

**Implications of climate change for Malaysian tropical montane bird
communities discernible over a 14-years interval.**

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1 **Abstract:** Tropical montane ecosystems are vulnerable to multiple threats, and severe
2 ecological impact on such systems has been documented. However, trends for local montane
3 biodiversity are often varied. Such discrepancy underscores the need to parse the spatial and
4 temporal dynamic of each habitat type within a montane landscape in terms of their species
5 richness, species turnover rate, and relative abundance. We studied species richness and
6 composition of two tropical montane bird communities at two localities in Peninsular
7 Malaysia in 2002-03 and 2016-17. The habitat types sampled at each locality represent a
8 disturbance gradient within a montane landscape. While the number of species generally
9 increased along the disturbance gradient, all study sites (bar tea plantation) had the same or
10 fewer species observed in 2016-17. At the community level observed from the two time
11 periods, Fraser's Hill – where development has been absent since 1920s – had a decrease in
12 species richness; and a higher proportion of species with a decline in their relative abundance,
13 compared to the more disturbed landscape in Cameron Highlands. Both the number of
14 species lost and the number of species gained also varied considerably between the two
15 communities. Our results suggest that climate change is a likely factor in negatively
16 impacting the montane bird communities in Peninsular Malaysia, and highlight the need to
17 monitor the temporal dynamic in the composition of local communities.

18

19 **Keywords:** climate change, community structure, conservation, habitat degradation,
20 Southeast Asia, tropical cloud forests

21

INTRODUCTION

23 The nature conservation of montane habitats in Malaysia was a recurrent theme in the 21st
24 anniversary Special Issue of MNJ (Molesworth Allen, 1961; Watson, 1961; Wyatt-Smith,
25 1961). Subsequently, habitat degradation and climate change have been identified as two
26 major threats to tropical birds (Sekercioglu *et al.*, 2012; Sodhi *et al.* 2004a), with those of
27 montane habitats particularly impacted. For example, the ranges, and population sizes, of
28 montane birds are projected to decrease by 19–42% and 19–62%, respectively, when both
29 habitat loss and climate change are considered together (Harris *et al.*, 2014). These threats
30 could also cause changes in tropical montane bird communities at species level. Lawler *et al.*
31 (2009) reported that hundreds of tropical montane species of restricted range are already
32 threatened by habitat degradation, and are therefore particularly vulnerable to extinction from
33 climate change. Conversely, those that are habitat generalists and have access to suitable
34 habitats spanning a wide elevation range are expected to be less affected (Anciaes &
35 Peterson, 2009).

36 While global biodiversity monitoring of tropical montane birds showed consistent
37 decline (e.g., Sekercioglu *et al.*, 2008), this effect may not translate into responses at
38 community level. Soh *et al.* (2019) reported that there are variable trends in local montane
39 communities, which could be due to species' equivocal or inverse responses to habitats with
40 intermediate level of degradation; citing higher resource availability (e.g., food resources or
41 breeding habitats) typically associated with those lightly disturbed habitats as the main cause
42 of higher species richness. The discrepancy could also be due to the occurrence of more
43 resilient species that are generalists, not threatened, have broad elevational distribution, are
44 introduced, or adaptable to climate change (Soh *et al.*, 2019; Dornelas *et al.*, 2014; Supp and
45 Ernest, 2014; Thomas, 2013). The contradictory findings at global and local community
46 levels underscore the need for better understanding of the effects of habitat degradation on

47 diversity of local communities, and for identifying actual local diversity consequences of the
48 observed climate changes. Despite the conservation importance of tropical montane bird
49 communities, there has been no previous study at this level on the extent and magnitude of
50 current changes in their diversity and distributions – and their differences in temporal trends –
51 in response to the synergistic effects of habitat degradation and climatic change.

52 We studied changes in species richness and composition of two tropical montane bird
53 communities at the landscape level in Peninsular Malaysia, between two time periods, 2002-3
54 and 2016-17. Combining observations from the two surveys, we produce a comprehensive
55 checklist of birds for two montane localities – namely Fraser's Hill and Cameron Highlands.
56 This allowed us to document the persistence of birds across a range of disturbed habitats
57 within these localities; identify species that are vulnerable to habitat degradation (i.e., species
58 that are confined to forests only). We also highlight the conservation value of degraded
59 habitats if they also harbour forest dependent species.

60 Our main questions are: (1) How has species richness of the montane bird community at each
61 locality (alpha diversity) changed over a period of 14 years? (2) Has species turnover (beta
62 diversity, i.e., change in community composition over time) differed between the two
63 localities? (3) Did the relative abundance change between the two time periods?

64 Based on the findings that Southeast Asian bird populations have shifted their
65 elevational distribution upwards, due to climate change (Peh, 2007), and the assumption that
66 community level change would mirror the population level responses to changing climate, we
67 hypothesise that (1) species richness has increased in both localities as some lowland species
68 expanded their upper elevational boundaries; (2) species turnover has been higher in
69 Cameron Highlands, as its landscape is more disturbed; and (3) forest dependent species have
70 become less common, relative to other species in the landscape.

