Noctuidae)	<i>Syngrapha interrogationis</i>	1.25 ± 0.43	Deciduous
Shoulder Striped Wainscot (Noctuidae) ^a	<i>Leucania comma</i>	0.01 ± 0.01	Moorland
Silver Ground Carpet (Geometridae)	<i>Xanthorhoe montanata</i>	0.01 ± 0.01	Generalist
Sixstriped Rustic (Noctuidae)	<i>Xestia sexstrigata</i>	0.17 ± 0.04	Generalist
Small Angleshades (Noctuidae)	<i>Euplexia lucipara</i>	0.01 ± 0.01	Generalist
Small Dotted Buff (Noctuidae)	<i>Photedes minima</i>	0.07 ± 0.03	Generalist
Small Fanfoot (Erebidae)	<i>Herminia grisealis</i>	0.09 ± 0.04	Generalist
Small Phoenix (Geometridae) ^a	<i>Ecliptopera silaceata</i>	0.07 ± 0.02	Deciduous
Small Rivulet (Geometridae)	<i>Perizoma alchemillata</i>	0.01 ± 0.01	Generalist
Small Square Spot (Noctuidae) ^a	<i>Diarsia rubi</i>	0.07 ± 0.04	Generalist
Small Wainscot (Noctuidae)	<i>Denticucullus pygmina</i>	0.21 ± 0.14	Generalist
Smokey Wainscot (Noctuidae)	<i>Mythimna impura</i>	0.01 ± 0.01	Generalist
Snout (Erebidae)	<i>Hypena proboscidalis</i>	0.08 ± 0.04	Generalist
Spruce Carpet (Geometridae)	<i>Thera britannica</i>	0.01 ± 0.01	Generalist
Square Spot Rustic (Noctuidae)	<i>Xestia xanthographa</i>	0.05 ± 0.03	Conifer
Square Spotted Clay (Noctuidae)	<i>Xestia stigmatica</i>	0.22 ± 0.10	Generalist
Straw Dot (Noctuidae)	<i>Rivula sericealis</i>	0.22 ± 0.11	Deciduous
Striped Twin Spot Carpet (Geometridae)	<i>Coenotephria salicata</i>	0.01 ± 0.01	Open ground

Swallow Prominent (Notodontidae)	<i>Pheosia tremula</i>	0.34 ± 0.10	Generalist
Tawny Barred Angle (Geometridae)	<i>Macaria liturata</i>	0.01 ± 0.01	Deciduous
The Clay (Noctuidae)	<i>Mythimna ferrago</i>	0.02 ± 0.02	Conifer
Treble Bar (Geometridae)	<i>Aplocera plagiata</i>	0.01 ± 0.01	Open ground
Triple Spotted Clay (Noctuidae)	<i>Xestia ditrapezium</i>	4.09 ± 0.82	Generalist
True Lovers Knot (Noctuidae)	<i>Lycophotia porphyrea</i>	0.07 ± 0.07	Deciduous
Twin Spot Carpet (Geometridae)	<i>Mesotype didymata</i>	0.01 ± 0.01	Moorland
Water Carpet (Geometridae)	<i>Lampropteryx suffumata</i>	0.04 ± 0.02	Open ground
Welsh Wave (Geometridae)	<i>Venusia cambrica</i>	0.05 ± 0.02	Generalist
White Ermine (Erebidae) ^a	<i>Spilosoma lubricipeda</i>	0.02 ± 0.01	Generalist
White Wave (Geometridae)	<i>Cabera pusaria</i>	4.07 ± 1.03	Generalist
Willow Beauty (Geometridae)	<i>Peribatodes rhomboidaria</i>	0.05 ± 0.03	Wood generalist
Wormwood Pug (Geometridae)	<i>Eupithecia absinthiata</i>	0.01 ± 0.01	Generalist

Table 2: List of micro moth species recorded:

