

1 Title: Early avian functional assemblages after fire, clearcutting, and post-fire salvage logging in  
2 North American forests

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## 25 ABSTRACT

26 Increased demand for timber, the reduction in the available timber resources, more frequent and  
27 severe forest fires under a changing climate have increased the use of salvage logging in North  
28 American forests despite concerns regarding impacts on biodiversity and long-term forest  
29 productivity. We aimed to complement previous approaches that used bird species richness or  
30 individual abundance in salvage-logged habitats to assess the sustainability of this practice. We  
31 looked for commonalities in the taxonomic, functional and phylogenetic components of bird  
32 assemblages among these three post-disturbance habitats across a broad geographic range. We  
33 compiled six North American datasets selected from primary and grey literature that documented  
34 species composition of avian assemblages in habitats after recent fire, post-fire salvage logging  
35 and traditional logging. Our results revealed contrasting patterns of bird trait assemblage among  
36 burned, post-fire salvage and traditionally logged habitats. In salvage-logged habitats, taxonomic  
37 diversity, functional diversity, functional and phylogenetic redundancy were significantly lower  
38 than in both burned and traditionally logged habitats. The frequency of insectivores was  
39 significantly lower after salvage logging than after both fire and traditional logging. These findings  
40 suggest that cumulative disturbances have a negative effect on early assembly of bird communities.  
41 The outcomes of this study encourage further assessments, at landscape level, of salvage logging  
42 intensity, burn size and fire severity on bird functional structure to better plan for their  
43 conservation.

44 **Keywords:** Phylogenetic redundancy, Forest disturbance, Functional redundancy, Taxonomic  
45 diversity, Timber supply

## 46 INTRODUCTION

47 Post-fire salvage logging is becoming more prevalent worldwide (Lindenmayer et al. 2008; Müller  
48 et al. 2019). It may become even more so if forecasted increases in the frequency and severity of  
49 natural disturbances under climate change scenarios hold true (Price et al. 2013) and burned forests  
50 become more accessible with the expansion of road networks and infrastructure. Minimizing the  
51 ecological effects of salvage logging while maintaining its economic benefits is challenging  
52 (Lindenmayer and Noss 2006; McIver and Starr 2000). A potentially significant factor associated  
53 with salvage logging is the cumulative effect of following natural (e.g. fire) disturbance with  
54 anthropogenic (e.g. clear cutting) disturbances (Lindenmayer et al. 2017) on ecosystem  
55 composition (i.e. biotic and abiotic components) and function (i.e. collective biotic activity of  
56 plants, animals, and microbes and their effects on the physical and chemical conditions of their  
57 habitat (Naeem 1998). After fire, forests are dominated by high snag densities in addition to  
58 patches of living trees of varying sizes according to fire severity (Lindenmayer et al. 2008; Nappi  
59 et al. 2004). By removing snags, post fire salvage logging modifies habitat structure (Lewis and  
60 Hartley 2006; Lindenmayer and Noss 2006) and reduces resource availability for species feeding  
61 and/or nesting in dead wood. After traditional logging, the majority of merchantable trees are  
62 removed and the harvested stands are dominated by shrubs, herbs, regenerating trees and some  
63 canopy trees within block residual retention (Swanson et al. 2011).

64 Several biodiversity indicators (e.g. species occurrence, species richness) have previously been  
65 used to assess the ecological impact of major forest disturbances (Lindenmayer and Noss 2006).  
66 The immediate stand-scale effect of salvage logging on biodiversity has received a lot of attention  
67 in the literature (Cahall and Hayes 2009; De Bello et al. 2007; Lindenmayer and Noss 2006; Nappi  
68 et al. 2004), with birds being one of the species groups having seen the most attention. Bird

69 communities play many roles in ecosystem function (Mikusiński et al. 2018), including regulating  
70 predator populations, controlling populations of their prey (Van Bael et al. 2003) and enhancing  
71 plant reproduction via pollination or seed dispersal (Anderson et al. 2011). Some birds act as  
72 keystone species (Drever and Martin 2010; Hutto and Gallo 2006; Koivula and Schmiegelow  
73 2007; Nappi et al. 2010). For example, woodpeckers generate habitat (i.e. excavation of cavities  
74 in trees) for secondary cavity users that can facilitate post-fire regeneration through seed dispersal,  
75 seed germination and regulation of insect populations, which in turn affect vegetative growth  
76 (Gregory et al. 2005; Vandewalle et al. 2010; Venier and Pearce 2004). Since bird populations  
77 show relatively rapid responses to environmental change, they represent sensitive indicators of  
78 habitat conditions (Gregory et al. 2005; Vandewalle et al. 2010; Venier and Pearce 2004).

79 Previous work on the response of birds to salvage logging has primarily focussed on cavity-nesting  
80 birds, which are thought to be particularly sensitive to post-fire logging because unlogged post-  
81 fire habitat provides cavity nesting habitat (Drever and Martin 2010; Hutto and Gallo 2006;  
82 Koivula and Schmiegelow 2007; Nappi et al. 2010). Aside from work on cavity nesters, most bird  
83 studies have focused on taxonomic diversity (i.e. species richness or abundance) (Cahall and Hayes  
84 2009; Morissette et al. 2002). Examining the impact of salvage logging on taxonomic diversity  
85 has the advantages of being sensitive to short-term dynamics and easily interpretable (Saint-  
86 Germain and Greene 2009). Unfortunately, the response of taxonomic diversity to a disturbance  
87 can rarely be generalized to other regions. It may not provide insight into the interaction between  
88 birds and their environment (Klingbeil and Willig 2016a; Lindenmayer and Noss 2006;  
89 Vandewalle et al. 2010) and other regions (Hobson and Schieck 1999). Examining the response of  
90 trait-based functional groups (as opposed to individual species) could complement single-  
91 parameter approaches (e.g. species richness) by providing possible mechanistic insights not

92 revealed through analyses of taxonomic diversity (Lindenmayer et al. 2014; Mouillot et al. 2013;  
93 Tarbill et al. 2015; Vandewalle et al. 2010). Functional traits are presumed to either have an effect  
94 on ecosystem function (e.g. seed dispersal, structural transformation, pest regulation) or  
95 characterize a response to environmental change (e.g. resource availability and disturbance)  
96 (Lavorel and Garnier 2002; Luck et al. 2012). For instance, foraging and nesting traits are related  
97 to resource use (e.g. insects, seeds) and changes in ecosystem structure (e.g. modification of soil  
98 and tree structure) and thereby provide habitat to other organisms. After a forest fire, it is well  
99 established that the activity of cavity excavating birds have cascading effects on forest succession  
100 (Tarbill et al. 2015). Less is known about whether other functional groups and interactions occur  
101 in early post-disturbance forests that may have ecological implications.

102 Bird communities can be described using many competing and/or complementary indices such as  
103 taxonomic diversity, functional diversity considering trait distance (FD, i.e. range in the number  
104 of ecological functions represented in a community) or functional diversity considering  
105 phylogenetic distance (PFD) or functional redundancy (FR, i.e. the number of species playing the  
106 same functional role) or phylogenetic redundancy (PFR). Although, high FD is often linked to  
107 greater resilience (Elmqvist et al. 2003; Ricotta et al. 2016), it has been predicted that conserving  
108 high FR should buffer communities following disturbance via species compensation (i.e.  
109 remaining species playing similar ecological roles as those lost from disturbance) (Concepción et  
110 al. 2016; Laliberté et al. 2010; Ricotta et al. 2016). Functional diversity and redundancy have a  
111 more direct relationship with community resilience to environmental change than taxonomic  
112 diversity (Mayfield et al. 2010; Petchey et al. 2007; Ricotta et al. 2016) and functional and  
113 taxonomic diversity may differentially respond to disturbance (Laliberte et al. 2010). Ultimately,  
114 examining the response of both taxonomic and functional diversity may provide more robust

115 measures of the potential implications of disturbance on bird communities' resilience to  
116 disturbances than taxonomic diversity alone (Klingbeil and Willig 2016b; Lavorel and Garnier  
117 2002; Ricotta et al. 2016).

118 In this study, we use a set of taxonomic diversity, functional diversity, functional and phylogenetic  
119 redundancy indices to test two hypotheses regarding the impact of disturbance on forest bird  
120 communities. We hypothesized that: 1) assemblages based upon the ecological traits of birds  
121 would differ among burned, traditionally logged and salvage-logged habitats due to the strong link  
122 between bird composition and habitat structure (Schieck and Song 2006); 2) salvage logging after  
123 fire should reduce taxonomic diversity, functional diversity, functional and phylogenetic  
124 redundancy of bird communities because of the cumulative effects of both disturbances  
125 (Lindenmayer and Noss 2006). To assess these predictions, we used legacy datasets of bird species  
126 composition and abundance in burned non-salvage, post-fire salvage and traditional logged North  
127 American forests, derived from peer-reviewed and grey literature.

## 128 **METHODS**

### 129 **Literature search**

130 Our literature survey included a search between September and November 2014 in ISI Web of  
131 Science and Google Scholar using the combination of different key words related to avian  
132 community in North America boreal forests and species composition changes after fire, post-fire  
133 salvage logging and traditional logging. We used the following search terms; ("boreal forest" OR  
134 "conifer forest" AND "salvage logging" OR "salvage cutting" OR "salvage harvesting" OR  
135 "commercial logging" OR "commercial cutting" OR "conventional harvesting" OR "traditional  
136 harvesting" OR harvesting OR logging OR fire OR wildfire AND bird\* OR "species abundance"

137 OR “functional diversity”). We also obtained relevant papers, theses and reports listed in the  
138 references of primary and grey literature from the aforementioned literature search. We only  
139 selected studies that explicitly considered changes in bird community within 5 years after  
140 disturbances (fire, post-fire salvage logging, and traditional logging) in Canadian forests. Studies  
141 targeting specific bird groups (such as cavity nesters, woodpeckers) or a limited number of species  
142 from the final data compilation were excluded to avoid biased comparisons.

### 143 **Data compilation**

144 We compiled six North American datasets selected from primary (peer-reviewed) and grey  
145 literature that documented species composition of avian assemblages in coniferous, deciduous and  
146 mixed wood forests after recent ( $\leq 5$  years) fire, post-fire salvage logging and traditional logging  
147 (Fig. 1; Table S1). We obtained raw data from three studies and extracted data from three published  
148 studies that provided a complete list of species and abundances. When a study had missing  
149 information that was important to consider in the study such as species abundance or occurrence,  
150 we contacted the authors for additional information. We excluded any study for which the species  
151 abundance or occurrence data were unavailable within the study. For these latter studies, we  
152 extracted the mean frequency of occurrence or species abundance for each site where point count  
153 sampling had been applied. We pooled data from stand types within the same treatment nested  
154 within individual study and treated these as our study sites. For the three studies in which we had  
155 access to the raw data, we calculated mean frequency of occurrence or abundance of species per  
156 treatment within each stand from raw data (Azeria et al. 2011; Song and Hannah unpublished; Van  
157 Wilgenburg and Hobson 2008), to make them comparable to estimates extracted from the four  
158 published studies. To ensure comparability between datasets surveyed using different methods (see  
159 Table S1), we calculated the relative contribution of each species to its community ( $p_i$ ). The data



160 table  $Y = [y_{ij}]$  presents sites in rows and species occurrence or abundance in columns. Each species  
161 mean frequency of occurrence or abundance ( $x_i$ ) was divided by the row sums  $y_{i+}$  following  
162 Legendre and Gallagher (Legendre and Gallagher 2001):  $p_i = x_i/y_{i+}$ . We had 17 sites in total  
163 distributed between the three treatments: fire (with no salvage logging or less than 33% area  
164 salvage-logged;  $n=6$ ), salvage logging (i.e., more than 33% area salvage-logged after fire;  $n=7$ )  
165 and traditional logging (i.e., clear-cut forest;  $n=4$ ).

### 166 **Bird trait selection**

167 We selected seven species traits which are potentially sensitive to forest disturbances (Luck et al.  
168 2013; Vandewalle et al. 2010). The selected traits are mainly related foraging and nesting  
169 behaviours, resource acquisition (foraging guild), mobility (body mass), reproduction (clutch size,  
170 nesting behaviour) and ecosystem engineering or structural transformation (nest location and  
171 guild). More details on potential response to environment and effect on ecosystem function of  
172 selected traits are reported in Table 1. We derived trait data from species accounts in the Birds of  
173 North America (Poole 2005) and phylogeny data in BirdTree.org (Jetz et al. 2014).

### 174 **Data analysis**

175 To assess community trait variation among disturbances, the relationship between bird species  
176 traits and disturbance type (traditional logging, fire, salvage logging, undisturbed) was tested using  
177 RLQ analysis (Dray et al. 2014). RLQ analysis is a three-matrix ordination method that relates  
178 species abundance and traits to environmental factors. The three matrices are: (1) L (community  
179 data) with species as columns and sampling units as rows; (2) Q (traits data) with traits as columns  
180 and species as rows; and (3) R (environmental data) with treatments (disturbance) as columns and  
181 sampling unit as rows. Prior to the analysis of trait data, we standardized body mass and clutch

182 size (using Z-score scaling) to reduce the high variation in magnitude of species body mass and  
183 clutch size. We use the *rlq* function implemented in the “ade4” package version 1.7-15 (Bougeard  
184 and Dray 2018; Dray and Dufour 2007) of R the statistical and programming environment version  
185 3.6.3 (R Core Team 2020) to perform RLQ analysis. We tested the significance of the relationship  
186 between treatments (R) and species functional traits (Q) using a “model 6” (Dray et al. 2014)  
187 Monte Carlo permutation test with 9999 permutations. Monte Carlo permutation tests were  
188 performed using the function *randtest* implemented in the “ade4” package version 1.7-15.  
189 Permutation “model 6” combines permutation “model 2” (i.e. permute entire rows of matrix L to  
190 test the null hypothesis that the distribution of species with fixed traits is not influenced by the  
191 environmental conditions) and “model 4” (i.e. permute entire columns of matrix L to test that the  
192 species composition of the sampling units is not influenced by the species traits) (Dray et al. 2014).