72

MATERIALS AND METHODS

73 *Study region*

74 The montane localities of Fraser's Hill and Cameron Highlands provide a range of habitat
75 types that reflect historical and current land-use in the tropical montane forest areas of
76 Peninsular Malaysia (Figure 1, Table 1). Fraser's Hill is gazetted as a wildlife sanctuary and
77 permanent forest reserve, and has remained relatively free from further development since it
78 was last developed as a holiday respite in the 1920s (Er *et al.*, 2013). In contrast, the
79 protected status of Cameron Highlands was lifted in 1960 to allow timber extraction,
80 agricultural and urban development (Chan, 2006). Since the 1960s, cultivation of temperate
81 fruit and vegetables, and tea, in Cameron Highlands has intensified (Chan, 2006; Peh *et al.*,
82 2011). GIS analyses revealed 2% loss of the forest cover in Cameron Highlands over one
83 decade (Ismail *et al.*, 2014), and an increase in mean annual temperature of 0.9 °C from 1970
84 to 2006 (Ismail *et al.*, 2011, 2014).

85

86 *Bird surveys*

87 We first surveyed our study sites in 2002 to 2003 (Soh *et al.*, 2006) and then resurveyed the
88 same sites 14 years later (Table 1). Bird occurrence and abundance surveys were conducted
89 over six periods within each sampling year in 2002-3 and 2016-17 (Table 2). During each
90 sampling period, six 10-minute point counts were conducted at each site (i.e., 36 point-counts
91 per habitat type: primary forest, secondary forest, edge forest, small fragment, tea plantation,
92 rural and urban areas), except for the fragment in Fraser's Hill where only three point counts
93 were conducted due to the small area (totaling 18 points). All points were spaced at least 300
94 m apart to ensure that observations were independent (Ralph *et al.*, 1993). All birds seen or
95 heard within a 25 m radius from the centre of each point over a 10-minute period were
96 recorded, but birds flying overhead were excluded (*sensu* Soh *et al.*, 2006). To maximize
97 detections, point counts were conducted between 0700 and 1100 hours on fair weather days

98 (i.e., no heavy rain). All point counts were conducted by M.C.K. Soh along forest trails and
99 along roads in rural and urban areas. Random sampling in the forested sites away from the
100 forest trails was deemed unsafe due to the steep terrain. Unfamiliar calls were recorded with
101 digital audio recorders (Olympus models DW-90 in 2002-3 and LS-14 in 2016-7) and later
102 identified to species by consulting expert ornithologists.

103

104 *Data analysis*

105 All statistical analyses were performed using R 3.2.2 (R Core Team, 2017). We determined if
106 species richness of each habitat type, as well as total species richness at each locality (i.e.,
107 landscape-scale), differ over time by comparing the numbers of species observed between the
108 two time periods (2002-3 and 2016-17).

109 To estimate the total species turnover of each habitat type and landscape, we divided
110 their sum of species gained and species lost by total species observed in both time periods, to
111 derive the proportion of species that differed between the two time periods (Diamond, 1969).
112 Since total species turnover incorporates both species that appeared (i.e., species gained) and
113 disappeared (i.e., species lost), we also report the proportion of species that appeared in 2016-
114 17 and that of species that disappeared in 2016-17, relative to the total number of species
115 observed in both time periods, in order to determine their relative contribution.

116 To determine if the relative abundance of each species differed between the two time
117 periods, we conducted Bayesian analysis using the “Bbinom” (Bayesian binomial simulation)
118 function from the “wqid” package (Meredith, 2017) to compare two sets of binomial data
119 (focal species or not). Our approach involved a sample of 50,000 Markov Chain Monte Carlo
120 [MCMC] simulations from the posterior for a binomial likelihood (i.e., probability of an
121 individual being the focal species being lower or higher in 2016-17 compared to 2002-3); and
122 we used a uniform prior in our analyses, instead of an informative prior, because we did not
123 have any prior information about the species’ relative abundance at landscape level. For each

124 species, we compared binomial likelihood of its identification between the two time periods
125 to deduce if it had increased or decreased (probability cut-off at 90%) in terms of relative
126 abundance.

127

128 **RESULTS AND DISCUSSION**

129 We observed 68 species at Fraser's Hill (60 in 2002-3; 57 in 2016-17) and 76 species at
130 Cameron Highlands (64 in 2002-3; 66 in 2016-17). In total, we recorded 80 species from both
131 localities in both time points (see checklist in Supporting Information Appendix 1). In
132 Fraser's Hill, the total number of species observed in both time points in primary forest,
133 forest fragment and forest edge were 39, 43 and 54, respectively. The total number of species
134 observed in both primary forest (48) and secondary forest (48) in Cameron Highlands fell
135 within that range. Both rural habitat in Fraser's Hill and Cameron Highlands had 52 and 47
136 species culminated from both time points, respectively. In comparison, we observed fewer
137 total species number in tea plantation (38) and urban habitat (20) in Cameron Highlands.

138

139 **Changes in species richness along disturbance gradient and between two time periods**

140 In general, the number of species increased along disturbance gradient in Fraser's Hill.
141 However, the most disturbed sites in Cameron Highlands (tea and urban) had the least
142 number of species (Fig. 2a). All sites had either the same number or fewer species observed
143 in 2016-17, with the exception of tea plantation (Fig. 2a). At landscape-scale, fewer species
144 were observed in Fraser's Hill as compared to Cameron Highlands/ This, was mainly due to
145 an increase in the number of species in tea plantation at the latter location (Fig. 2a).