Common name (Family)	Latin Name	Habitat preference	Abundance per trap (\pm SE)
Water Veneer (Crambidae)	<i>Acentria ephemerella</i>	Water	0.05 \pm 0.03
Caledonian Button (Tortricidae)	<i>Acleris caledoniana</i>	Moorland	0.01 \pm 0.01
Notched winged Tortricid (Tortricidae)	<i>Acleris emargana</i>	Deciduous	0.04 \pm 0.02
Dark-triangle Button (Tortricidae)	<i>Acleris laterana</i>	Open ground	0.01 \pm 0.01
Rhomboid Tortrix (Tortricidae)	<i>Acleris rhombana</i>	Generalist	0.01 \pm 0.01
Thistle Conch (Tortricidae)	<i>Aethes cnicana</i>	Grassland	0.04 \pm 0.03
Burdock Conch (Tortricidae)	<i>Aethes rubigana</i>	Open ground	0.01 \pm 0.01
Hook-marked Straw Moth (Tortricidae)	<i>Agapeta hamana</i>	Open ground	0.01 \pm 0.01
Hemlock Moth (Depressariidae)	<i>Agonopterix alstromeriana</i>	Open ground	0.01 \pm 0.01
Angelica Flat-body (Depressariidae)	<i>Agonopterix angelicella</i>	Generalist	0.01 \pm 0.01
Brindled Flat-body (Depressariidae)	<i>Agonopterix arenella</i>	Generalist	0.01 \pm 0.01
Gorse Tip Moth (Depressariidae)	<i>Agonopterix nervosa</i>	Generalist	0.02 \pm 0.01
Coastal Flat-body (Depressariidae)	<i>Agonopterix yeatiana</i>	Generalist	0.01 \pm 0.01
Barred Grass Veneer (Crambidae)	<i>Agriphila inquinatella</i>	Grassland	0.02 \pm 0.01
Pearl Veneer (Crambidae)	<i>Agriphila straminella</i>	Grassland	0.68 \pm 0.18
Common Grass Veneer (Crambidae)	<i>Agriphila tristella</i>	Grassland	0.01 \pm 0.01
Broken Barred Roller (Tortricidae)	<i>Ancylis unguicella</i>	Moorland	0.01 \pm 0.01
Birch Marble (Tortricidae)	<i>Apotomis betuletana</i>	Deciduous	0.05 \pm 0.03
Rush Marble (Tortricidae)	<i>Bactra lancealana</i>	Open ground	0.22 \pm 0.07
(Blastobasidae)	<i>Blastobasis decolorella</i>	Wood generalist	0.01 \pm 0.01
Dark Groundling (Gelechiidae)	<i>Bryotropha affinis</i>	Generalist	0.01 \pm 0.01
(Gelechiidae)	<i>Bryotropha boreella</i>	Generalist	0.01 \pm 0.01
Cinereous Groundling (Gelechiidae)	<i>Bryotropha terrella</i>	Grassland	0.01 \pm 0.01
Pearl-band Grass Veneer (Crambidae)	<i>Catoptria margaritella</i>	Moorland	0.21 \pm 0.10
Pearl Grass Veneer (Crambidae)	<i>Catoptria pinella</i>	Moorland	0.01 \pm 0.01
Dark Strawberry Tortrix (Tortricidae)	<i>Celypha lacunana</i>	Generalist	1.23 \pm 0.30
Garden Grass Veneer (Crambidae)	<i>Chrystoteuchia culmella</i>	Grassland	0.14 \pm 0.06
Flax Tortrix (Tortricidae)	<i>Cnephasia asseclana</i>	Generalist	0.01 \pm 0.01
Hedge Case-bearer (Coleophoridae)	<i>Coleophora striatipennella</i>	Deciduous	0.01 \pm 0.01
Hook-streaked Grass Veneer (Crambidae)	<i>Crambus lathoniellus</i>	Grassland	0.01 \pm 0.01
Grass Veneer (Crambidae)	<i>Crambus pascuella</i>	Grassland	0.31 \pm 0.09
Grey Gorse Piercer (Tortricidae)	<i>Cydia ulicetana</i>	Open ground	0.23 \pm 0.07
Northern Tubic (Oecophoridae)	<i>Denisia similella</i>	Deciduous	0.01 \pm 0.01
Little Grey (Crambidae)	<i>Dipleurina lacustrata</i>	Deciduous	0.01 \pm 0.01
(Crambidae)	<i>Donacaula micronellus</i>	Moorland	0.04 \pm 0.02
Dotted Shade (Tortricidae)	<i>Eana osseana</i>	Open ground	0.01 \pm 0.01
Brown China Mark (Crambidae)	<i>Elophila nymphaeata</i>	Water	0.01 \pm 0.01
Knapweed Bell (Tortricidae)	<i>Epiblema cirsiiana</i>	Deciduous	0.01 \pm 0.01
Thistle Bell (Tortricidae)	<i>Epiblema scutulana</i>	Open ground	0.01 \pm 0.01

