



Title	Efficacy of aural detection methods for detecting Northern Pika (<i>Ochotona hyperborea</i>) occupancy in rocky and densely vegetated habitats
Author(s)	Sakiyama, Tomoki; Molinos, Jorge Garcia
Citation	Journal of mammalogy https://doi.org/10.1093/jmammal/gyad066
Issue Date	2023-06-24
Doc URL	http://hdl.handle.net/2115/92673
Rights	This is a pre-copyedited, author-produced version of an article accepted for publication in Journal of mammalogy following peer review. The version of record is available online at: 10.1093/jmammal/gyad066 .
Type	article (author version)
Additional Information	There are other files related to this item in HUSCAP. Check the above URL.
File Information	108070.pdf



[Instructions for use](#)

1 **Running header:** Aural detection methods to survey pikas

2

3 **Efficacy of aural detection methods for detecting Northern Pika (*Ochotona hyperborea*)**
4 **occupancy in rocky and densely vegetated habitats**

5

6

7

8 Tomoki Sakiyama* ¹, and Jorge García Molinos²

9

10 ¹ *Graduate School of Environmental Science, Hokkaido University, Kita-10, Nishi-5, Kita-ku,*
11 *Sapporo, Hokkaido, 060-0810, Japan*

12

13 ² *Arctic Research Center, Hokkaido University, Kita-21, Nishi-11, Kita-ku, Sapporo,*
14 *Hokkaido, 001-0021, Japan*

15

16 **To whom correspondence should be addressed: wisdom.tree.1994@gmail.com*

17

18

19 **Teaser text:** Pikas are considered vulnerable to warming climates, but detecting their
20 presence in rocky patches covered by dense vegetation is challenging. Here, we assessed the
21 efficacy of aural detection methods, namely auditory observation and playback, to survey
22 Northern Pikas in Japan.

23

24 Wildlife monitoring plays a key role in species conservation, with growing importance under
25 the threat of climate change. The Northern Pika (*Ochotona hyperborea*) is a cold-adapted
26 species found in Hokkaido, Japan, presumably vulnerable to such threats. However, its high
27 elusiveness hinders detailed population surveys, and visual detection methods—often used for
28 other pika species—are not applicable to its densely vegetated habitats. In this study, we
29 assessed the efficacy of aural detection methods to survey the occupancy of Northern Pika
30 through their distinct vocalizations. We conducted two types of point-count surveys—
31 auditory observation and playback—during 2021 at 18 sites in and around Daisetsuzan
32 National Park. We then assessed the efficacy of these methods in detecting presence and
33 compared time until first detection of the animal. The Northern Pika was present at 11 of all
34 surveyed sites, with a predominance at higher elevations. Our results suggest that both
35 auditory observation and playback are effective at detecting presence, but playback is more
36 time-efficient. We discuss the advantages and disadvantages of each method given these
37 results. In conclusion, our results demonstrate that both survey methods are applicable for
38 Northern Pika population surveys—even in densely vegetated habitats—representing valid
39 and affordable survey methods that can help to improve current monitoring and conservation
40 efforts, and will be of increasing value given potentially negative effects of climate change on
41 persistence of the species.

42

43 Key words: aural detection, auditory observation, Northern Pika, playback, wildlife

44 monitoring

45

46 Cumulative evidence suggests that wildlife populations are declining globally, and monitoring
47 has been critical in detecting such changes (Collen et al. 2009; Almond et al. 2020).
48 Understanding the drivers of these declines poses a significant challenge informing
49 conservation policy and managing wildlife populations adaptively under global environmental
50 changes (Yoccoz et al. 2001; Di Fonzo et al. 2016). However, some species are more elusive
51 to detect than others (Thompson 2004), making monitoring surveys of rare and elusive
52 species prone to inaccurate detections and hindering our ability to obtain unbiased estimates
53 of occupancy and abundance.