146 Our results generally do not support our hypothesis that species richness increased in
147 Fraser's Hill and Cameron Highlands from first to second survey. While we reasoned that
148 global warming would have encouraged species with lowland affinities to expand their

149 vertical distribution, this increase was apparently negated by a greater loss in montane
150 specialists due to contractions in their attitudinal ranges. Upwards shifts and narrowing
151 vertical distributions are well documented for Neotropical montane birds and are a
152 consequence of a reduction in their climatic niches (e.g.. Bender *et al.*, 2019; Forero-Medina
153 *et al.*, 2011; Neate-Clegg *et al.*, 2018). More worryingly, higher altitudinal specialists are also
154 at risk of local extirpations in some mountain tops (Freeman *et al.*, 2018). In Peninsular
155 Malaysia uplands, such vulnerable species include upper-montane specialists such as Rufous-
156 bellied niltava (*Niltava sundara*) and Chestnut-tailed minla (*Actinodura strigula*) which
157 probably deserve closer conservation attention.

158 Since we were careful to repeat the sampling protocol in the same localities and the
159 same observer (M.C.K. Soh) conducted the surveys in both periods, we doubt that the
160 observed phenomenon was an artefact of sampling error. Further, the decline in species
161 richness was a consistent trend across all habitat types, except for the tea plantation. Any
162 climatic changes are expected to be slight, since the lapse between our sampling periods was
163 only 14 years. Thus, it is not surprising that declines in species richness in each habitat type
164 were incremental. Nonetheless, if the current climatic trends remain, species declines are
165 likely to progress over a protracted period. Further, our sampled sites remained relatively
166 unchanged in terms of level of disturbance since the survey in 2002-3 was conducted.

167 On a more positive note, the increase in species richness at the tea plantation is
168 encouraging and demonstrates the potential conservation value of such estates -- assuming
169 certain landscape features such as remnant pockets of forest, native riparian vegetation along
170 streams for irrigation, and natural hedges along roads remain prominent. Aside from these
171 features, the closeness to forest habitats may also encourage more edge species to venture
172 into the tea plantation to forage (Barlow *et al.*, 2007; Lucey & Hill, 2012). Some species may
173 also use small forest remnants as 'stepping stones' to move between forest patches (Baum *et*

174 *al.*, 2004; Uezu *et al.*, 2008; Saura *et al.*, 2014). Thus, by maintaining natural corridors to
175 facilitate movement and remnant forest patches, well-managed plantations that are
176 biodiversity friendly can still contribute towards the conservation of the montane ecosystem.

177

178 **Species turnover rates between two time periods**

179 The primary forests of Fraser's Hill and urban areas of Cameron Highlands had the lowest
180 proportion of species that appeared in 2016-17 relative to the total number of
181 species observed in both time points (Fig. 2b). All disturbed sites in Cameron Highlands (bar
182 urban areas) had the lowest proportion of species that disappeared in 2016-17 relative to the
183 total number of species observed in both time points (Fig. 2b). The edge forests in Fraser's
184 Hill and tea plantation in Cameron Highlands had the highest total turnover (Fig. 2b). There
185 was no difference in total species turnover at both Fraser's Hill and Cameron Highlands, but
186 Fraser's Hill had a higher proportion of species lost and Cameron Highlands had a higher
187 proportion of species gained (Fig. 2b).

188 Our hypothesis that species turnover would be higher in Cameron Highlands since it
189 was more disturbed was not supported; rather the species turnovers in both Fraser's Hill and
190 Cameron Highlands were similar. However, the gains did exceed losses in Cameron
191 Highlands, primarily due to a greater number of new species detected in the more recent
192 survey at the tea plantation for reasons explained earlier. Conversely, the losses were higher
193 than gains in Fraser's Hill, which we postulate is a likely consequence of progressively
194 warmer climate. The higher turnover losses were unlikely to be caused by increased habitat
195 degradation in Fraser's Hill, since the landscape cover remained largely unchanged during
196 the interval of 14 years (Er *et al.*, 2013). Fraser's Hill is still protected and, unlike Cameron
197 Highlands, its steep terrain is generally unsuitable for extensive agriculture (Chan, 2006;
198 Ismail *et al.*, 2011, 2014). Thus, the absence of previously detected species in Fraser's Hill

199 was probably due to more species retreating to higher elevations. That said, the higher species
200 turnover losses seen in most habitat types in Cameron Highlands could also be attributed to a
201 warmer climate, compounded by continual deforestation and habitat degradation.

202

203 **Changes in relative abundance between two time periods**

204 The rural areas at Fraser's Hill, and tea plantation and urban areas at Cameron Highlands had
205 the highest proportion of species with an increase in relative abundance (21-25%; Fig. 2c).

206 The secondary forest and rural areas at Cameron Highlands had the highest proportion of
207 species with a decrease in relative abundance (19-21%; Fig. 2c). Generally, there was a
208 decline along the disturbance gradient in the proportion of species that had no change in their
209 relative abundance, though the trend is less pronounced at Cameron Highlands (Fig. 2c).

210 There was no difference between Fraser's Hill and Cameron Highlands at landscape-scale in
211 terms of the proportion of species that had no change in their relative abundance. However,
212 Fraser's Hill had a higher proportion of species with a decrease in relative abundance, and
213 Cameron Highlands had a higher proportion of species with an increase in relative abundance
214 (Fig. 2c).