193 We used a set of complementary indices to describe the bird communities. (1) Taxonomic diversity  
194 (SD) is a measure of diversity that considers species relative abundance. 2) Functional diversity  
195 (FD) is the range in the number of ecological functions represented in a community considering  
196 trait distance (i.e. trait divergence between species) and 3) PFD when considering phylogenetic  
197 distance. 4) Functional redundancy (FR) is the number of species with similar functional roles  
198 considering trait distance and 5) phylogenetic redundancy (PFR) when considering phylogenetic  
199 distance. We used *rao.diversity* function implemented in “SYNCSA” package version 1.3.4  
200 (Debastiani and Pillar 2012) to compute the indices. We calculated community taxonomic  
201 diversity within each treatment using the Gini-Simpson index (Simpson 1949). The rationale  
202 behind the use of Gini-Simpson index (SD) is its direct relationship with the functional diversity  
203 (FD) and functional redundancy (FR) indices (i.e.  $FR = SD - FD$ ) (De Bello et al. 2007). We  
204 calculated FD and PFD using Rao’s quadratic entropy index (Rao 1982). Functional diversity (FD)

205 and FR were computed separately for each of the seven selected traits and for the overall trait  
206 assemblages by treatment, while SD, PFD and PFR were computed overall trait and phylogenetic  
207 assemblage. The values of the metrics (SD, FD, PFD, FR and PFR) range between 0 (no diversity)  
208 and 1 (infinite diversity).

209 The statistical analyses of the diversity and redundancy indices followed a protocol for data  
210 exploration described by Zuur et al. (2010), and protocols for conducting and presenting  
211 regression-type analyses (Stasinopoulos et al. 2017; Zuur and Ieno 2016; Zuur et al. 2010). We  
212 investigated what theoretical distribution came closest to the marginal distribution of the  
213 dependent variable (i.e. SD, FD, PFD, FR and PFR). We used the *fitDist* function of gamlss  
214 package version 5.1-4 (Rigby and Stasinopoulos 2005) in order to find the best fit (based on the  
215 Akaike's Information Criterion (AIC) of the different fitting attempts) among a list of distributions  
216 adapted to positive real number (ranged between 0 and 1) data available in the gamlss.dist package  
217 version 5.1-4 (Stasinopoulos and Rigby 2019). The simplex distribution had the best fit based on  
218 AIC for SD, FD and PFD, while beta distribution has the best fit for FR and PFD (Table S2).

219 Since the data were from different locations and stand types, we therefore applied mixed effect  
220 models with the combination of Location (4 levels) and Stand\_type (3 levels) being treated as a  
221 random effect. To model the community diversity indices as function of disturbance, we used a  
222 Generalized Additive Model for Location, Scale and Shape (GAMLSS) with the best-fit family  
223 distribution (Table S2) mentioned above for each response variable. GAMLSS family distributions  
224 have up to four parameters ( $\mu$ ,  $\sigma$ ,  $\nu$ ,  $\tau$ ). The terms location, scale, and shape refer to these parameters  
225 and are connected, but necessarily equal, to the four moments of a distribution, namely the mean  
226 ( $\mu$ ), the variance ( $\sigma$ ), the skewness ( $\nu$ ), and the kurtosis ( $\tau$ ). GAMLSS is a modern distribution-  
227 based approach to regression analysis that expands traditional approaches to accommodate

228 distribution parameters that are modeled as additive functions of predictor variables (Barber 2018).  
229 We included disturbance type as a fixed effect factor in our models. Model assumptions were  
230 verified by plotting residuals against fitted values to assess homoscedasticity; making QQ plots  
231 and computing filliben correlation coefficients to verify normality and using worm plots to  
232 visualize model fit to the data (Stasinopoulos et al. 2017). We performed likelihood-ratio tests to  
233 assess significance of the fixed and random factors.

234 We investigated the functional rarity within communities between disturbance types. Rare species  
235 perform different functions in ecosystems, some being redundant with those of many other rare  
236 and common species, while others are unique (Violle et al. 2017). We computed the functional  
237 rarity (Uniqueness) and taxonomic scarcity (Scarcity) indices. Uniqueness (UI) measures how  
238 functionally rare a species is, with values ranging from 0 indicating that the focal species shares  
239 the exact same traits as other species in the pool and conversely to 1 indicating the species shares  
240 no traits as others in the pool. Scarcity (SI) measures how abundant a species is in comparison to  
241 other species within a community with values ranging from 0 indicating that the focal species is  
242 abundant to 1 indicating the opposite. We used the “funrar” package version 1.2.1 (Grenié et al.  
243 2017) to compute the functional uniqueness and the taxonomic scarcity indices.

244 All graphics were produced using the “ggplot2” package version 3.3.0 (Wickham 2016).

245

## 246 **RESULTS**

### 247 **Bird traits response to disturbances**

248 The first axis of the RLQ analysis captured 78% of the total association between traits and  
249 disturbance classes (i.e. co-inertia), accounting for most of their relation. The Monte Carlo  
250 permutation “model2” test was significant ( $P=0.048$ ) indicating that trait assemblages were  
251 significantly associated to disturbances. In addition, the Monte Carlo permutation “model 4” was  
252 not significant ( $P=0.172$ ) indicating that species composition of the sampling units was not  
253 significantly influenced by species characteristics. The summary statistics showed that the axis 1  
254 describes 40% of the covariance between disturbances and traits. The RLQ axis 1 split the bird  
255 trait assemblage along a gradient from naturally disturbed (positive side of axis 1) to  
256 anthropogenically disturbed forest types (negative side of axis 1, Figure 2), with logged forests  
257 falling between non-salvaged and salvaged habitats. For example, there were cavity nesters on the  
258 positive side vs non-cavity nesters on the negative side of axis 1; tree nesters vs shrub and ground  
259 nesters; omnivores and insectivores vs herbivores; bark foragers, foliage gleaners and aerial  
260 foragers vs ground foragers. Axis 2 of the RLQ analysis captured 22% of the association between  
261 traits and disturbances and described 20% of covariance between them. The second axis  
262 corresponded to a gradient of trait assemblage between traditionally logged and both non-salvaged  
263 and salvaged forests. Carnivores, aerial and ground nesters were associated with traditionally  
264 logged landscapes (i.e. fell on negative side of the axis 2), while herbivores and shrub nesters  
265 more associated with salvage logged landscape (i.e. falling on negative side of axis 2).

266 Bark foragers that are mainly omnivores or insectivores and tree cavity nesters were the most  
267 frequent in burned and non-salvage habitat assemblages. Species associated with salvage-logged  
268 habitats were predominantly ground foragers that are herbivores and shrub nesters. The analysis  
269 also associated suggested body mass was positively correlated with salvage logging. Species using  
270 open habitats, ground nesters and carnivores characterized harvested habitats.

## 271 **Birds' community metric response to forest disturbances**

272 Bird taxonomic diversity (SD) differed substantially among disturbances (Table 2A). The  
273 validation of the model describing taxonomic diversity indicates no violation of assumptions (i.e.  
274 independence and absence of residual patterns, Figure S1), and the worm plots indicates an  
275 appropriate model fit (i.e. 95% of the worm plot lying within the dashed confidence bands). The  
276 taxonomic diversity in burned habitats was significantly lower compared to traditionally logged  
277 habitats (0.89 vs 0.93), while it was significantly higher (0.89 vs 0.77) in burned habitats compared  
278 to salvage-logged habitats (Figure 3.A, Table S3A). The effect of disturbance on taxonomic  
279 diversity was largely masked by high variation among measurements (Table S3A; Figure 3B). The  
280 highest coefficient of variation ( $\sigma$ ) was observed in salvage-logged habitats (2.84) while there were  
281 similar in burned and traditionally logged habitats (0.92 and 0.93 respectively).

282 Global functional diversity (FD, univariate index) considering trait distance, did not statistically  
283 differ among disturbance types (Table 2B; Figure 3C; Table S3B). However, the coefficient of  
284 variation was significantly higher in salvage-logged habitats compared to the other disturbances  
285 (Table 2B, Figure 3D). Analysis of individual traits showed that foraging strategy FD was  
286 significantly lower in salvage-logged habitats than in both burned or traditionally logged (0.51 vs  
287 0.53) habitats (Table 3A). Additionally, nesting guild FD in burned habitats was significantly  
288 higher than in salvage-logged habitats (0.31 vs 0.24), while the latter was significantly higher than  
289 in traditionally logged ones (0.24 vs 0.19). Functional diversity, considering phylogenetic distance,  
290 was significantly higher in burned habitats than in both harvested (0.01 vs 0.00) and salvage-  
291 logged (0.01 vs 0.00) habitats (Table 2C; Figure 3E, Table S3C) and the coefficient of variation  
292 followed the same pattern (Figure 3F). Model validations suggested good fit (Figure S2 & S3).

293 Global functional redundancy (FR, Univariate index) in burned habitats was significantly lower  
294 than in traditionally logged habitats (0.37 vs 0.33), while it was significantly higher in harvested  
295 habitats (0.40) than in burned ones (Table 2D; Figure 3G, Table S3D). The coefficient of variation  
296 exhibited converse pattern (Figure 3H.). The phylogenetic redundancy of the bird community  
297 followed the same pattern of the global functional redundancy (Table 2E; Figure 3I & J, Table  
298 S3E). Analysis of individual traits showed that functional redundancy in salvage-logged habitats  
299 was significantly lower than in burned habitats (e.g. foraging guild, habitat affinity, nest location  
300 and body mass) and in traditionally logged habitats (clutch size; Table 3B). Only foraging strategy  
301 and nest guild FR did not exhibit significant differences among disturbances. Model validation  
302 suggested the data were well fit by the models (Figure S4 & S5).

303 The functional uniqueness analysis revealed that insectivores with higher uniqueness values (i.e.  
304 high functional distance from other species) in salvage-logged habitats were *Cardellina pusilla*,  
305 *Catharus ustulatus*, *Certhia americana*, *Empidonax flaviventris*, *Empidonax minimus*, *Empidonax*  
306 *traillii*, *Mniotilta varia*, *Poecile atricapillus*, *Troglodytes aedon* and *Troglodytes hiemalis* (Table  
307 4; Table A1). The taxonomic scarcity analysis show that at least 12 insectivores (*Chaetura vauxi*,  
308 *Contopus cooperi*, *Ixoreus naevius*, *Pheucticus melanocephalus*, *Poecile gambeli*, *Setophaga*  
309 *townsendi*, *Sialia currucoides*, *Sialia mexicana*, *Sphyrapicus nuchalis*, *Sphyrapicus thyroideus*,  
310 *Sturnus vulgaris* and *Vireo gilvus*) were abundant (SI<0.01) in non-salvage habitats, while in  
311 salvage-logged habitats these species were scarce (SI>0.80) or not represented. Two species  
312 (*Bucephala albeola* and *Setophaga fusca*) were scarce (SI>0.90) in non-salvage habitats, but they  
313 were absent in salvage-logged habitats.

314

## 315 **DISCUSSION**

316 Consistent with our first hypothesis, our study revealed contrasting patterns of bird community  
317 composition among burned and non-salvage, post-fire salvage and traditionally logged habitats.  
318 The RLQ results showed that species displaying specific ecological traits are differentially  
319 associated with natural and anthropogenic disturbances, with traditional logging exhibiting  
320 intermediate effects between fire and salvage logging on bird species differing in their nest location  
321 and habitat affinities. This contrast is mainly due to the differences in the remaining configuration  
322 of habitat (e.g. canopy cover, snag density) left by each disturbance (Thorn et al. 2016). Forest fire  
323 generates “keystone structures” (i.e. distinct spatial structures providing resources, shelter or goods  
324 and services that are crucial for other species (Tews et al. 2004)) for cavity nester and insectivores.  
325 These structures are characterised by stand structural complexity, habitats connectivity and  
326 landscape heterogeneity (Burton et al. 2016; Lindenmayer et al. 2006). However, salvage-logging  
327 after fire reduces these keystone structures quantity and/or quality (Lindenmayer et al. 2008)  
328 favoring ground and shrub nesters.

329 A meta-analysis comparing changes in boreal forest bird assembly after fire and harvesting  
330 reported that post-fire communities still differed from post-harvest ones 30 years after disturbance  
331 (Schieck and Song 2006). Convergence between communities started to be observed around 60  
332 years after disturbance (Schieck and Hobson 2000). To our knowledge, no studies have yet  
333 examined convergence between fire and salvage-logged stands. The reduction of keystone  
334 structures in salvaged habitats and the attendant impacts on insectivores (i.e. loss of perches,  
335 potential nest sites and resource upon which insects forage) could exacerbate the widespread  
336 declines that North American insectivores are experiencing (Kelly et al. 2013; Nebel et al. 2010;  
337 Sauer et al. 2017) if salvage logging becomes more widely used. While the long-term



338 consequences of declining insectivore populations on forest ecosystem health are unclear, it may  
339 reduce the extent of natural pest control (e.g. some bark beetles or long horn beetles) (Dolbeer  
340 2008; Naylor and Ehrlich 1997).

341 Our results also showed that salvage-logged habitats had the lowest taxonomic diversity, global  
342 functional diversity, functional and phylogenetic redundancy among all the disturbances  
343 considered. This is consistent with our expectations that salvage logging would reduce avian  
344 taxonomic diversity, functional diversity, functional and phylogenetic redundancy after fire. Thorn  
345 et al. (2017) found that bird taxonomic diversity was significantly lower in salvage-logged habitat  
346 than in non-salvage habitats. Our results also revealed a decrease in bird functional diversity after  
347 salvage logging. For instance, individual traits such as foraging strategy and nest guild had lower  
348 functional diversity in salvage-logged habitats than in burned and non-salvaged habitats. This is  
349 consistent with previous studies (Cahall and Hayes 2009; Hutto et al. 2015; Nappi and Drapeau  
350 2011) that reported negative effect of salvage logging on cavity nesters, bark and aerial foragers  
351 due of the reduction of the keystone structures.

352 We observed that functional and phylogenetic redundancy of bird communities was lowest in  
353 salvage-logged habitats, suggesting that many species in these habitats are functionally and  
354 phylogenetically unique compared to both burned and harvested habitats. In fact, post fire habitats  
355 have high structural complexity (i.e. spatial heterogeneity) that results in greater representation of  
356 species displaying ecological traits such as insectivory and cavity nesting (Hutto et al. 2015;  
357 Morissette et al. 2002; Nappi et al. 2010). In contrast, traditionally (green) harvested habitats  
358 displayed patterns of community composition that were intermediate between fire and salvage  
359 logging with respect to the ecological traits of the constituent species.

360 The under scarcity of insectivores and cavity nesters species (e.g. *Bucephala albeola*, *Certhia*  
361 *americana*, *Chaetura vauxi*, *Contopus cooperi*, *Empidonax flaviventris*, *Empidonax minimus*,  
362 *Ixoreus naevius*, *Pheucticus melanocephalus*, *Pica hudsonia*, *Poecile atricapillus*, *Poecile*  
363 *gambeli*, *Poecile rufescens*, *Setophaga fusca*, *Setophaga townsendi*, *Sialia mexicana*, *Sphyrapicus*  
364 *nuchalis*, *Sphyrapicus thyroideus*, *Sphyrapicus vulgaris*, *Troglodytes aedon* and *Troglodytes*  
365 *hiemalis*) in salvage-logged habitats compared to post fire ones, suggests a loss of some functional  
366 roles after salvage-logging. Salvage logging disrupts the specific functional roles of some bird  
367 species and their associated partners post-fire (e.g. marked reduction of insect population control  
368 or marked reduction in seed dispersal). The disruption of these specific functional roles of some  
369 birds depends on both fire severity and salvage logging intensity, since salvage logging targets the  
370 least burned forests (Koivula and Spence 2006). The least burned habitats can function as refugia  
371 for species associated with unburned habitats. If most of the burned forests are salvage-logged, it  
372 could have a negative impact on the conservation of some bird species.