Bramble Shoot Moth (Tortricidae)	<i>Epiblema uddmanniana</i>	Deciduous	0.01 ± 0.01
Square Barred Bell (Tortricidae)	<i>Epinotia fraternella</i>	Conifer	0.01 ± 0.01
Common Birch Bell (Tortricidae)	<i>Epinotia immundana</i>	Deciduous	0.16 ± 0.09
Grey Poplar Bell (Tortricidae)	<i>Epinotia nisella</i>	Deciduous	0.03 ± 0.03
Small Birch Bell (Tortricidae)	<i>Epinotia ramella</i>	Deciduous	0.01 ± 0.01
Variable Bell (Tortricidae)	<i>Epinotia solandriana</i>	Deciduous	0.01 ± 0.01
Common Spruce Bell (Tortricidae)	<i>Epinotia tedella</i>	Conifer	0.01 ± 0.01
White Blotch Bell (Tortricidae)	<i>Epinotia trigonella</i>	Wood generalist	0.11 ± 0.04
Bright Bell (Tortricidae)	<i>Eucosma hohenwartiana</i>	Grassland	0.02 ± 0.02
Two-coloured Bell (Tortricidae)	<i>Eucosma obumbratana</i>	Open ground	0.04 ± 0.02
Pied Grey (Crambidae)	<i>Eudonia delunella</i>	Deciduous	0.01 ± 0.01
Small Grey (Crambidae)	<i>Eudonia mercurella</i>	Deciduous	0.01 ± 0.01
Brassy Tortrix (Tortricidae)	<i>Eulia ministrana</i>	Deciduous	0.25 ± 0.16
Lilac Leafminer (Gracillariidae)	<i>Gracillaria syringella</i>	Generalist	0.01 ± 0.01
Small Fanfoot (Erebidae)	<i>Herminia grisealis</i>	Deciduous	0.02 ± 0.01
Marsh Oblique-barred (Erebidae)	<i>Hypenodes humidalis</i>	Open ground	0.99 ± 0.44
Red Piercer (Tortricidae)	<i>Lathronympha strigana</i>	Wood generalist	0.01 ± 0.01
Rust-blotch Cosmet (Momphidae)	<i>Mompha lacteella</i>	Grassland	0.01 ± 0.01
Little Cosmet (Momphidae)	<i>Mompha raschkiella</i>	Grassland	0.01 ± 0.01
Carrion Moth (tineidae)	<i>Monopis weaverella</i>	Generalist	0.01 ± 0.01
Heather Groundling (Gelechiidae)	<i>Neofaculta ericetella</i>	Moorland	0.01 ± 0.01
Beautiful China Mark (Crambidae)	<i>Nymphula stagnata</i>	Water	0.08 ± 0.03
Sorrel Bent-wing (Opostegidae)	<i>Opostega salaciella</i>	Grassland	0.01 ± 0.01
Woodland Marble (Tortricidae)	<i>Orthotaenia undulana</i>	Generalist	0.01 ± 0.01
Barred Fruit Tree Tortrix (Tortricidae)	<i>Pandemis cerasana</i>	Deciduous	0.04 ± 0.03
White-faced Tortrix (Tortricidae)	<i>Pandemis cinnamomeana</i>	Deciduous	0.01 ± 0.01
Dark Fruit Tree Tortrix (Tortricidae)	<i>Pandemis hepararia</i>	Deciduous	0.03 ± 0.01
Large Marble (Tortricidae)	<i>Phiaris schulziana</i>	Moorland	0.01 ± 0.01
Small Clouded Knot-horn (Pylalidae)	<i>Phycitodes saxicola</i>	Open ground	0.01 ± 0.01
Light Streak (Oecophoridae)	<i>Pleurota bicostella</i>	Moorland	0.01 ± 0.01
Diamondback Moth (Plutellidae)	<i>Plutella xylostella</i>	Generalist	0.01 ± 0.01
Ash Bud Moth (Praydicae)	<i>Prays fraxinella</i>	Deciduous	0.01 ± 0.01
White Plume Moth (Pterophoridae)	<i>Pterophorus pentadactyla</i>	Generalist	0.01 ± 0.01
Common Purple and Gold (Crambidae)	<i>Pyrausta purpuralis</i>	Grassland	0.01 ± 0.01
Holly Tortrix Moth (Tortricidae)	<i>Rhopobota naevana</i>	Generalist	0.04 ± 0.02
Pinion Streaked Snout (Hypenodinae)	<i>Schrankia costaestrigalis</i>	Open ground	0.57 ± 0.15
Common Grey (Crambidae)	<i>Scoparia ambigualis</i>	Deciduous	3.26 ± 0.62
Meadow Grey (Crambidae)	<i>Scoparia pyralella</i>	Open ground	0.05 ± 0.02
Brown Plume (Pterophoridae)	<i>Stenoptilia pterodactyla</i>	Generalist	0.01 ± 0.01
Fulvous Clothes Moth (tineidae)	<i>Tinea semifulvella</i>	Generalist	0.01 ± 0.01
Birds-nest Moth (tineidae)	<i>Tinea trinotella</i>	Generalist	0.01 ± 0.01
Pale Straw Pearl (Crambidae)	<i>Udea lutealis</i>	Generalist	0.17 ± 0.07
Olive Pearl (Crambidae)	<i>Udea olivalis</i>	Generalist	0.02 ± 0.01
Dusky Pearl (Crambidae)	<i>Udea prunalis</i>	Generalist	0.01 ± 0.01
Spindle Ermine (Yponomeutidae)	<i>Yponomeuta cagnagella</i>	Deciduous	0.01 ± 0.01
Bird-cherry Ermine (Yponomeutidae)	<i>Yponomeuta evonymella</i>	Deciduous	0.44 ± 0.43