215 The increased bird abundances in the tea plantation and urban areas at Cameron
216 Highlands indicate that some species probably thrive in human modified habitats. Many such
217 species are affiliated to lowland habitats: marked increases in abundance were observed for
218 Black-naped oriole (*Oriolus chinensis*), Common myna (*Acridotheres tristis*), Eurasian tree
219 sparrow (*Passer montanus*), Large-billed crow (*Corvus macrohynchos*), Oriental magpie-
220 robin (*Copsychus saularis*), and Spotted dove (*Spilopelia chinensis*). The results suggest that
221 the species composition in highly developed areas in montane environments are not unlike
222 those at lower altitudes, demonstrating the effects of biotic homogenisation, whereby a few
223 highly adaptable species dominate the community (Lever, 1987; Marzluff, 2001; Soh *et al.*,

224 2006). Yet, the abundance of some montane species also increased in the tea plantation,
225 including Blue-winged minla (*Actinodura cyanouroptera*), Fire-breasted flowerpecker
226 (*Dicaeum ignipectus*), Long-tailed sibia (*Heterophasia picaoides*), Mountain bulbul (*Ixos*
227 *mcclellandii*) and Silver-eared mesia (*Leiothrix argentauris*). This result implies that the tea
228 plantation not only attracted new bird species but also more individuals; thus, providing
229 further evidence of its increased conservation value since the 2002-3 surveys. However, the
230 preservation of large tracts of contiguous montane forests is still a priority as the montane
231 birds that are utilising in the tea plantation are all more adaptable edge species (Robson,
232 2008).

233 Apart from the observed decline in species richness and greater species turnover
234 losses, the larger number of bird species with a reduction in abundance at Fraser's Hill may
235 additionally signal an effect of global warming. This result reiterates the need to monitor to
236 the population dynamics of montane specialists over the long-term, in order to better
237 comprehend their responses to climate change.

238

239 **Limitations and future research**

240 Our results provide a preliminary analysis of the changes in the montane bird community in
241 Peninsular Malaysia which include alpha and beta diversities, and relative abundance. We did
242 not correct for imperfect detection, which can arise from imperfect sampling design and
243 environment constraints (e.g., not detecting a species behind an observer, or dense vegetation
244 obstructing a clear line of sight) (MacKenzie *et al.*, 2002). Estimating detection probabilities
245 to correct for species occupancy probabilities and abundance estimates can be done, using
246 multispecies occupancy modelling or N-mixture models respectively if their assumptions
247 such as sampling closure are met (Kéry & Royle, 2014).

248 We may also have missed cryptic and/or rare species during our surveys. Such
249 sampling deficiencies can be supplemented by deploying autonomous sound recorders. These
250 may be more successful than traditional sampling by point counts and line transects in
251 detecting species that tend to avoid human observers, and can be scheduled to record for
252 much longer periods (Tegeler *et al.*, 2012; Zwart *et al.*, 2015).

253 Our results are also indicative of the responses of birds, a group relatively more
254 mobile than other taxa that may be more vulnerable to disturbances such as amphibians and
255 reptiles (Hopkins, 2007; Bishop *et al.*, 2012). Apart from deforestation and land-use
256 conversion, the montane landscape at Cameron Highlands is increasingly fragmented; this
257 may compromise gene flow in some isolated populations (Habel *et al.* 2014; Husemann *et al.*,
258 2015). Studies for certain species vulnerable to such impacts can help inform future land-use
259 planners if the preservation of natural corridors to encourage greater gene flow may be
260 needed.

261

262 CONCLUSION

263 Our results suggest that climate change was a likely factor in negatively impacting two
264 montane bird communities in Peninsular Malaysia. This was more clearly demonstrated at
265 our Fraser's Hill sampling sites, where community-specific changes were observed without
266 further habitat degradation since 2002-3. An increase in species richness, in the tea plantation
267 suggest that agricultural landscapes can increase in conservation value if interspersed with
268 remnants and/or corridors of native vegetation. The relatively high species turnover in local
269 montane bird communities in our study suggests the need to monitor the temporal dynamics
270 in the composition of local communities. While our study indicates the effect of climate
271 change and habitat degradation on montane bird communities, more research to investigate

272 the impacts on other taxa, and population genomics of vulnerable species is crucial to better
273 comprehend such responses to environmental change.

274

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Table 1: Description and elevation of habitat types across two montane localities in Peninsular Malaysia surveyed in 2002 and 2017.

Habitat type and locality/localities	Elevation (m) of each locality as listed in previous column	Description
Primary forest CH, FH	1621, 1280	Primary montane forest
Secondary forest CH	1506	Exotic softwood plantation ~2 ha surrounded by roads and native forests left to regenerate more than 50 years ago
Tea CH	1591	Mature tea plantation
Rural CH, FH	1477, 1243	Roadsides beyond the perimeter of town centres and flanked by vegetation
Urban CH	1475	Roads within the town centres which are mostly devoid of vegetation
Edge forest FH	1236	Part of continuous montane forest 100–150 m away from edge of road
Small fragment FH	1260	Small forest patch <5 ha isolated by narrow roads and golf course

CH = Cameron Highlands, FH = Fraser's Hill

Table 2: Dates that point counts were conducted

2002-3	2016-7
26 th July to 29 th September 2002	21 st May to 15 th June 2016
12 th October to 8 th November 2002	30 th July to 23rd August 2016
23 rd November to 9 th December 2002	30 th October to 25 th November 2016
17 th February 2003 to 23 rd March 2003	27 th December 2016 to 22 nd January 2017
5 th May 2003 to 24 th June 2003	14 th March to 7 th April 2017
1 st July 2003 to 13 th August 2003	9 th May to 2 nd June 2017

Figure legends

Figure 1: Map of Peninsular Malaysia showing two study localities and sites along the Main Range.