373 Limiting or reducing logging intensity to keep a suitable residual structure of fire legacy connected  
374 to green forest might help to mitigate salvage logging impact. As some ecological traits are  
375 positively associated with increased fire severity, the retention of the most severely burned stands  
376 which have reduced timber value may preserve components of the bird community and help  
377 maintain ecological functions performed by birds on the landscape (Koivula and Schmiegelow  
378 2007; Smucker et al. 2005). However, the most severely burned stands are also the least colonized  
379 by saproxylic insects, which are the food for woodpeckers (Saint-Germain et al. 2004; Saint-  
380 Germain et al. 2007). These stands could be useful for at least one idiosyncratic longhorned beetle  
381 (*Gnathacmeops pratensis*) but they would not provide diverse and abundant food for bark foragers.  
382 Some authors suggested keeping the periphery of burned stands to make sure ecotones are

383 maintained (Boucher et al. 2016; Nappi and Drapeau 2011). Maintaining ecotones along the burn  
384 periphery could facilitate species colonizing burned stands from adjacent green forest as it would  
385 reduce dispersal distances for birds and insects, many of which use both habitats. Since any single  
386 disturbance will disadvantage species not possessing traits adapted to the ecological conditions  
387 created by that disturbance, the precautionary approach would be to vary the level of retention  
388 between salvage logged stands (Bunnell and Houde 2010) and to maximize the connectivity  
389 between residual stands (Boucher et al. 2016; Nappi and Drapeau 2011).

390 Although we recommend limiting post fire salvage logging to conserve the functional roles played  
391 by birds, data on how trait assemblage patterns vary with fire size and severity are lacking and  
392 needed. Fire severity and size are known to have a strong effect on bird communities as well as  
393 the insects on which they feed (Boucher et al. 2016; Luck et al. 2012; Stephens et al. 2015).  
394 Understanding the relationship between bird community dynamics and fire size and severity would  
395 lead to the identification of high conservation value forests for burn-associated birds and would  
396 improve retention strategies for salvage logging operations. In conclusion, we show that in North  
397 American forests, ecological trait composition of bird communities varies between post  
398 disturbance habitats. Importantly, salvage logging significantly reduces bird taxonomic diversity,  
399 functional diversity, and both functional and phylogenetic redundancy after fire. Although we  
400 believe that we can broadly generalize our findings to other North American forests, studies are  
401 needed to test the generality of this finding in other regions in the world and to determine the  
402 variation around this estimate.

403

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### LITERATURE CITED

- 412 Anderson, S.H., Kelly, D., Ladley, J.J., Molloy, S., and Terry, J. 2011. Cascading effects of bird  
 413 functional extinction reduce pollination and plant density [Article]. *Science* **331**(6020):  
 414 1068-1071. doi:10.1126/science.1199092.
- 415 Azeria, E.T., Ibarzabal, J., Hébert, C., Boucher, J., Imbeau, L., and Savard, J.-P.L. 2011.  
 416 Differential response of bird functional traits to post-fire salvage logging in a boreal  
 417 forest ecosystem. *Acta Oecologica* **37**(3): 220-229. doi:10.1016/j.actao.2011.02.005.
- 418 Barber, X. 2018. Flexible Regression and Smoothing: Using GAMLSS in R. *Journal of statistical*  
 419 *software* **85**. doi:10.18637/jss.v085.b02.
- 420 Boucher, J., Hebert, C., Ibarzabal, J., Bauce, E.R., Leather, S., and Jonsell, M. 2016. High  
 421 conservation value forests for burn-associated saproxylic beetles: an approach for  
 422 developing sustainable post-fire salvage logging in boreal forest. *Insect Conservation and*  
 423 *Diversity* **9**(5): 402-415.
- 424 Bougeard, S. and Dray, S. 2018. Supervised Multiblock Analysis in R with the ade4 Package.  
 425 *Journal of Statistical Software* **86**(1): 1-17. doi:10.18637/jss.v086.i01.
- 426 Bunnell, F.L. and Houde, I. 2010. Down wood and biodiversity — implications to forest  
 427 practices [Review]. *Environmental Reviews* **18**(NA): 397-421. doi:10.1139/a10-019.
- 428 Burton, P.J., Messier, C., Adamowicz, W.L., and Kuuluvainen, T. 2016. Sustainable  
 429 management of Canada's boreal forests: Progress and prospects. *Écoscience* **13**(2): 234-  
 430 248. doi:10.2980/i1195-6860-13-2-234.1.
- 431 Brandt 2009. The extent of the North American boreal zone. *Environ. Rev.* **17**: 101–161
- 432 Cahall, R.E. and Hayes, J.P. 2009. Influences of postfire salvage logging on forest birds in the  
 433 Eastern Cascades, Oregon, USA. *Forest Ecology and Management* **257**(3): 1119-1128.  
 434 Available from <Go to ISI>://CCC:000263215700040 [accessed].
- 435 Concepción, E.D., Götzenberger, L., Nobis, M.P., de Bello, F., Obrist, M.K., and Moretti, M.  
 436 2017. Contrasting trait assembly patterns in plant and bird communities along  
 437 environmental and human-induced land-use gradients. *Ecography* **40**(6): 753-763.  
 438 doi:10.1111/ecog.02121.
- 439 De Bello, F., Lepš, J., Lavorel, S., and Moretti, M. 2007. Importance of species abundance for  
 440 assessment of trait composition: an example based on pollinator communities [Article].  
 441 *Community Ecology* **8**(2): 163-170. doi:10.1556/ComEc.8.2007.2.3.

- 442 Debastiani, V.J. and Pillar, V.D. 2012. SYNCSA — R tool for analysis of metacommunities  
443 based on functional traits and phylogeny of the community components. *Bioinformatics*  
444 **28**: 2067-2068.
- 445 Dolbeer, R.A. 2008. Ornithology and integrated pest management: Red-winged Blackbirds  
446 *Agelaius phoeniceus* and corn [Article]. *Ibis* **132**(2): 309-322. doi:10.1111/j.1474-  
447 919X.1990.tb01048.x.
- 448 Dray, S. and Dufour, A.B. 2007. The ade4 package: implementing the duality diagram for  
449 ecologists. *Journal of Statistical Software* **22**(4): 1-20.
- 450 Dray, S., Choler, P., Doledec, S., Peres-Neto, P.R., Thuiller, W., Pavoine, S., and ter Braak, C.J.  
451 2014. Combining the fourth-corner and the RLQ methods for assessing trait responses to  
452 environmental variation [Article]. *Ecology* **95**(1): 14-21. doi:10.1890/13-0196.1.
- 453 Drever, M.C. and Martin, K. 2010. Response of woodpeckers to changes in forest health and  
454 harvest: Implications for conservation of avian biodiversity [Article]. *Forest Ecology and*  
455 *Management* **259**(5): 958-966. doi:10.1016/j.foreco.2009.11.038.
- 456 Elmqvist, T., Folke, C., Nyström, M., Peterson, G., Bengtsson, J., Walker, B., and Norberg, J.  
457 2003. Response diversity, ecosystem change, and resilience [Review]. *Frontiers in*  
458 *Ecology and the Environment* **1**(9): 488-494. Available from  
459 [https://www.scopus.com/inward/record.uri?eid=2-s2.0-](https://www.scopus.com/inward/record.uri?eid=2-s2.0-33845673122&partnerID=40&md5=f6f5a6d356c582a09c7163beab83d3cf)  
460 [33845673122&partnerID=40&md5=f6f5a6d356c582a09c7163beab83d3cf](https://www.scopus.com/inward/record.uri?eid=2-s2.0-33845673122&partnerID=40&md5=f6f5a6d356c582a09c7163beab83d3cf) [accessed.
- 461 Gregory, R.D., van Strien, A., Vorisek, P., Gmelig Meyling, A.W., Noble, D.G., Foppen, R.P.,  
462 and Gibbons, D.W. 2005. Developing indicators for European birds. *Philos Trans R Soc*  
463 *Lond B Biol Sci* **360**(1454): 269-288. doi:10.1098/rstb.2004.1602.
- 464 Grenié, M., Denelle, P., Tucker, C.M., Munoz, F., and Violle, C. 2017. funrar: An R package to  
465 characterize functional rarity. R package version 1.2.1.
- 466 Hobson, K.A. and Schieck, J. 1999. Changes in bird communities in boreal mixedwood forest:  
467 Harvest and wildfire effects over 30 years [Article]. *Ecological Applications* **9**(3): 849-  
468 863. Available from [http://www.scopus.com/inward/record.url?eid=2-s2.0-](http://www.scopus.com/inward/record.url?eid=2-s2.0-0033178333&partnerID=40&md5=88e4f74f2a0398678f0ac07a17f09626)  
469 [0033178333&partnerID=40&md5=88e4f74f2a0398678f0ac07a17f09626](http://www.scopus.com/inward/record.url?eid=2-s2.0-0033178333&partnerID=40&md5=88e4f74f2a0398678f0ac07a17f09626) [accessed.
- 470 Hutto, R.L. and Gallo, S.M. 2006. The Effects of Postfire Salvage Logging on Cavity-Nesting  
471 Birds. *The Condor* **108**(4): 817-831. doi:10.1093/condor/108.4.817.
- 472 Hutto, R.L., Bond, M.L., and DellaSala, D.A. 2015. Using Bird Ecology to Learn About the  
473 Benefits of Severe Fire. *In* *The Ecological Importance of Mixed-Severity Fires*. pp. 55-  
474 88.
- 475 Jetz, W., Thomas, G., Joy, J., Hartmann, K., and Mooers, A. 2014. A global phylogeny of birds.  
476 2014. BirdTree.org <http://birdtree.org/downloads/>.
- 477 Kelly, J.F., Bridge, E.S., Frick, W.F., and Chilson, P.B. 2013. Ecological energetics of an  
478 abundant aerial insectivore, the Purple Martin [Article]. *PLoS One* **8**(9): e76616.  
479 doi:10.1371/journal.pone.0076616.
- 480 Klingbeil, B.T. and Willig, M.R. 2016a. Matrix composition and landscape heterogeneity  
481 structure multiple dimensions of biodiversity in temperate forest birds [Article].  
482 *Biodiversity and Conservation* **25**(13): 2687-2708. doi:10.1007/s10531-016-1195-6.
- 483 Klingbeil, B.T. and Willig, M.R. 2016b. Community assembly in temperate forest birds: habitat  
484 filtering, interspecific interactions and priority effects [Article]. *Evolutionary Ecology*  
485 **30**(4): 703-722. doi:10.1007/s10682-016-9834-7.
- 486 Koivula, M.J. and Schmiegelow, F.K.A. 2007. Boreal woodpecker assemblages in recently  
487 burned forested landscapes in Alberta, Canada: Effects of post-fire harvesting and burn

- 488 severity. *Forest Ecology and Management* **242**(2-3): 606-618.  
 489 doi:10.1016/j.foreco.2007.01.075.
- 490 Laliberte, E., Wells, J.A., Declerck, F., Metcalfe, D.J., Catterall, C.P., Queiroz, C., Aubin, I.,  
 491 Bonser, S.P., Ding, Y., Fraterrigo, J.M., McNamara, S., Morgan, J.W., Merlos, D.S.,  
 492 Vesk, P.A., and Mayfield, M.M. 2010. Land-use intensification reduces functional  
 493 redundancy and response diversity in plant communities [Article]. *Ecol Lett* **13**(1): 76-86.  
 494 doi:10.1111/j.1461-0248.2009.01403.x.
- 495 Lavorel, S. and Garnier, E. 2002. Predicting changes in community composition and ecosystem  
 496 functioning from plant traits: revisiting the Holy Grail [Review]. *Functional Ecology*  
 497 **16**(5): 545-556. doi:10.1046/j.1365-2435.2002.00664.x.
- 498 Legendre, P. and Gallagher, E.D. 2001. Ecologically meaningful transformations for ordination  
 499 of species data [Article]. *Oecologia* **129**(2): 271-280. doi:10.1007/s004420100716.
- 500 Lewis, K. and Hartley, I. 2006. Rate of deterioration, degrade, and fall of trees killed by  
 501 mountain pine beetle. *BC Journal of Ecosystems and Management* **7**(2): 11-19.
- 502 Lindenmayer, D., Thorn, S., and Banks, S. 2017. Please do not disturb ecosystems further. *Nat*  
 503 *Ecol Evol* **1**(2): 31. doi:10.1038/s41559-016-0031.
- 504 Lindenmayer, D.B. and Noss, R.F. 2006. Salvage logging, ecosystem processes, and biodiversity  
 505 conservation. *Conserv Biol* **20**(4): 949-958. doi:10.1111/j.1523-1739.2006.00497.x.
- 506 Lindenmayer, D.B., Franklin, J.F., and Fischer, J. 2006. General management principles and a  
 507 checklist of strategies to guide forest biodiversity conservation. *Biological Conservation*  
 508 **131**(3): 433-445. doi:10.1016/j.biocon.2006.02.019.
- 509 Lindenmayer, D.B., Burton, P.J., and Franklin, J.F. 2008. Salvage logging and its ecological  
 510 consequences. Island Press, Washington, D.C., USA. 246 p.
- 511 Lindenmayer, D.B., Blanchard, W., McBurney, L., Blair, D., Banks, S.C., Driscoll, D.A., Smith,  
 512 A.L., Gill, A.M., and Cumming, G. 2014. Complex responses of birds to landscape-level  
 513 fire extent, fire severity and environmental drivers [Article]. *Diversity and Distributions*  
 514 **20**(4): 467-477. doi:10.1111/ddi.12172.
- 515 Luck, G.W., Carter, A., and Smallbone, L. 2013. Changes in bird functional diversity across  
 516 multiple land uses: interpretations of functional redundancy depend on functional group  
 517 identity [Article]. *PLoS One* **8**(5): e63671. doi:10.1371/journal.pone.0063671.
- 518 Luck, G.W., Lavorel, S., McIntyre, S., and Lumb, K. 2012. Improving the application of  
 519 vertebrate trait-based frameworks to the study of ecosystem services [Article]. *J Anim*  
 520 *Ecol* **81**(5): 1065-1076. doi:10.1111/j.1365-2656.2012.01974.x.
- 521 Mayfield, M.M., Bonser, S.P., Morgan, J.W., Aubin, I., McNamara, S., and Vesk, P.A. 2010.  
 522 What does species richness tell us about functional trait diversity? Predictions and  
 523 evidence for responses of species and functional trait diversity to land-use change  
 524 [Article]. *Global Ecology and Biogeography* **19**(4): 423-431. doi:10.1111/j.1466-  
 525 8238.2010.00532.x.
- 526 McIver, J. and Starr, L. 2000. Environmental Effects of Postfire Logging: Literature Review and  
 527 Annotated Bibliography, United States Department of Agriculture (USDA), Forest  
 528 Service, Pacific Northwest Research Station. General Technical Report PNW-GTR-486.
- 529 Mikusiński, G., Roberge, J., and Fuller, R. 2018. Ecology and Conservation of Forest Birds.  
 530 Ecology, Biodiversity and Conservation. Cambridge University Press, Cambridge.
- 531 Morissette, J.L., Cobb, T.P., Brigham, R.M., and James, P.C. 2002. The response of boreal forest  
 532 songbird communities to fire and post-fire harvesting. *Canadian Journal of Forest*  
 533 *Research* **32**(12): 2169-2183. doi:10.1139/x02-134.