White-shouldered Smudge (Ypsolophidae)	<i>Ypsolopha parenthesesella</i>	Deciduous	0.01 ± 0.01
Larch Tortrix (Tortricidae)	<i>Zeiraphera griseana</i>	Conifer	0.15 ± 0.12
Spruce Bud Moth (Tortricidae)	<i>Zeiraphera ratzeburgiana</i>	Conifer	0.17 ± 0.06

Supplementary data 4:

Stand characteristics for each management stage and stand features associated with management.

*Diameter at Breast Height – estimate of tree maturity

Stand Age	Management Stage	Key stand features
40 – 60 years	Mature	Occasionally thinned, stand density between 500 and 2200 stems ha ⁻¹ , average stand density: 1267 stems ha ⁻¹ , canopy closure between 80 and 100%, average closure 99%
20 – 40 years	Thin	Trees more densely packed, losing midstem branches and some trees dying off (self thinned). Occasionally thinned through management. Stand density between 600 – 2800 stems ha ⁻¹ , average stand density: 1624 stems ha ⁻¹ . Canopy closure between 50 and 100%, average closure: 95%
10 – 20 years	Thicket	Very dense, retain midstem branches, no undergrowth. Stand density between 300 – 3000 stems ha ⁻¹ , average stand density: 1850 stems ha ⁻¹ . Canopy closure between 16 and 100%, average closure: 69%
5 – 10 years	Young	Small, nearly all trees < 7cm DBH*, no canopy closure, lots of vegetation and ground cover
Clearfell	Felled < 5 years ago	Lots of dead wood and brash, standing water and undergrowth
Native	Unmanaged	Broadleaf stand, planted as part of plantation restructuring