Figure 2: (a) Species richness; (b) species turnover; and (c) relative abundance of all habitat types and landscapes at Fraser's Hill and Cameron Highlands.

Figure 1

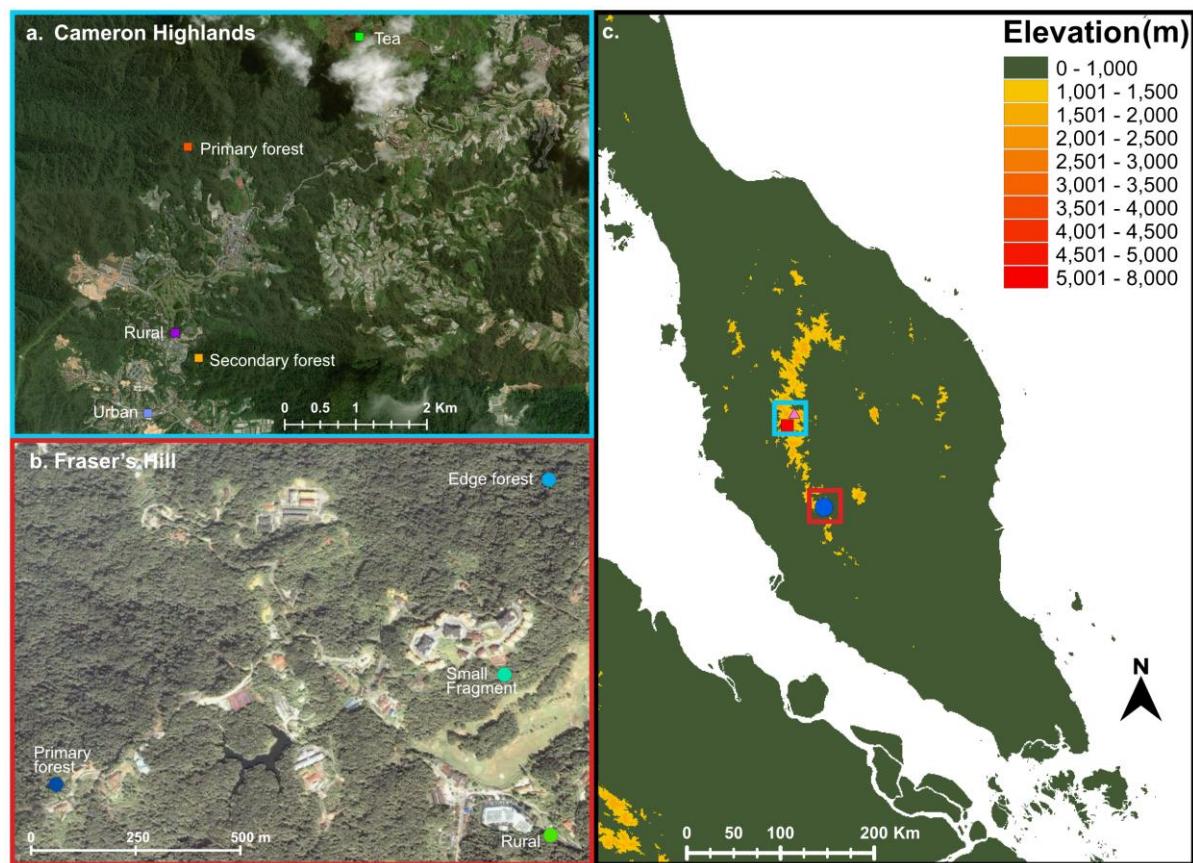
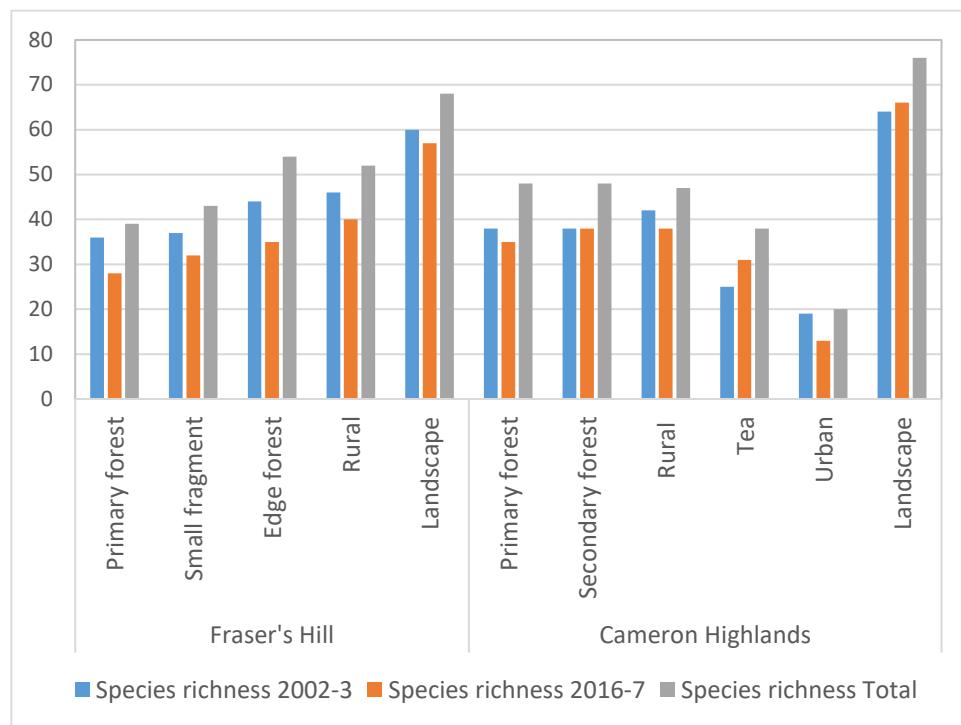
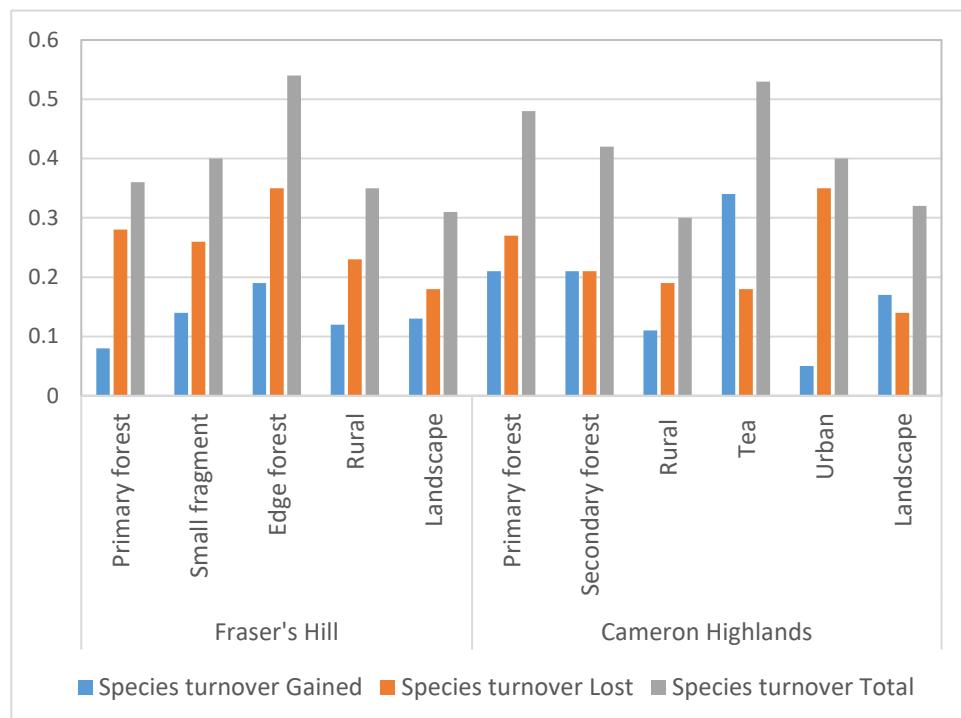