- 534 Mouillot, D., Graham, N.A., Villeger, S., Mason, N.W., and Bellwood, D.R. 2013. A functional  
535 approach reveals community responses to disturbances [Review]. *Trends Ecol Evol*  
536 **28**(3): 167-177. doi:10.1016/j.tree.2012.10.004.
- 537 Müller, J., Noss, R.F., Thorn, S., Bäessler, C., Leverkus, A.B., and Lindenmayer, D. 2019.  
538 Increasing disturbance demands new policies to conserve intact forest. *Conservation*  
539 *Letters* **12**(1). doi:10.1111/conl.12449.
- 540 Naeem, S. 1998. Species Redundancy and Ecosystem Reliability [Article]. *Conservation Biology*  
541 **12**(1): 39-45. doi:10.1046/j.1523-1739.1998.96379.x.
- 542 Nappi, A. and Drapeau, P. 2009. Reproductive success of the black-backed woodpecker  
543 (*Picoides arcticus*) in burned boreal forests: Are burns source habitats? [Article].  
544 *Biological Conservation* **142**(7): 1381-1391. doi:10.1016/j.biocon.2009.01.022.
- 545 Nappi, A. and Drapeau, P. 2011. Pre-fire forest conditions and fire severity as determinants of  
546 the quality of burned forests for deadwood-dependent species: the case of the black-  
547 backed woodpecker [Article]. *Canadian Journal of Forest Research* **41**(5): 994-1003.  
548 doi:10.1139/x11-028.
- 549 Nappi, A., Drapeau, P., and Savard, J.P.L. 2004. Salvage logging after wildfire in the boreal  
550 forest: Is it becoming a hot issue for wildlife? *Forestry Chronicle* **80**(1): 67-74. Available  
551 from <Go to ISI>://CCC:000220659900050 [accessed].
- 552 Nappi, A., Drapeau, P., Saint-Germain, M., and Angers, V.A. 2010. Effect of fire severity on  
553 long-term occupancy of burned boreal conifer forests by saproxylic insects and wood-  
554 foraging birds [Article]. *International Journal of Wildland Fire* **19**(4): 500-511.  
555 doi:10.1071/wf08109.
- 556 Naylor, R. and Ehrlich, P. 1997. The value of natural pest control services in agriculture. *In*  
557 *Nature's Services: Societal Dependence on Natural Ecosystems*. Edited by G. Daily.  
558 Island Press, Washington, D.C. pp. 151-174.
- 559 Nebel, S., Mills, A., McCracken, J.D., and Taylor, P.D. 2010. Declines of aerial insectivores in  
560 North America follow a geographic gradient [Article]. *Avian Conservation and Ecology*  
561 **5**(2). Available from <https://www.scopus.com/inward/record.uri?eid=2-s2.0-78651105296&partnerID=40&md5=1726e0429925bad7a5c37c013230c776> [accessed].
- 562 Petchey, O.L., Evans, K.L., Fishburn, I.S., and Gaston, K.J. 2007. Low functional diversity and  
563 no redundancy in British avian assemblages. *Journal of Animal Ecology* **76**: 977-985.
- 564 Poole, A. 2005. The birds of North America online: <http://bna.birds.cornell.edu/BNA/>. Cornell  
565 Laboratory of Ornithology, Ithaca.
- 566 Price, D.T., Alfaro, R.I., Brown, K.J., Flannigan, M.D., Fleming, R.A., Hogg, E.H., Girardin,  
567 M.P., Lakusta, T., Johnston, M., McKenney, D.W., Pedlar, J.H., Stratton, T., Sturrock,  
568 R.N., Thompson, I.D., Trofymow, J.A., and Venier, L.A. 2013. Anticipating the  
569 consequences of climate change for Canada's boreal forest ecosystems. *Environmental*  
570 *Reviews* **21**(4): 322-365. Available from <Go to ISI>://CCC:000328470300006  
571 [accessed].
- 572 R Core Team. 2020. R: A language and environment for statistical computing. R Foundation for  
573 Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0, URL [http://www.R-](http://www.R-project.org/)  
574 [project.org/](http://www.R-project.org/).
- 575 Rao, C.R. 1982. Diversity and dissimilarity coefficients: A unified approach [Article].  
576 *Theoretical Population Biology* **21**(1): 24-43. doi:10.1016/0040-5809(82)90004-1.
- 577 Ricotta, C., Bello, F., Moretti, M., Caccianiga, M., Cerabolini, B.E.L., Pavoine, S., and  
578 Peres-Neto, P. 2016. Measuring the functional redundancy of biological communities: a  
579

- 580 quantitative guide. *Methods in Ecology and Evolution* **7**(11): 1386-1395.  
 581 doi:10.1111/2041-210x.12604.
- 582 Rigby, R.A. and Stasinopoulos, D.M. 2005. Generalized additive models for location, scale and  
 583 shape,(with discussion). *Appl. Statist.* **54**: 507-554.
- 584 Saint-Germain, M. and Greene, D.F. 2009. Salvage logging in the boreal and cordilleran forests  
 585 of Canada: Integrating industrial and ecological concerns in management plans. *The*  
 586 *Forestry chronicle* **85**(1): 120-134.
- 587 Saint-Germain, M., Drapeau, P., and Hébert, C. 2004. Xylophagous insect species composition  
 588 and patterns of substratum use on fire-killed black spruce in central Quebec. *Canadian*  
 589 *Journal of Forest Research* **34**(3): 677-685. doi:10.1139/x03-235.
- 590 Saint-Germain, M., Drapeau, P., and M. Buddle, C. 2007. Host-use patterns of saproxylic  
 591 phloeophagous and xylophagous Coleoptera adults and larvae along the decay gradient in  
 592 standing dead black spruce and aspen. *Ecography* **30**(6): 737-748.  
 593 doi:10.1111/j.2007.0906-7590.05080.x.
- 594 Sauer, J.R., Niven, D.K., Hines, J.E., Ziolkowski Jr., D.J., Pardieck, K.L., Fallon, J.E., and Link,  
 595 W.A. 2017. The North American breeding bird survey, results and analysis 1966-2015.  
 596 Available: <http://www.mbr-pwrc.usgs.gov/bbs/> Accessed 30.03.2017.
- 597 Schieck, J. and Hobson, K.A. 2000. Bird communities associated with live residual tree patches  
 598 within cut blocks and burned habitat in mixedwood boreal forests [Article]. *Canadian*  
 599 *Journal of Forest Research* **30**(8): 1281-1295. Available from  
 600 [https://www.scopus.com/inward/record.uri?eid=2-s2.0-  
 601 0033780243&partnerID=40&md5=8fd79c66f64c3a4d61853a11c7f5da7f](https://www.scopus.com/inward/record.uri?eid=2-s2.0-0033780243&partnerID=40&md5=8fd79c66f64c3a4d61853a11c7f5da7f) [accessed].
- 602 Schieck, J. and Song, S.J. 2006. Changes in bird communities throughout succession following  
 603 fire and harvest in boreal forests of western North America: literature review and meta-  
 604 analyses [Article]. *Canadian Journal of Forest Research* **36**(5): 1299-1318.  
 605 doi:10.1139/x06-017.
- 606 Simpson, E. 1949. Measurement of diversity. . *Nature*: 163:688.
- 607 Smucker, K.M., Hutto, R.L., and Steele, B.M. 2005. Changes in bird abundance after wildfire:  
 608 Importance of fire severity and time since fire [Article]. *Ecological Applications* **15**(5):  
 609 1535-1549. Available from [https://www.scopus.com/inward/record.uri?eid=2-s2.0-  
 610 26044475582&partnerID=40&md5=17f9021202696d76233ee50a23cbc8c1](https://www.scopus.com/inward/record.uri?eid=2-s2.0-26044475582&partnerID=40&md5=17f9021202696d76233ee50a23cbc8c1) [accessed].
- 611 Song, S.J. and Hannah, J. unpublished. Unpublished data.
- 612 Stasinopoulos, M. and Rigby, R.A. 2019. *gamlss.dist: Distributions for Generalized Additive*  
 613 *Models for Location Scale and Shape*. R package version 5.1-4. [https://CRAN.R-  
 614 project.org/package=gamlss.dist](https://CRAN.R-project.org/package=gamlss.dist).
- 615 Stasinopoulos, M., Rigby, R.A., Heller, G.Z., Voudouris, V., and De Bastiani, F. 2017. *Flexible*  
 616 *Regression and Smoothing Using GAMLSS in R*. Boca Raton, FL: CRC Press.
- 617 Stephens, J.L., Ausprey, I.J., Seavy, N.E., and Alexander, J.D. 2015. Fire severity affects mixed  
 618 broadleaf–conifer forest bird communities: Results for 9 years following fire [Article].  
 619 *The Condor* **117**(3): 430-446. doi:10.1650/condor-14-58.1.
- 620 Swanson, M.E., Franklin, J.F., Beschta, R.L., Crisafulli, C.M., DellaSala, D.A., Hutto, R.L.,  
 621 Lindenmayer, D.B., and Swanson, F.J. 2011. The forgotten stage of forest succession:  
 622 early-successional ecosystems on forest sites. *Frontiers in Ecology and the Environment*  
 623 **9**(2): 117-125. doi:10.1890/090157.



- 624 Tarbill, G.L., Manley, P.N., and White, A.M. 2015. Drill, baby, drill: the influence of  
625 woodpeckers on post-fire vertebrate communities through cavity excavation [Article].  
626 *Journal of Zoology* **296**(2): 95-103. doi:10.1111/jzo.12220.
- 627 Tews, J., Brose, U., Grimm, V., Tielborger, K., Wichmann, M.C., Schwager, M., and Jeltsch, F.  
628 2004. Animal species diversity driven by habitat heterogeneity/diversity: the importance  
629 of keystone structures. *Journal of Biogeography* **31**(1): 79-92. Available from <Go to  
630 ISI>://CCC:000187549700008 [accessed].
- 631 Thorn, S., Werner, S.A.B., Wohlfahrt, J., Bässler, C., Seibold, S., Quillfeldt, P., and Müller, J.  
632 2016. Response of bird assemblages to windstorm and salvage logging — Insights from  
633 analyses of functional guild and indicator species [Article]. *Ecological Indicators* **65**:  
634 142-148. doi:10.1016/j.ecolind.2015.06.033.
- 635 Van Bael, S.A., Brawn, J.D., and Robinson, S.K. 2003. Birds defend trees from herbivores in a  
636 Neotropical forest canopy [Article]. *Proc Natl Acad Sci U S A* **100**(14): 8304-8307.  
637 doi:10.1073/pnas.1431621100.
- 638 Van Wilgenburg, S.L. and Hobson, K.A. 2008. Landscape-scale disturbance and boreal forest  
639 birds: Can large single-pass harvest approximate fires? *Forest Ecology and Management*  
640 **256**(1-2): 136-146. doi:10.1016/j.foreco.2008.04.017.
- 641 Vandewalle, M., de Bello, F., Berg, M.P., Bolger, T., Doledec, S., Dubs, F., Feld, C.K.,  
642 Harrington, R., Harrison, P.A., Lavorel, S., da Silva, P.M., Moretti, M., Niemela, J.,  
643 Santos, P., Sattler, T., Sousa, J.P., Sykes, M.T., Vanbergen, A.J., and Woodcock, B.A.  
644 2010. Functional traits as indicators of biodiversity response to land use changes across  
645 ecosystems and organisms. *Biodiversity and Conservation* **19**(10): 2921-2947. Available  
646 from <Go to ISI>://CCC:000280807700009 [accessed].
- 647 Venier, L.A. and Pearce, J.L. 2004. Birds as indicators of sustainable forest management  
648 [Review]. *Forestry Chronicle* **80**(1): 61-66. Available from  
649 [https://www.scopus.com/inward/record.uri?eid=2-s2.0-  
650 2442572111&partnerID=40&md5=6265fdb7cb57ca4c6523d2df4baeac6](https://www.scopus.com/inward/record.uri?eid=2-s2.0-2442572111&partnerID=40&md5=6265fdb7cb57ca4c6523d2df4baeac6) [accessed].
- 651 Violle, C., Thuiller, W., Mouquet, N., Munoz, F., Kraft, N.J.B., Cadotte, M.W., Livingstone,  
652 S.W., and Mouillot, D. 2017. Functional Rarity: The Ecology of Outliers [Review].  
653 *Trends Ecol Evol* **32**(5): 356-367. doi:10.1016/j.tree.2017.02.002.
- 654 Wickham, H. 2016. *ggplot2: Elegant Graphics for Data Analysis*. Springer-Verlag New York.
- 655 Zuur, A.F. and Ieno, E.N. 2016. A protocol for conducting and presenting results of  
656 regression-type analyses. *Methods in Ecology and Evolution* **7**(6): 636-645.  
657 doi:10.1111/2041-210X.12577.
- 658 Zuur, A.F., Ieno, E.N., and Elphick, C.S. 2010. A protocol for data exploration to avoid common  
659 statistical problems. *Methods in Ecology and Evolution* **1**(1): 3-14. doi:10.1111/j.2041-  
660 210X.2009.00001.x.

661

**Table 1.** Description of the selected traits

Functional Trait	Class	Example of responses and effects on ecosystem function
Foraging strategy	Ground forager Bark forager Foliage gleaner Aerial forager	Resource use (e.g. reduced primary diet) and competitiveness (e.g. enhanced pollinisation by insects)
Foraging guild	Herbivorous Insectivorous Omnivorous Carnivorous	Resource use, seed dispersal (e.g. reduced seed germination and seedling survival) and predation (e.g. pest regulation)
Habitat affinity	Open canopy Forest	Resource use (e.g. reduced habitat specialists), reproductive effort (e.g. reduce reproduction sites), ecosystem engineering (e.g. tree cavities excavation providing habitat features for seed dispersers, insectivores and small predators)
Nest location	Ground Shrub Tree	Reproductive effort (e.g. reduced response capacity due to low reproductive sites), ecosystem engineering (e.g. modification of soil, trees and canopy structures, creating substantial nutrient concentration)
Nest guild	Non-cavity-nester Cavity-nester	Reproductive effort (e.g. loss of large cavity bearing tree reduce cavity nesting bird), ecosystem engineering (e.g. modification of large trees and canopy structures, provision of nesting habitats for other organisms)
Clutch size (Mean)	Quantitative	Reproductive effort (e.g. reduced response capacity due to low reproductive rates) and trophic interactions (e.g. prey availability and

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			predation pressure)
Body (Average)	mass	Quantitative	Resource use (e.g. reduced response capacity due to low turnover rates), mobility (e.g. habitat fragmentation)

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Trait description and examples of potential responses to disturbance and effects on ecosystem function according to literature (Luck et al., 2012; Concepción et al., 2016a; Williams and Purves, 2011) are provided.

**Table 2.** Likelihood ration tests (LRT) for the predictors of Gini-Simpson index (A), Rao quadratic entropy considering trait distance (B), Rao quadratic entropy considering phylogenetic distance (C), Functional redundancy (D), and Phylogenetic redundancy (E)

	df	AIC	LRT	p-value
A. Gini-Simpson index (SIMPLEX, $\mu$ and $\sigma$ )				
$\mu$				
Starting model		-62.31		
Disturbance	5.46	-49.02	24.21	<0.001
Random factor	3.47	-046.56	20.67	<0.001
$\sigma$				
Starting model		-62.31		
Disturbance	2.70	-56.73	10.98	0.009
B. Rao quadratic entropy with trait distance (SIMPLEX, $\mu$ and $\sigma$ )				
$\mu$				
Starting model		-57.88		
Disturbance	1.94	-60.39	1.38	0.488
Random factor	3.57	-46.61	18.41	<0.001
$\sigma$				
Starting model		-57.88		
Disturbance	2.50	-52.47	10.41	0.009
C. Rao quadratic entropy with phylogenetic distance (BE, $\mu$ and $\sigma$ )				
$\mu$				
Starting model		-154.15		
Disturbance	1	-148.46	7.96	0.006
Random factor	<0.01	-153.67	0.48	<0.001
$\sigma$				
Starting model		-154.15		
Disturbance	-1.19	-149.96	1.81	<0.001
D. Function redundancy (BE, $\mu$ and $\sigma$ )				
$\mu$				
Starting model		-66.51		
Disturbance	5.08	-57.96	18.72	0.002
Random factor	3.08	-63.50	9.18	0.029
$\sigma$				
Starting model		-66.51		
Disturbance	5.08	-57.95	18.72	0.002
E. Phylogenetic Redundancy (BE, $\mu$ and $\sigma$ )				
$\mu$				
Starting model		-60.76		
Disturbance	5.26	-48.45	22.64	<0.001

Random factor	3.26	-48.31	18.98	<0.001
$\sigma$				
Starting model		-60.76		
Disturbance	2.37	-55.44	10.07	0.010

**Table 3.** Predicted values (Mean  $\pm$  SE) across post-disturbed habitats of individual traits analysis for  $\mu$  component of: (A) the (SIMPLEX,  $\mu$  and  $\sigma$ ) model for FD; (B) the (BE,  $\mu$  and  $\sigma$ ) model for FR . Bolded values indicate traits where a significant difference was detected between disturbance types (see Results).

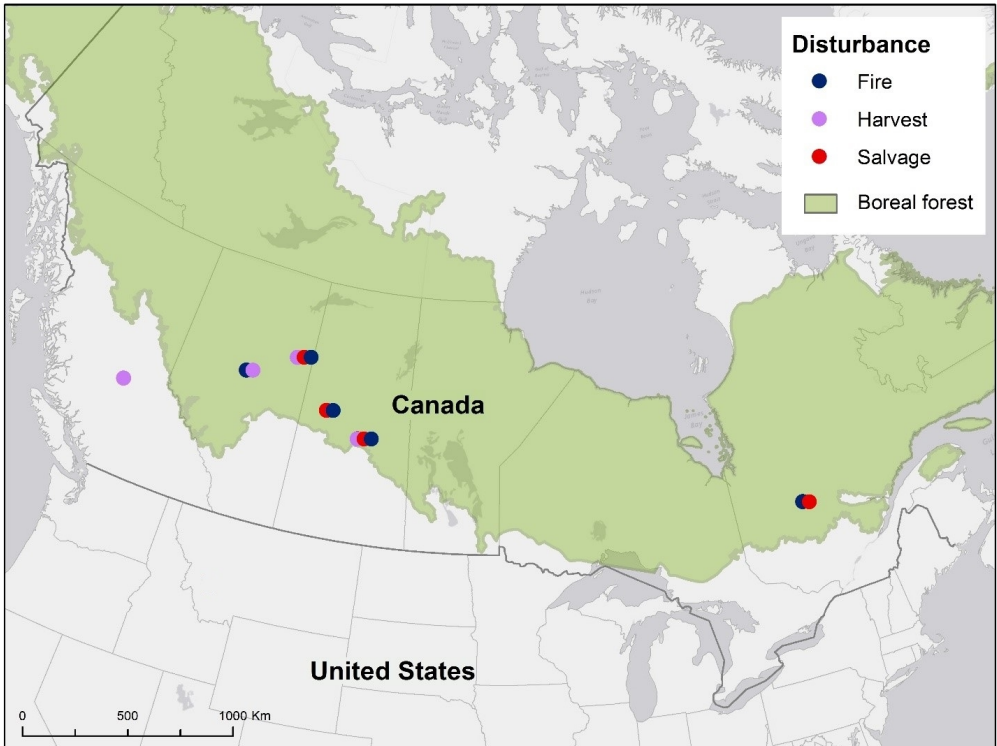
	Fire (n=7)	Traditional logging (n=4)	Salvage logging (n=6)
<b>A. Functional diversity measured as Rao' s quadratic entropy index (FD)</b>			
Foraging strategy	<b>0.61<math>\pm</math>0.06<sup>a</sup></b>	<b>0.59<math>\pm</math>0.06<sup>a</sup></b>	<b>0.54<math>\pm</math>0.11<sup>b</sup></b>
Foraging guild	0.48 $\pm$ 0.06 <sup>a</sup>	0.50 $\pm$ 0.06 <sup>a</sup>	0.48 $\pm$ 0.11 <sup>a</sup>
Habitat affinity	0.54 $\pm$ 0.06 <sup>a</sup>	0.55 $\pm$ 0.06 <sup>a</sup>	0.54 $\pm$ 0.11 <sup>a</sup>
Nest Location	0.56 $\pm$ 0.06 <sup>a</sup>	0.55 $\pm$ 0.06 <sup>a</sup>	0.56 $\pm$ 0.11 <sup>a</sup>
Nest guild	<b>0.31<math>\pm</math>0.06<sup>a</sup></b>	<b>0.19<math>\pm</math>0.06<sup>c</sup></b>	<b>0.24<math>\pm</math>0.11<sup>b</sup></b>
Body Mass	0.06 $\pm$ 0.06 <sup>a</sup>	0.07 $\pm$ 0.06 <sup>a</sup>	0.09 $\pm$ 0.11 <sup>a</sup>
Clutch size	0.20 $\pm$ 0.06 <sup>a</sup>	0.20 $\pm$ 0.06 <sup>a</sup>	0.16 $\pm$ 0.11 <sup>a</sup>
<b>B. Functional redundancy index (FR)</b>			
Foraging strategy	0.31 $\pm$ 0.09 <sup>a</sup>	0.36 $\pm$ 0.10 <sup>a</sup>	0.34 $\pm$ 0.07 <sup>a</sup>
Foraging guild	<b>0.42<math>\pm</math>0.12<sup>a</sup></b>	<b>0.44<math>\pm</math>0.05<sup>ab</sup></b>	<b>0.36<math>\pm</math>0.08<sup>b</sup></b>
Habitat affinity	<b>0.52<math>\pm</math>0.12<sup>a</sup></b>	<b>0.50<math>\pm</math>0.11<sup>ab</sup></b>	<b>0.40<math>\pm</math>0.13<sup>b</sup></b>
Nest Location	<b>0.36<math>\pm</math>0.08<sup>a</sup></b>	<b>0.38<math>\pm</math>0.11<sup>ab</sup></b>	<b>0.30<math>\pm</math>0.14<sup>b</sup></b>
Nest guild	0.66 $\pm$ 0.16 <sup>a</sup>	0.75 $\pm$ 0.22 <sup>a</sup>	0.66 $\pm$ 0.10 <sup>a</sup>
Body Mass	<b>0.84<math>\pm</math>0.09<sup>b</sup></b>	<b>0.86<math>\pm</math>0.10<sup>a</sup></b>	<b>0.74<math>\pm</math>0.07<sup>c</sup></b>
Clutch size	<b>0.70<math>\pm</math>0.05<sup>ab</sup></b>	<b>0.73<math>\pm</math>0.04<sup>a</sup></b>	<b>0.65<math>\pm</math>0.13<sup>b</sup></b>

**Table 4.** Summary of abundant or scarce species with high functional uniqueness. All species scarcity and functional uniqueness indices are reported in Table A1.

Species	Species scarcity index			Functional uniqueness index		
	Harvest	Fire	Salvage logging	Harvest	Fire	Salvage logging
<i>Bucephala albeola</i>	0.974	0.948	NA	0.017	0.017	NA
<i>Cardellina pusilla</i>	0.000	0.485	0.655	0.145	0.145	0.146
<i>Catharus ustulatus</i>	0.079	0.220	0.521	0.021	0.021	0.145
<i>Certhia americana</i>	0.760	0.281	0.572	0.146	0.144	0.146
<i>Chaetura vauxi</i>	NA	0.000	NA	NA	0.018	NA
<i>Contopus cooperi</i>	0.000	0.012	NA	0.005	0.005	NA
<i>Empidonax flaviventris</i>	0.818	0.535	0.764	0.146	0.146	0.146
<i>Empidonax minimus</i>	0.241	0.394	0.497	0.143	0.018	0.143
<i>Empidonax traillii</i>	NA	0.007	0.985	NA	0.002	0.145
<i>Ixoreus naevius</i>	0.195	0.000	NA	0.017	0.017	NA
<i>Mniotilta varia</i>	0.810	0.641	0.897	0.144	0.144	0.144
<i>Pheucticus melanocephalus</i>	NA	0.005	NA	NA	0.011	NA
<i>Poecile atricapillus</i>	0.374	0.279	0.812	0.144	0.001	0.144
<i>Poecile gambeli</i>	NA	0.000	NA	NA	0.001	NA
<i>Setophaga fusca</i>	0.889	0.974	NA	0.003	0.003	NA
<i>Setophaga townsendi</i>	NA	0.000	NA	NA	0.003	NA
<i>Sialia currucoides</i>	0.000	0.003	0.856	0.006	0.004	0.007
<i>Sialia mexicana</i>	NA	0.002	NA	NA	0.004	NA
<i>Sphyrapicus nuchalis</i>	NA	0.000	NA	NA	0.002	NA

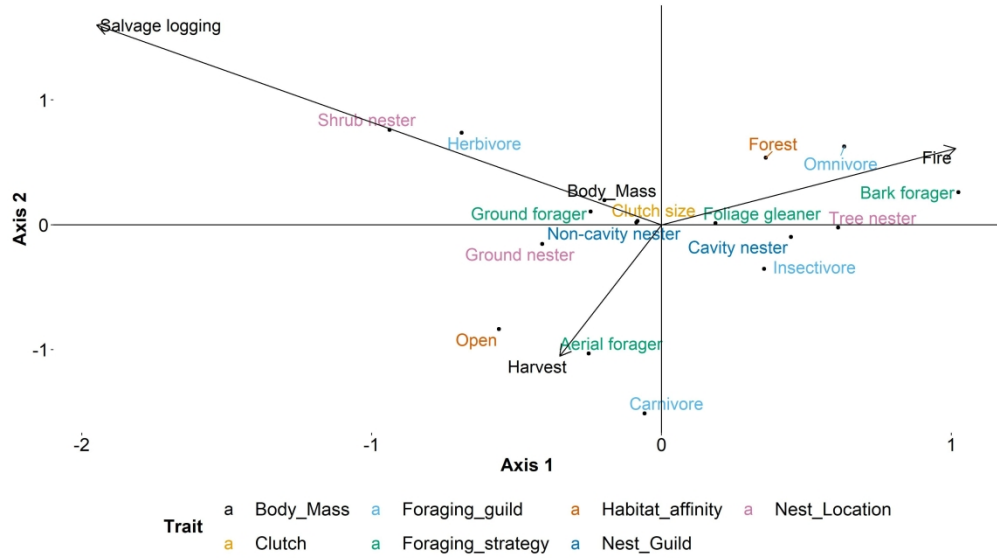
<i>Sphyrapicus thyroideus</i>	NA	0.000	NA	NA	0.001	NA
<i>Sturnus vulgaris</i>	NA	0.000	NA	NA	0.010	NA
<i>Troglodytes aedon</i>	0.736	0.155	0.080	0.146	0.146	0.146
<i>Troglodytes hiemalis</i>	0.334	0.246	0.580	0.146	0.146	0.146
<i>Vireo gilvus</i>	0.494	0.000	0.985	0.145	0.145	0.030