Figure 2

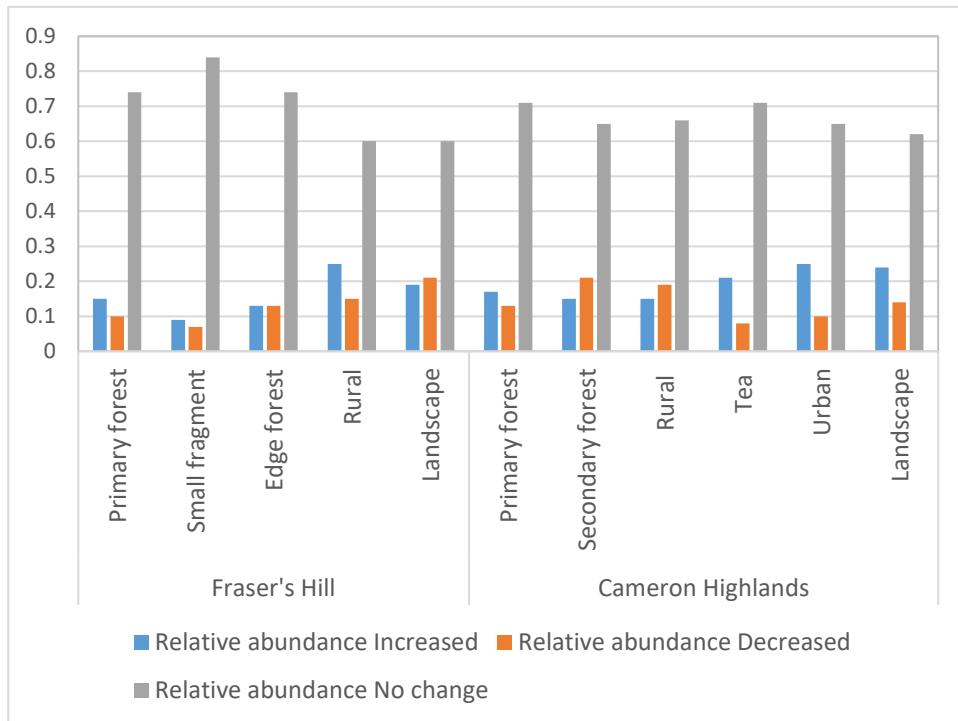
(a)



(b)



(c)



APPENDIX

Appendix 1: Species checklist of resident birds detected in 2002-3 and 2016-7. Acronyms indicate the study region and habitat type, followed by the year of survey, i.e., 02 refers to 2002-3 and 16 refers to 2016-7. FP – Fraser’s Hill primary forest, FE – Fraser’s Hill edge forest, FF – Fraser’s hill small fragment forest, FR – Fraser’s Hill rural areas, CP – Cameron Highlands primary forest, CS – Cameron Highlands secondary forest, CT – Cameron Highlands tea plantation, CR – Cameron Highlands rural areas, and CU – Cameron Highlands urban areas. Common and scientific names follow IOC world bird list (Gill *et al.*, 2020).

1. Ashy Bulbul; *Hemixos flavala*; FR16; CS02
2. Black Laughingthrush; *Melanocichla lugubris*; FE02
3. Blue Nuthatch; *Sitta azurea*; FF02; FR02; CP02 & CP16; CS02
4. Bay Woodpecker; *Blythipicus pyrrhotis*; CP16
5. Buff-breasted Babbler; *Pellorneum tickelli*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16
6. Black-browed Barbet; *Psilopogon oorti*; FP02 & FP16; FE02 & FE16; FR16
7. Black-crested Bulbul; *Rubigula flaviventris*; CR16
8. Black-and-Crimson Oriole; *Oriolus cruentus*; FP02; FE02 & FE16; FF02 & FF16; FR02 & FR16; CS02 & CS16; CR02
9. Black-eared Shrike-babbler; *Pteruthius melanotis*; FP02; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02; CS02 & CS16; CR02 & CR16
10. Blyth's Hawk-Eagle; *Nisaetus alboniger*; CS16
11. Black-naped Oriole; *Oriolus chinensis*; CT16; CR02
12. Blyth's Shrike-Babbler; *Pteruthius aeralatus*; FP02 & FP16; FE02 & FE16; FF02; FR02; CP02; CS02 & CS16; CR02
13. Bar-throated Minla; *Actinodura strigula*; CP02
14. Black-throated Sunbird; *Aethopyga saturata*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16; CU02 & CU16

15. Blue-winged Minla; *Actinodura cyanouroptera*; FP02 & FP16; FE02; FF02; CP02; CS16; CT16; CR02
16. Bar-wing Flycatcher-shrike; *Hemipus picatus*; FE16; FR02 & FR16; CU16
17. Common Myna; *Acridotheres tristis*; CR02 & CR16; CU02 & CU16
18. Collared Owlet; *Glaucidium brodiei*; FP02 & FP16; FE16; FF02; FR16; CP02 & CP16
19. Common Tailorbird; *Orthotomus sutorius*; FR02 & FR16; CT02 & CT16; CR02 & CR16; CU02 & CU16
20. Chestnut-capped Laughingthrush; *Pterorhinus mitratus*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CR02 & CR16; CU02
21. Chestnut-crowned Warbler; *Phylloscopus castaniceps*; FE02; FF02; FR02; CP02
22. Common Green Magpie; *Cissa chinensis*; FE02; FF16; FR02 & FR16; CP16; CS02
23. Crested Serpent Eagle; *Spilornis cheela*; CP02; CS02; CT02
24. Diard's Trogan; *Harpactes diardii*; CP16
25. Dark-necked Tailorbird; *Orthotomus atrogularis*; FE02; FR02 & FR16; CT02; CR02
26. Everett's White-eye; *Zosterops everetti*; FP16; CS02; CR02 & CR16; CU02 & CU16
27. Eurasian Tree Sparrow; *Passer montanus*; CT16; CR02 & CR16; CU02
28. Fire-breasted Flowerpecker; *Dicaeum ignipectus*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16; CU02
29. Fire-tufted Barbet; *Psilopogon pyrolophus*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16
30. Golden Babbler; *Cyanoderma chrysaeum*; FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CR02 & CR16
31. Greater Yellownape; *Chrysophlegma flavinucha*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16

32. Green-billed Malkoha; *Phaenicophaeus tristis*; FR02 & FR16; CT16; CR02 & CR16; CU02

33. Grey-chinned Minivet; *Pericrocotus solaris*; FP02; FE02 & FE16; FF02 & FF16; CP02

34. Grey-headed Canary-flycatcher; *Culicicapa ceylonensis*; FP02; CT02 & CT16

35. Grey-throated Babbler; *Stachyris nigriceps*; FP02; FE02 & FE16; FF02; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02

36. Golden-throated Barbet; *Psilopogon franklinii*; FP02; CP16; CS16; CR02

37. Javan Myna; *Acridotheres javanicus*; CU02

38. Large Cuckooshrike; *Coracina macei*; FE16; FR16; CP16; CS16; CR16

39. Large Niltava; *Niltava grandis*; FP02; FE02; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CR02 & CR16

40. Lesser Shortwing; *Brachypteryx leucophrys*; FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS16; CT02 & CT16; CR02 & CR16

41. Little Spiderhunter; *Arachnothera longirostra*; FP02 & FP16; FE02; FR02 & FR16

42. Lesser Yellownape; *Picus chlorolophus*; FE02 & FE16; FF02

43. Large-billed Crow; *Corvus macrorhynchos*; FF02 & FF16; FR02; CP02; CS02 & CS16; CT02 & CT16; CR02 & CR16; CU02 & CU16