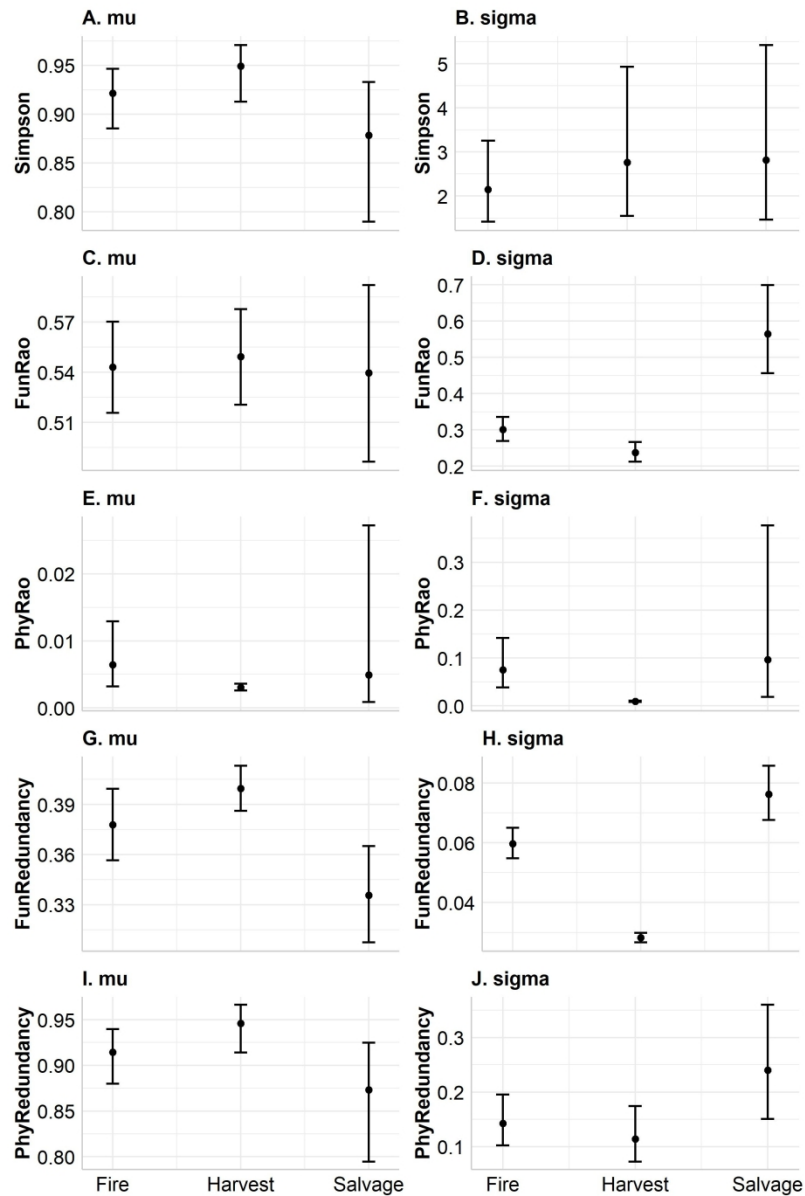
Study geographical distribution (adapted from Brandt 2009, created in ArcGIS). No permissions required from the original source/copyright holders to publish this figure.

165x125mm (220 x 220 DPI)



Relationship between traits and disturbances using RLQ analysis. Disturbance classes (Fire, Harvest, and Salvage,) are presented as vectors. Functional traits are presented as group centroids

349x199mm (300 x 300 DPI)



Predicted values and confidence intervals of additive terms of Gini-Simpson index (A.  $\mu$  and B.  $\sigma$ ), Rao quadratic entropy considering trait distance (C.  $\mu$  and D.  $\sigma$ ), Rao quadratic entropy considering phylogenetic distance (E.  $\mu$  and F.  $\sigma$ ), Functional redundancy (G.  $\mu$  and H.  $\sigma$ ) and Phylogenetic redundancy (I.  $\mu$  and J.  $\sigma$ ) for both  $\mu$  and  $\sigma$  components of the (SIMPLEX,  $\mu$  and  $\sigma$ ) and (BE,  $\mu$  and  $\sigma$ ) models

199x299mm (300 x 300 DPI)

**Table A1. List of avia**

<b>Species</b>	<b>Species_Code</b>	<b>Species_ID</b>
<i>Accipiter cooperii</i>	ACCO	COHA
<i>Accipiter striatus</i>	ACST	SSHA
<i>Actitis macularius</i>	ACMA	SPSA
<i>Aegolius acadicus</i>	AEAC	NSWO
<i>Aegolius funereus</i>	AEFU	BOOW
<i>Agelaius phoeniceus</i>	AGPH	RWBL
<i>Ammospiza caudacuta</i>	AMCAU	SALS
<i>Ammospiza leconteii</i>	AMLE	LCSP
<i>Anas americana</i>	ANAME	AMWI
<i>Anas crecca</i>	ANCRE	GWTE
<i>Anas cyanoptera</i>	ANCY	CITE
<i>Anas discors</i>	ANDIS	BWTE
<i>Anas platyrhynchos</i>	ANPLA	MALL
<i>Anas rubripes</i>	ANRUB	ABDU
<i>Archilochus colubris</i>	ARCO	RTHU
<i>Asio otus</i>	ASOTU	LEOW
<i>Aythya americana</i>	AYAME	REDH
<i>Aythya collaris</i>	AYCOLL	RNDU
<i>Bombycilla cedrorum</i>	BOCE	CEDW
<i>Bonasa umbellus</i>	BOUM	RUGR
<i>Botaurus lentiginosus</i>	BOLEN	AMBI
<i>Branta canadensis</i>	BRCAN	CANG
<i>Bubo virginianus</i>	BUVI	GHOW
<i>Bucephala albeola</i>	BUALB	BUFF
<i>Bucephala clangula</i>	BUCLA	COGO
<i>Buteo jamaicensis</i>	BUJA	RTHA
<i>Buteo platypterus</i>	BUPL	BWHA
<i>Cardellina canadensis</i>	CACA	CAWA
<i>Cardellina pusilla</i>	CAPU	WIWA
<i>Catharus fuscescens</i>	CAFU	VEER
<i>Catharus guttatus</i>	CAGU	HETH
<i>Catharus ustulatus</i>	CAUS	SWTH
<i>Certhia americana</i>	CEAM	BRCR
<i>Chaetura vauxi</i>	CHVA	VASW
<i>Charadrius vociferus</i>	CHVO	KILL
<i>Chlidonias niger</i>	CHNIG	BLTE
<i>Chordeiles minor</i>	CHMI	CONI
<i>Cinclus mexicanus</i>	CIME	AMDI
<i>Circus cyaneus</i>	CICY	NOHA
<i>Coccythraustes vespertinus</i>	COVE	EVGR
<i>Colaptes auratus</i>	COAU	NOFL
<i>Contopus cooperi</i>	COCO	OSFL
<i>Contopus sordidulus</i>	COSO	WEWP
<i>Corvus brachyrhynchos</i>	COBR	AMCR
<i>Corvus corax</i>	CORCOR	CORA
<i>Cyanocitta cristata</i>	CYCR	BLJA
<i>Cyanocitta stelleri</i>	CYST	STJA
<i>Dendragapus obscurus</i>	DEOB	DUGR

<i>Dryocopus pileatus</i>	DRPI	PIWO
<i>Empidonax alnorum</i>	EMAL	ALFL
<i>Empidonax flaviventris</i>	EMFL	YBFL
<i>Empidonax hammondi</i>	EMHA	HAFL
<i>Empidonax minimus</i>	EMMI	LEFL
<i>Empidonax oberholseri</i>	EMOB	DUFL
<i>Empidonax traillii</i>	EMTR	WIFL
<i>Euphagus carolinus</i>	EUCA	RUBL
<i>Euphagus cyanocephalus</i>	EUCY	BRBL
<i>Falcipectens canadensis</i>	FACA	SPGR
<i>Falco columbarius</i>	FACO	MERL
<i>Falco sparverius</i>	FASP	AMKE
<i>Fulica americana</i>	FUAME	AMCO
<i>Gallinago delicata</i>	GADE	WISN
<i>Gavia immer</i>	GAIM	COLO
<i>Geothlypis philadelphia</i>	GEPH	MOWA
<i>Geothlypis tolmiei</i>	GETO	MGWA
<i>Geothlypis trichas</i>	GETR	COYE
<i>Grus canadensis</i>	GRCA	SACR
<i>Haemorhous cassinii</i>	HACA	CAFI
<i>Haemorhous purpureus</i>	HAPU	PUFI
<i>Haliaeetus leucocephalus</i>	HALEU	BAEA
<i>Hirundo rustica</i>	HIRUS	BARS
<i>Icterus galbula</i>	ICGAL	BAOR
<i>Ixoreus naevius</i>	IXNA	VATH
<i>Junco hyemalis</i>	JUHY	DEJU
<i>Larus argentatus</i>	LAARG	HERG
<i>Larus delawarensis</i>	LADEL	RBGU
<i>Leucophaeus pipixcan</i>	LEPIP	FRGU
<i>Loxia curvirostra</i>	LOCU	RECR
<i>Loxia leucoptera</i>	LOLE	WWCR
<i>Megasceryle alcyon</i>	MEAL	BEKI
<i>Melospiza georgiana</i>	MEGE	SWSP
<i>Melospiza lincolni</i>	MELI	LISP
<i>Melospiza melodia</i>	MEME	SOSP
<i>Mergus merganser</i>	MEMER	COME
<i>Mniotilta varia</i>	MNVA	BAWW
<i>Molothrus ater</i>	MOAT	BHCO
<i>Myadestes townsendi</i>	MYTO	TOSO
<i>Myiarchus crinitus</i>	MYCRI	GCFL
<i>Nucifraga columbiana</i>	NUCO	CLNU
<i>Oporornis agilis</i>	OPAG	CONW
<i>Oreothlypis celata</i>	ORCE	OCWA
<i>Oreothlypis peregrina</i>	ORPE	TEWA
<i>Oreothlypis ruficapilla</i>	ORRU	NAWA
<i>Pandion haliaetus</i>	PAHA	OSPR
<i>Parkesia noveboracensis</i>	PANO	NOWA
<i>Passerculus sandwichensis</i>	PASA	SAVS
<i>Passerella iliaca</i>	PAIL	FOSP
<i>Passerina amoena</i>	PAAM	LAZB
<i>Perisoreus canadensis</i>	PECA	CAJA
<i>Petrochelidon pyrrhonota</i>	PEPY	CLSW
<i>Pheucticus ludovicianus</i>	PHLU	RBGR

<i>Pheucticus melanocephalus</i>	PHME	BHGR
<i>Pica hudsonia</i>	PIHU	BBMA
<i>Picoides arcticus</i>	PIAR	BBWO
<i>Picoides dorsalis</i>	PIDO	ATTW
<i>Picoides pubescens</i>	PIPU	DOWO
<i>Picoides villosus</i>	PIVI	HAWO
<i>Pinicola enucleator</i>	PIEN	PIGR
<i>Pipilo chlorurus</i>	PICH	GTTO
<i>Pipilo erythrophthalmus</i>	PIER	EATO
<i>Piranga ludoviciana</i>	PILU	WETA
<i>Podiceps grisegena</i>	POGRI	RNGR
<i>Podilymbus podiceps</i>	POPOD	PBGR
<i>Poecile atricapillus</i>	POAT	BCCH
<i>Poecile gambeli</i>	POGA	MOCH
<i>Poecile hudsonicus</i>	POHU	BOCH
<i>Poecile rufescens</i>	PORU	CBCH
<i>Poocetes gramineus</i>	POGR	VESP
<i>Porzana carolina</i>	POCAR	SORA
<i>Quiscalus quiscula</i>	QUQU	COGR
<i>Regulus calendula</i>	RECA	RCKI
<i>Regulus satrapa</i>	RESA	GCKI
<i>Salpinctes obsoletus</i>	SAOB	ROWR
<i>Sayornis phoebe</i>	SAPHO	EAPH
<i>Seiurus aurocapilla</i>	SEAU	OVEN
<i>Selasphorus calliope</i>	SECAL	CAHU
<i>Selasphorus rufus</i>	SERU	RUHU
<i>Setophaga castanea</i>	SECAS	BBWA
<i>Setophaga coronata</i>	SECO	YRWA
<i>Setophaga fusca</i>	SEFU	BLBW
<i>Setophaga magnolia</i>	SEMA	MAWA
<i>Setophaga palmarum</i>	SEPA	PAWA
<i>Setophaga pensylvanica</i>	SEPE	CSWA
<i>Setophaga petechia</i>	SEPET	YEWA
<i>Setophaga ruticilla</i>	SERUT	AMRE
<i>Setophaga tigrina</i>	SETI	CMWA
<i>Setophaga townsendi</i>	SETO	TOWA
<i>Setophaga virens</i>	SEVI	BTNW
<i>Sialia currucoides</i>	SICU	MOBL
<i>Sialia mexicana</i>	SIME	WEBL
<i>Sialia sialis</i>	SISI	EABL
<i>Sitta canadensis</i>	SICA	RBNU
<i>Sitta carolinensis</i>	SICAR	WBNU
<i>Sphyrapicus nuchalis</i>	SPNU	RNSA
<i>Sphyrapicus ruber</i>	SPRU	RBSA
<i>Sphyrapicus thyroideus</i>	SPTH	WISA
<i>Sphyrapicus varius</i>	SPVA	YBSA
<i>Spinus pinus</i>	SPPI	PISI
<i>Spinus tristis</i>	SPTR	AMGO
<i>Spizella pallida</i>	SPPA	CCSP
<i>Spizella passerina</i>	SPPAS	CHSP
<i>Strix nebulosa</i>	STNE	GGOW
<i>Strix varia</i>	STVAR	BADO
<i>Sturnella neglecta</i>	STNEG	WEME

<i>Sturnus vulgaris</i>	STVU	EUST
<i>Surnia ulula</i>	SUUL	NHOW
<i>Tachycineta bicolor</i>	TABI	TRES
<i>Tringa flavipes</i>	TRFL	LEYE
<i>Tringa melanoleuca</i>	TRME	GRYE
<i>Tringa solitaria</i>	TRSO	SOSA
<i>Troglodytes aedon</i>	TRAE	HOWR
<i>Troglodytes hiemalis</i>	TRHI	WIWR
<i>Turdus migratorius</i>	TUMI	AMRO
<i>Tyrannus tyrannus</i>	TYTY	EAKI
<i>Vireo cassinii</i>	VICA	CAVI
<i>Vireo gilvus</i>	VIGI	WAVI
<i>Vireo olivaceus</i>	VIOL	REVI
<i>Vireo philadelphicus</i>	VIPH	PHVI
<i>Vireo solitarius</i>	VISO	BHVI
<i>Xanthocephalus xanthocephalus</i>	XAXAN	YHBL
<i>Zenaida macroura</i>	ZEMA	MOD0
<i>Zonotrichia albicollis</i>	ZOAL	WTSP
<i>Zonotrichia leucophrys</i>	ZOLE	WCSP