44. Long-billed Spiderhunter; *Arachnothera robusta*; CP02

45. Little Cuckoo-dove; *Macropygia ruficeps*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16

46. Large Cuckooshrike; *Coracina macei*; FP02; FE02; FF02; FR02; CP02; CS02

47. Large Hawk-Cuckoo; *Hierococcyx sparverioides*; FP02; FE02; FR02; CP16

48. Little Pied Flycatcher; *Ficedula westermanni*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02; CR02 & CR16

49. Lesser Racket-tailed Drongo; *Dicrurus remifer*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CR02 & CR16

50. Large Scimitar Babbler; *Erythrogenys hypoleucos*; FE02; FF16; FR02; CP16

51. Long-tailed Broadbill; *Psarisomus dalhousiae*; FE02 & FE16; FR02 & FR16

52. Long-tailed Shrike; *Lanius schach*; CT16

53. Long-tailed Sibia; *Heterophasia picaoides*; FP02 & FP16; FE02; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT16; CR02 & CR16

54. Mountain Bulbul; *Ixos mcclellandii*; FP02 & FP16; FE16; FF02 & FF16; FR02 & FR16; CP02; CS02 & CS16; CT16; CR02 & CR16; CU02 & CU16

55. Mountain Fulvetta; *Alcippe peracensis*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16

56. Malayan Laughingthrush; *Trochalopteron peninsulae*; FP02 & FP16; FE16; FR02; CP02 & CP16; CS02 & CS16; CT02

57. Malaysian Partridge; *Arborophila campbelli*; CP16; CS16

58. Mountain Tailorbird; *Phyllergates cucullatus*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16

59. Maroon Woodpecker; *Blythipicus rubiginosus*; FE02

60. Mountain Imperial Pigeon; *Ducula badia*; FP02 & FP16; FE02; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02; CR02 & CR16

61. Mountain Leaf-warbler; *Phylloscopus trivirgatus*; FP02; FE02 & FE16; FF02 & FF16; CP02 & CP16; CS02 & CS16; CT02; CR02 & CR16

62. Ochraceous Bulbul; *Alophoixus ochraceus*; FE16

63. Oriental Cuckoo; *Cuculus optatus*; FR02

64. Indian White-eye; *Zosterops palpebrosus*; CT16

65. Orange-bellied Leafbird; *Chloropsis hardwickii*; FP02; FE02; FR02 & FR16; CP02; CS16; CR02 & CR16

66. Oriental Magpie-Robin; *Copsychus saularis*; FF02; FR02 & FR16; CS16; CT02 & CT16; CR02 & CR16; CU02 & CU16

67. Pacific Swallow; *Hirundo tahitica*; CT16; CR02 & CR16; CU02 & CU16

68. Pale Blue Flycatcher; *Cyornis unicolor*; CR02 & CR16

69. Pygmy Flycatcher; *Ficedula hodgsoni*; FE02; CP02; CS02 & CS16; CT16; CR02 & CR16

70. Rock Dove; *Columba livia*; CR02 & CR16; CU02 & CU16

71. Rhinoceros Hornbill; *Buceros rhinoceros*; FE16

72. Rufous-browed Flycatcher; *Anthipes solitaris*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP16; CS02 & CS16; CR02 & CR16

73. Red-headed Trogon; *Harpactes erythrocephalus*; FE16; CS02

74. Rusty-naped Pitta; *Pitta oatesi*; FE16

75. Red-whiskered Bulbul; *Pycnonotus jocosus*; CR02 & CR16

76. Spotted Dove; *Spilopelia chinensis*; FF16; CT02; CR02 & CR16; CU02 & CU16

77. Streaked Spiderhunter; *Arachnothera magna*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16; CU02 & CU16

78. Snowy-browed Flycatcher; *Ficedula hyperythra*; FE16; FR02; CS16

79. Slaty-backed Forktail; *Enicurus schistaceus*; FF16; FR02; CS02 & CS16; CT16

80. Scaly-breasted Munia; *Lonchura punctulata*; FR02 & FR16; CT16; CR02 & CR16; CU02

81. Silver-eared Mesia; *Leiothrix argentauris*; FP02 & FP16; FE02; FF02 & FF16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16

82. Stripe-throated Bulbul; *Pycnonotus finlaysoni*; FF16; FR02 & FR16; CT02 & CT16

83. Streaked Wren-babbler; *Gypsophila brevicaudata*; FP02 & FP16; FE02 & FE16; FF16; FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16

84. Verditer Flycatcher; *Eumyias thalassinus*; FF02; FR16

85. White-rumped Munia; *Lonchura striata*; FR02

86. White-throated Fantail; *Rhipidura albicollis*; FP02 & FP16; FE02 & FE16; FF02 & FF16; FR02 & FR16; CP02 & CP16; CS02 & CS16; CT02 & CT16; CR02 & CR16; CU02

87. White-tailed Robin; *Myiomela leucura*; FE02; FF16; CP16; CS16; CT02; CR02 & CR16

88. Yellow-bellied Prinia; *Prinia flaviventris*; CT16

89. Yellow-breasted Warbler; *Phylloscopus montis*; FE02; FF02; CP02 & CP16; CS02

90. Yellow-vented Bulbul; *Pycnonotus goiavier*; FR02 & FR16; CS02; CT02 & CT16; CR02 & CR16; CU02 & CU16