## n species with their functional traits, and their taxonomic scarcity and function

English_Name	French_Name	Foraging strategy	Foraging guild
Cooper's Hawk	Épervier de Cooper	A	C
Sharp-shinned Hawk	Épervier brun	G	C
Spotted Sandpiper	Chevalier grivelé	G	C
Northern Saw-whet Owl	Petite Nyctale	G	C
Boreal Owl	Nyctale de Tengmalm	G	C
Red-winged Blackbird	Carouge à épaulettes	G	I
Saltmarsh Sparrow	Bruant à queue aiguë	G	I
LeConte's Sparrow	Bruant de LeConte	G	I
American Wigeon	Canard d'Amérique	G	H
Green-winged Teal	Sarcelle d'hiver	G	H
Cinnamon Teal	Sarcelle cannelle	G	H
Blue-winged Teal	Sarcelle à ailes bleues	G	H
Mallard	Canard colvert	G	H
American Black Duck	Canard noir	G	I
Ruby-throated Hummingbird	Colibri à gorge rubis	F	H
Long-eared Owl	Hibou moyen-duc	G	C
Redhead	Fuligule à tête rouge	G	H
Ring-necked Duck	Fuligule à collier	G	H
Cedar Waxwing	Jaseur d'Amérique	F	H
Ruffed Grouse	Gélinotte huppée	F	O
American Bittern	Butor d'Amérique	G	C
Canada Goose	Bernache du Canada	G	H
Great Horned Owl	Grand-duc d'Amérique	G	C
Bufflehead	Petit Garrot	G	I
Common Goldeneye	Garrot à oeil d'or	G	I
Red-tailed Hawk	Buse à queue rousse	G	C
Broad-winged Hawk	Petite Buse	G	C
Canada Warbler	Paruline du Canada	F	I
Wilson's Warbler	Paruline à calotte noire	F	I
Veery	Grive fauve	G	I
Hermit Thrush	Grive solitaire	G	I
Swainson's Thrush	Grive à dos olive	F	I
Brown Creeper	Grimpereau brun	B	I
Vaux's Swift	Martinet de Vaux	A	I
Killdeer	Pluvier kildir	G	I
Black Tern	Guifette noire	G	I
Common Nighthawk	Engoulevent d'Amérique	G	I
American Dipper	Cinle d'Amérique	G	I
Northern Harrier	Busard des marais	G	C
Evening Grosbeak	Gros-bec errant	G	H
Northern Flicker	Pic flamboyant	G	I
Olive-sided Flycatcher	Moucherolle à côtés olive	A	I
Western Wood-Pewee	Pioui de l'Ouest	A	I
American Crow	Corneille d'Amérique	G	O
Common Raven	Grand Corbeau	G	O
Blue Jay	Geai bleu	G	O
Steller's Jay	Geai de Steller	G	O
Dusky Grouse	Tétras sombre	G	H



Pileated Woodpecker	Grand Pic	B	I
Alder Flycatcher	Moucherolle des aulnes	A	I
Yellow-bellied Flycatcher	Moucherolle à ventre jaune	A	I
Hammond's Flycatcher	Moucherolle de Hammond	A	I
Least Flycatcher	Moucherolle tchébec	A	I
Dusky Flycatcher	Moucherolle sombre	A	I
Willow Flycatcher	Moucherolle des saules	A	I
Rusty Blackbird	Quiscale rouilleux	G	I
Brewer's Blackbird	Quiscale de Brewer	G	H
Spruce Grouse	Tétras du Canada	F	H
Merlin	Faucon émerillon	A	C
American Kestrel	Crécerelle d'Amérique	G	C
American Coot	Foulque d'Amérique	G	H
Wilson's Snipe	Bécassine de Wilson	G	I
Common Loon	Plongeon huard	G	C
Mourning Warbler	Paruline triste	F	I
MacGillivray's Warbler	Paruline des buissons	F	I
Common Yellowthroat	Paruline masquée	F	I
Sandhill Crane	Grue du Canada	G	O
Cassin's Finch	Roselin de Cassin	G	H
Purple Finch	Roselin pourpré	F	H
Bald Eagle	Pygargue à tête blanche	G	C
Barn Swallow	Hirondelle rustique	A	I
Baltimore Oriole	Oriole de Baltimore	F	I
Varied Thrush	Grive à collier	G	I
Dark-eyed Junco	Junco ardoisé	G	H
Herring Gull	Goéland argenté	G	O
Ring-billed Gull	Goéland à bec cerclé	G	O
Franklin's Gull	Mouette de Franklin	G	O
Red Crossbill	Bec-croisé des sapins	F	H
White-winged Crossbill	Bec-croisé bifascié	F	H
Belted Kingfisher	Martin-pêcheur d'Amérique	G	C
Swamp Sparrow	Bruant des marais	G	I
Lincoln's Sparrow	Bruant de Lincoln	G	I
Song Sparrow	Bruant chanteur	G	I
Common Merganser	Grand Harle	G	C
Black-and-white Warbler	Paruline noir et blanc	B	I
Brown-headed Cowbird	Vacher à tête brune	G	H
Townsend's Solitaire	Solitaire de Townsend	A	I
Great Crested Flycatcher	Tyran huppé	A	I
Clark's Nutcracker	Cassenoix d'Amérique	F	O
Connecticut Warbler	Paruline à gorge grise	G	I
Orange-crowned Warbler	Paruline verdâtre	F	I
Tennessee Warbler	Paruline obscure	F	I
Nashville Warbler	Paruline à joues grises	F	I
Osprey	Balbuzard pêcheur	G	C
Northern Waterthrush	Paruline des ruisseaux	G	I
Savannah Sparrow	Bruant des prés	G	I
Fox Sparrow	Bruant fauve	G	I
Lazuli Bunting	Passerin azuré	G	I
Canada Jay	Mésangeai du Canada	G	O
Cliff Swallow	Hirondelle à front blanc	A	I
Rose-breasted Grosbeak	Cardinal à poitrine rose	G	O

Black-headed Grosbeak	Cardinal à tête noire	F	I
Black-billed Magpie	Pie d'Amérique	G	I
Black-backed Woodpecker	Pic à dos noir	B	I
American Three-toed Woodpecker	Pic à dos rayé	B	I
Downy Woodpecker	Pic mineur	B	I
Hairy Woodpecker	Pic chevelu	B	I
Pine Grosbeak	Durbec des sapins	F	H
Green-tailed Towhee	Tohi à queue verte	G	H
Eastern Towhee	Tohi à flancs roux	G	O
Western Tanager	Piranga à tête rouge	F	I
Red-necked Grebe	Grèbe jougris	G	C
Pied-billed Grebe	Grèbe à bec bigarré	G	C
Black-capped Chickadee	Mésange à tête noire	F	I
Mountain Chickadee	Mésange de Gambel	F	I
Boreal Chickadee	Mésange à tête brune	F	I
Chestnut-backed Chickadee	Mésange à dos marron	F	I
Vesper Sparrow	Bruant vespéral	G	I
Sora	Marouette de Caroline	G	H
Common Grackle	Quiscale bronzé	G	O
Ruby-crowned Kinglet	Roitelet à couronne rubis	F	I
Golden-crowned Kinglet	Roitelet à couronne dorée	F	I
Rock Wren	Troglodyte des rochers	G	I
Eastern Phoebe	Moucherolle phébi	A	I
Ovenbird	Paruline couronnée	G	I
Calliope Hummingbird	Colibri calliope	F	H
Rufous Hummingbird	Colibri roux	F	H
Bay-breasted Warbler	Paruline à poitrine baie	F	I
Yellow-rumped Warbler	Paruline à croupion jaune	F	I
Blackburnian Warbler	Paruline à gorge orangée	F	I
Magnolia Warbler	Paruline à tête cendrée	F	I
Palm Warbler	Paruline à couronne rousse	G	I
Chestnut-sided Warbler	Paruline à flancs marron	F	I
Yellow Warbler	Paruline jaune	F	I
American Redstart	Paruline flamboyante	F	I
Cape May Warbler	Paruline tigrée	F	I
Townsend's Warbler	Paruline de Townsend	F	I
Black-throated Green Warbler	Paruline à gorge noire	F	I
Mountain Bluebird	Merlebleu azuré	A	I
Western Bluebird	Merlebleu de l'Ouest	A	I
Eastern Bluebird	Merlebleu de l'Est	G	I
Red-breasted Nuthatch	Sittelle à poitrine rousse	B	I
White-breasted Nuthatch	Sittelle à poitrine blanche	B	I
Red-naped Sapsucker	Pic à nuque rouge	B	I
Red-breasted Sapsucker	Pic à poitrine rouge	B	I
Williamson's Sapsucker	Pic de Williamson	B	I
Yellow-bellied Sapsucker	Pic maculé	B	I
Pine Siskin	Tarin des pins	F	H
American Goldfinch	Chardonneret jaune	F	H
Clay-colored Sparrow	Bruant des plaines	F	H
Chipping Sparrow	Bruant familier	G	H
Great Gray Owl	Chouette lapone	G	I
Barred Owl	Chouette rayée	G	C
Western Meadowlark	Sturnelle de l'Ouest	G	I

European Starling	Étourneau sansonnet	G	I
Northern Hawk Owl	Chouette épervière	G	C
Tree Swallow	Hirondelle bicolore	A	I
Lesser Yellowlegs	Petit Chevalier	G	I
Greater Yellowlegs	Grand Chevalier	G	I
Solitary Sandpiper	Chevalier solitaire	G	I
House Wren	Troglodyte familier	F	I
Winter Wren	Troglodyte des forêts	G	I
American Robin	Merle d'Amérique	G	I
Eastern Kingbird	Tyran tritri	A	I
Cassin's Vireo	Viréo de Cassin	F	I
Warbling Vireo	Viréo mélodieux	F	I
Red-eyed Vireo	Viréo aux yeux rouges	F	I
Philadelphia Vireo	Viréo de Philadelphie	F	I
Blue-headed Vireo	Viréo à tête bleue	F	I
Yellow-headed Blackbird	Carouge à tête jaune	G	I
Mourning Dove	Tourterelle triste	G	H
White-throated Sparrow	Bruant à gorge blanche	G	H
White-crowned Sparrow	Bruant à couronne blanche	G	I

## Final uniqueness values in non-salvage, salvage and traditionally logged h

Habitat affinity	Nest location	Nest guild	Body Mass (g)	Clutch size (Mean)	SI_Harvest
F	T	NCN	473	4.2	0.993502429
F	T	NCN	183	4.3	0.800583824
O	G	NCN	48.1	4	0.993502429
F	T	CN	131.1	5.67	0.711144773
F	T	CN	166.8	3.25	NA
O	S	NCN	43.8	4	0.737429659
O	G	NCN	19	6	0.987041506
O	G	NCN	13.2	4.51	0.565007609
O	G	NCN	1330	10	0.980625407
O	G	NCN	500	10	0.993502429
O	G	NCN	394	10	2.29E-06
O	G	NCN	545	14	0.961573353
O	G	NCN	1114	8.72	0.924724927
O	G	NCN	1600	14	NA
O	T	NCN	3.4	2	0.987041506
F	T	CN	430	10	0.993502429
O	G	NCN	1500	8	0.987041506
O	G	NCN	900	4	0.942509787
O	T	NCN	32.7	4.5	0.239173332
F	G	NCN	500.1	10.5	0.249411643
O	G	NCN	500	5	0.974251015
O	G	NCN	9000	8	0.582116222
O	T	NCN	1706	2	0.711144773
O	T	CN	600	15	0.974251015
O	T	CN	1300	16	0.919244748
O	T	NCN	1224	2.5	0.61495463
F	T	NCN	437	2.5	0.993502429
F	G	NCN	10.2	4.5	0.801128678
O	G	NCN	7.72	5	5.89E-11
F	G	NCN	32.4	4	0.993502429
O	G	NCN	30.1	4	0.254597362
F	S	NCN	29.6	3.5	0.078597311
F	T	NCN	8.4	5.5	0.760245448
F	T	NCN	18.8	6	NA
O	G	NCN	101.5	4	0.847511413
O	G	NCN	60	3	0.924724927
O	G	NCN	66.5	2	0.980625407
O	G	NCN	53.6	4.1	NA
O	G	NCN	513	4.4	1.57E-15
F	T	NCN	62.6	3	0.553622659
O	T	CN	131.83	5.5	0.283081587
O	T	NCN	31.9	3	1.57E-15
O	T	NCN	12.2	3	0.450457099
O	T	NCN	474	5	0.945635233
O	G	NCN	1055	4.5	0.317146291
F	T	NCN	70.53	4.5	0.792065933
F	T	NCN	128	3.06	NA
F	G	NCN	839	7.1	NA

F	T	CN	270	3.5	0.544874948
O	T	NCN	10.2	4	0.156880258
F	T	CN	10.83	3.5	0.818266088
F	G	NCN	10.3	4	6.10E-11
F	T	NCN	10.3	4	0.241409306
O	S	NCN	11.06	3.6	1.57E-15
O	S	NCN	12.3	3.68	NA
F	T	NCN	64	4.5	0.493520753
O	S	NCN	58.1	4.98	NA
F	G	NCN	424	5	1.37E-06
O	T	NCN	243.6	4.2	0.496751215
O	T	CN	120	4	0.322640091
O	G	NCN	700	10	0.993502429
O	G	NCN	109	4	0.28000026
O	G	NCN	4657	2	0.69859306
F	G	NCN	12.27	3.5	0.271899799
F	S	NCN	10.4	4.12	5.50E-11
O	S	NCN	9.9	4	0.372669828
O	G	NCN	3460	2	0.891476194
F	T	NCN	27.7	4	NA
F	T	NCN	24.6	3.5	0.477687649
F	T	NCN	6300	3	0.803434358
O	T	CN	20	4	0.967917979
O	T	NCN	40	6	NA
F	T	NCN	31.7	3.2	0.19471461
F	G	NCN	20.25	4	0.098904379
O	G	NCN	1200	2	NA
O	G	NCN	700	4	0.707764624
O	G	NCN	300	4	0.900914199
F	T	NCN	26.54	3	0.490328498
F	T	NCN	24.9	4	0.207484117
O	G	NCN	151.6	6	0.929644277
O	G	NCN	17	4.5	0.520948801
O	G	NCN	18.1	4	0.194713551
O	S	NCN	23.4	4.5	0.429013888
O	T	CN	1185	11.5	NA
F	G	NCN	10.7	4.5	0.809592312
O	T	NCN	32	40	0.88409647
O	G	NCN	32.6	4	NA
O	T	CN	40	8	0.980625407
F	T	NCN	129	3.06	NA
F	G	NCN	15.32	4.5	0.591824771
F	G	NCN	9.2	4.6	0.338090391
F	G	NCN	8.9	5	0.001974553
F	G	NCN	8.6	4.5	0.484859891
O	T	NCN	1900	3	0.562297904
F	G	NCN	16.1	4	8.79E-09
O	G	NCN	20	4	NA
F	G	NCN	33.9	3.5	0.496751215
O	S	NCN	14.9	3.13	0.993502429
F	T	NCN	67.6	3	0.248892809
O	G	NCN	24.15	3.48	NA
F	T	NCN	45.1	3.5	0.355625885

F	T	NCN	48.2	3.4	NA
O	T	NCN	25	4	NA
F	T	CN	68	3.5	0.450457099
F	T	CN	52	3.87	0.45340438
F	T	CN	27.64	4.5	0.955375298
F	T	CN	65.96	3.5	0.431391912
O	T	NCN	48.1	4	0.496751215
O	S	NCN	29.4	3.65	NA
O	G	NCN	39.3	4	NA
F	T	NCN	29.8	4	0.360188598
O	G	NCN	1600	8	0.961626189
O	G	NCN	500	10	0.974251015
F	T	CN	11	6.5	0.374372156
F	T	CN	10.78	6.38	NA
F	T	NCN	9.8	6	0.767329029
F	T	CN	9.1	6.2	NA
O	G	NCN	22.95	3.6	NA
O	G	NCN	100	12	0.918713907
O	T	NCN	100.8	4.5	NA
F	T	NCN	6.4	8.5	0.046187079
F	T	NCN	6.1	8.5	0.43598151
O	G	NCN	18	5.6	NA
O	T	CN	20	6	0.961626189
F	G	NCN	21.7	4.5	0.076031225
O	T	NCN	2.85	2	1.57E-15
O	T	NCN	3.58	2	1.57E-15
F	T	NCN	11.1	5.5	0.792993341
F	T	NCN	12.2	3.5	0.011152337
F	T	NCN	10.3	4	0.889239707
F	T	NCN	8.4	4	0.325828437
O	G	NCN	10.3	4.5	0.830079778
O	S	NCN	12	4	0.686098104
O	S	NCN	9.6	4.5	0.809592312
F	T	NCN	8.7	4	0.492468396
F	T	NCN	10.2	6.5	0.850015297
F	T	NCN	8.6	5.7	NA
F	T	NCN	9	3.5	0.735676047
O	T	CN	29.6	5.53	1.57E-15
O	T	CN	27.09	4.78	NA
O	T	CN	30	4.5	NA
F	T	CN	10	5.5	0.273708198
F	T	CN	21.1	7.3	0.974251015
F	T	CN	49.2	4.8	NA
F	T	CN	57.7	4.69	8.47E-14
F	T	CN	47.6	4.38	NA
F	T	CN	50.3	4.5	0.405404256
O	T	NCN	12.91	4	0.121991039
O	S	NCN	11.97	5.5	0.873300581
O	S	NCN	12.2	4	0.318602487
O	S	NCN	12.3	4	0.014711867
F	T	NCN	1267	4.6	0.711144773
F	T	CN	1000	5	0.958342892
O	G	NCN	100	6	NA

O	T	CN	79.46	5.1	NA
F	T	CN	339.8	7	0.993502429
O	T	CN	21.1	5.5	0.393455169
O	G	NCN	85.2	4	0.967917979
O	G	NCN	176	3.7	0.450756915
O	T	NCN	48.4	4	0.840597684
O	T	CN	12	6.8	0.736008058
F	T	CN	8.6	5.5	0.333695929
O	T	NCN	75	3.5	0.331941358
O	T	NCN	41.6	3	0.980625407
F	T	NCN	14.6	3.87	1.78E-09
O	T	NCN	13.98	3.5	0.493520753
F	T	NCN	19.9	3.5	0.068594943
F	T	NCN	12.2	4	0.716403383
F	T	NCN	15.4	4	0.432768827
O	S	NCN	100	4	NA
O	T	NCN	112	2	NA
F	G	NCN	25.1	4	0.002530435
O	G	NCN	25.47	4.58	NA

## abitats

SI_Fire	SI_Salvage	UI_Harvest	UI_Fire	UI_Salvage
0.564829518	NA	0.150651615	0.160069043	NA
0.201033415	0.722245394	0.022199383	0.15385626	0.148619241
0.477519836	NA	0.027871448	0.027871448	NA
NA	NA	0.021885383	NA	NA
NA	0.551665614	NA	NA	0.028028946
0.231662785	0.313663119	0.012994194	0.012994194	0.014153321
NA	0.772256657	0.007611057	NA	0.007816712
0.708312459	0.385021479	0.004435768	0.004435768	0.004894452
NA	NA	0.007954059	NA	NA
NA	NA	0.004224147	NA	NA
NA	NA	0.004224147	NA	NA
NA	NA	0.016565469	NA	NA
0.923499164	0.920653644	0.007954059	0.013049658	0.013908758
NA	0.555178626	NA	NA	0.081936597
0.973820305	NA	0.000914611	0.000914611	NA
NA	NA	0.015452182	NA	NA
NA	NA	0.007981392	NA	NA
0.923499164	NA	0.021526346	0.021526346	NA
0.340662615	0.346750945	0.008721584	0.008721584	0.020075283
0.341480145	0.677454341	0.166460501	0.166460501	0.166765723
0.923499164	NA	0.002710712	0.018796992	NA
0.368865584	0.245543193	0.031766743	0.039748134	0.043611168
NA	NA	0.005668893	NA	NA
0.94832605	NA	0.017467558	0.017467558	NA
0.857014866	0.678657781	0.017467558	0.017467558	0.084280811
0.464400877	0.764247209	0.007766329	0.009675824	0.01048888
NA	NA	0.022199383	NA	NA
0.530802563	0.905552385	0.00220533	0.00220533	0.002396117
0.48527596	0.655366619	0.145378912	0.145378912	0.145641906
NA	NA	0.002682069	NA	NA
0.119422079	0.350758362	0.009017385	0.005141677	0.005450502
0.220446303	0.520630584	0.020875177	0.020875177	0.144868683
0.281126585	0.572437739	0.145948316	0.144026203	0.146270694
1.87E-07	NA	NA	0.018186781	NA
0.899322238	0.771341483	0.00126391	0.00126391	0.001395722
0.766987656	0.96208944	0.005582994	0.005582994	0.005773176
0.521225866	0.690815564	0.005582994	0.005582994	0.005773176
0.289590038	NA	NA	0.00613513	NA
NA	NA	0.002710712	NA	NA
0.449661119	0.856765146	0.144219515	0.018214726	0.144361596
0.343241845	0.491926191	0.062581644	0.010479414	0.032741534
0.011756225	NA	0.004707041	0.004707041	NA
0.230440815	0.302386116	0.006933808	0.006933808	0.007264867
0.718840017	0.759148852	0.153528739	0.029325966	0.032188329
0.309342996	0.437292864	0.00915256	0.00915256	0.009911044
0.746908153	0.793035604	0.006391357	0.006391357	0.00646981
2.45E-09	NA	NA	0.01154438	NA
0.002036658	NA	NA	0.073872526	NA



0.326838588	0.82290299	0.024447263	0.024447263	0.02699686
0.255837043	0.273895971	0.006933808	0.006933808	0.007264867
0.535444358	0.764247209	0.145626433	0.145626433	0.145719208
7.76E-13	NA	0.142857143	0.142857143	NA
0.394120138	0.496795028	0.142857143	0.018186781	0.143048153
2.76E-40	NA	0.145796045	0.002184749	NA
0.0070329	0.984542143	NA	0.002184749	0.144543591
0.73663459	0.660972711	0.017343268	0.017343268	0.058827564
0.045129344	NA	NA	0.017076723	NA
0.941970664	0.551665614	0.166460501	0.162851762	0.166765723
NA	NA	0.154621774	NA	NA
0.36476278	0.939587579	0.150163181	0.068795827	0.073030025
0.923499164	0.96208944	0.005965436	0.013049658	0.013908758
0.370470691	0.525490052	0.00126391	0.00126391	0.001395722
0.695006304	0.617523655	0.041497865	0.041497865	0.043473266
0.23778634	0.35018121	0.007035244	0.007035244	0.00737688
1.14E-38	NA	0.020875177	0.020875177	NA
0.319693316	0.440753091	0.002425261	0.002425261	0.002482157
0.620958233	0.873687614	0.030456146	0.01877445	0.020732431
4.40E-62	NA	NA	0.018214726	NA
0.852850706	0.856765146	0.002094603	0.002094603	0.002117015
NA	NA	0.049188213	NA	NA
NA	NA	0.006588341	NA	NA
NA	0.984542143	NA	NA	0.029980266
7.76E-13	NA	0.017343268	0.017343268	NA
0.136148267	0.208303315	0.003806724	0.003806724	0.004203726
0.544727947	0.558691639	NA	0.011681697	0.011919811
0.973820305	0.96208944	0.00915256	0.00915256	0.009911044
0.923499164	NA	0.015022046	0.015022046	NA
5.79E-71	0.856765146	0.003225474	0.003225474	0.003365825
0.552833552	0.368681133	0.002094603	0.002094603	0.002117015
0.602712764	0.555178626	0.02491695	0.02491695	0.03840166
0.587241513	0.438604993	0.002991305	0.002991305	0.003107234
0.311767984	0.294224354	0.002991305	0.00176975	0.003107234
0.213986345	0.31344408	0.012994194	0.012994194	0.013987548
0.467977064	0.555178626	NA	0.068795827	0.073030025
0.641362324	0.896685657	0.143705599	0.143705599	0.143794085
0.237890675	0.653687114	0.268321737	0.165067824	0.270428539
1.15E-52	NA	NA	0.007274189	NA
NA	NA	0.014624114	NA	NA
5.62E-43	NA	NA	0.142995115	NA
0.261769344	0.711619078	0.002760141	0.002760141	0.006816262
0.486910154	0.939587579	0.00157163	0.00157163	0.001696329
0.200673681	0.409113905	0.002091527	0.002091527	0.002152825
0.52707538	0.605525218	0.00157163	0.00157163	0.001696329
0.353427385	0.612429865	0.005668893	0.009675824	0.01048888
6.46E-07	NA	0.002760141	0.002760141	NA
3.37E-60	NA	NA	0.00176975	NA
NA	NA	0.002682069	NA	NA
0.002036658	0.965706335	0.013152961	0.013152961	0.013987548
0.179656596	0.372406245	0.006391357	0.006391357	0.00646981
4.15E-06	NA	NA	0.007274189	NA
0.775474041	0.781135041	0.009055226	0.009055226	0.00980356

0.004932909	NA	NA	0.010780811	NA
0.289590038	NA	NA	0.011712438	NA
0.226695515	0.483349531	0.000540022	0.000540022	0.000596341
0.4156907	0.632107807	0.002957721	0.002957721	0.003019179
0.496480575	0.925063568	0.010615294	0.010088252	0.01172236
0.353333266	0.592421117	0.000540022	0.000540022	0.000596341
0.0070329	NA	0.008721584	0.008721584	NA
1.20E-06	NA	NA	0.016765069	NA
0.045129344	NA	NA	0.036035945	NA
0.484176739	0.830474	0.009038612	0.009038612	0.009785213
0.973820305	0.955079116	0.028140712	0.028140712	0.030291366
0.899322238	0.92561609	0.018796992	0.018796992	0.030291366
0.278846693	0.812237326	0.144195843	0.000809309	0.144335456
7.35E-45	NA	NA	0.000809309	NA
0.529179582	0.702538481	0.002588968	0.002588968	0.002662938
0.15583873	NA	NA	0.003680367	NA
6.46E-07	0.031085015	NA	0.003943065	0.005723948
0.821131444	0.668382318	0.031828949	0.042018532	0.045616499
0.544727947	0.631626521	NA	0.029325966	0.032188329
0.194237363	0.421869939	0.000851173	0.000851173	0.000939941
0.43063547	0.882824818	0.000851173	0.000851173	0.000939941
0.045129344	NA	NA	0.00514872	NA
NA	0.939587579	0.002828943	NA	0.002927939
0.218473966	0.508046685	0.00617253	0.00617253	0.006816262
0.000589796	NA	0.003128459	0.003128459	NA
6.51E-14	NA	0.000914611	0.000914611	NA
0.512496507	0.954339624	0.004088116	0.004088116	0.004318431
0.113042363	0.386872395	0.001879699	0.001879699	0.001879699
0.973820305	NA	0.002993085	0.002993085	NA
0.282332985	0.737916135	0.000622147	0.000622147	0.00068703
0.814083341	0.922457129	0.004435768	0.004435768	0.004894452
0.354156842	0.734046515	0.00341063	0.00341063	0.003766323
0.284256156	0.65822571	0.002425261	0.002425261	0.002482157
0.455843844	0.905552385	0.000622147	0.000622147	0.00068703
0.945148357	0.958214532	0.002588968	0.002588968	0.002662938
7.08E-10	NA	NA	0.003443631	NA
0.923499164	0.92561609	0.002480752	0.002480752	0.002543436
0.003340567	0.855742604	0.006114193	0.004390542	0.006740078
0.002036658	NA	NA	0.004390542	NA
0.489353985	0.534315694	NA	0.019525032	0.032741534
0.196730989	0.583695136	0.020005214	0.021784461	0.02366429
NA	NA	0.015313082	NA	NA
2.07E-17	NA	NA	0.001519841	NA
NA	NA	0.003147677	NA	NA
0.00031739	NA	NA	0.001429297	NA
0.664484009	0.628235083	0.002957721	0.001429297	0.003019179
0.34884614	0.764689783	0.018356882	0.018356882	0.020075283
0.620324691	0.793035604	0.005976531	0.005976531	0.006011721
0.126710326	0.23336388	0.005976531	0.005976531	0.006011721
0.021519339	0.010803745	0.143001873	0.016765069	0.143016967
0.970551732	0.984542143	0.053307361	0.053307361	0.058827564
NA	NA	0.02665583	NA	NA
NA	0.76297118	NA	NA	0.007810291

8.45E-09	NA	NA	0.010479414	NA
0.419868992	0.665406724	0.015452182	0.163979704	0.028028946
0.242376162	0.351670822	0.002828943	0.006114193	0.002927939
NA	0.886675142	0.003103654	NA	0.003427334
0.804867306	0.589638832	0.009622595	0.009622595	0.010508512
0.650351057	0.92561609	0.00964495	0.00964495	0.010454786
0.154851198	0.079938879	0.145527618	0.145527618	0.145688501
0.246272733	0.58003591	0.145531136	0.145531136	0.145810006
0.191026963	0.32502978	0.00964495	0.00964495	0.010454786
0.708312459	0.544410571	0.004707041	0.004707041	0.024016056
NA	NA	0.001434512	NA	NA
2.74E-33	0.984542143	0.145017464	0.145271736	0.029980266
0.237885939	0.418276509	0.006424659	0.006424659	0.006898652
0.611058781	0.826887966	0.001879699	0.001879699	0.001879699
0.28544884	0.68027066	0.001434512	0.004129727	0.004560415
0.899322238	NA	NA	0.014636229	NA
0.404390272	NA	NA	0.154123613	NA
0.043200345	0.107924505	0.003806724	0.003806724	0.004203726
5.06E-26	0.984542143	NA	0.005141677	0.005450502