54 Pikas are small lagomorphs found at high latitudes or altitudes in Asia and North
55 America. Because of their adaptation to cold environments (Yang et al. 2008; Wang et al.
56 2020), they are considered to be vulnerable to warming climates (Beever et al. 2011; Billman
57 et al. 2021). Therefore, efforts to monitor the status of pika populations are needed to better
58 understand how climate change affects population distribution and abundance. However,
59 some pika species are highly elusive given their preference to live in rocky patches (Henry
60 and Russello 2011), and while research exists for the American Pika (*Ochotona princeps*), it
61 is lacking for many of other rock-dwelling species. This is true for the Northern Pika
62 (*Ochotona hyperborea*), a species found in northeastern Eurasia that has the widest
63 distribution among all pika species (Smith et al. 2018). Although a recent study in the eastern
64 subspecies *O. h. yesoensis* in Hokkaido, Japan suggested its vulnerability to climate change
65 due to thermal constraints (Sakiyama et al. 2021), the effects of climate change on local
66 populations remain largely unknown. An absence of population studies on the Northern Pika
67 in Hokkaido is not only due to its habitat preference for rocky patches, but also to the dense
68 vegetation that covers their rocky habitats. Moreover, most habitats are located on steep
69 slopes, making accessibility and site navigation extremely difficult and time-consuming.

70 Therefore, establishing robust yet affordable methods to assess the occupancy of Northern
71 Pikas in Hokkaido and localities with similar habitat characteristics are needed.

72 Previous studies have surveyed the occurrence of rock-dwelling pikas either by
73 detecting sign of animal presence visually or aurally, or indirectly through presence of feces
74 and haypiles (Beever et al. 2010; Erb et al. 2011; Stewart et al. 2015; Hall et al. 2016;
75 Korolev 2017). Such multiple-evidence approaches likely aid in detecting the presence of this
76 elusive animal, but the major assumption of the survey (although not mentioned) is that the
77 visibility and navigability of the study sites allow for detection of these signs. This
78 assumption, however, does not apply to the environments found in Hokkaido because of the
79 aforementioned difficulties.

80 Here, we advocate instead for the use of survey methods based solely on aural detection
81 as an affordable and reliable alternative, and test their utility in the Hokkaido locality.
82 Northern Pikas are highly vocal, and their vocalizations are detectable from distant locations
83 (up to 200 m; Kawamichi 1969) even when the vocalizing individual is not visible
84 (Kawamichi 1970). Moreover, vocalizations are heard frequently during and after the
85 reproductive season (Kawamichi 1971), permitting population surveys in the snow-free
86 season (i.e., summer) even though their habitats—typically located in remote settings such as
87 deep forests and alpines—are inaccessible at other times of the year. Previous efforts have
88 been made to survey a Northern Pika population based on their vocalizations (Korolev 2017),
89 but that survey was conducted only at sites of known presence since the study focused on
90 abundance estimation. Occupancy surveys, in contrast, aim to establish whether the Northern
91 Pika is present or absent at a given site.

92 We conducted field experiments to evaluate the efficacy of two aural detection
93 methods—auditory observation and playback—for establishing occupancy of Northern Pikas
94 in Hokkaido. Auditory observation refers to a passive survey that involves listening to

95 naturally occurring vocalizations and is the most commonly used aural detection method for
96 pikas (Billman et al. 2021). Playback uses acoustic cues to trigger vocalizations from the
97 target animal, enabling detection of their presence. The playback method has been often used
98 in avian studies (Conway and Gibbs 2005; Kawamura et al. 2016); however, recent studies
99 have applied it to survey mammals including ungulates (Enari et al. 2017) and primates
100 (Dacier et al. 2011; Gestich et al. 2017). Several studies have used playback with pikas
101 (Somers 1973; Conner 1983, 1985a; b; Trefry and Hik 2009), but for purposes related to the
102 communicative nature of vocalizations rather than for conducting population surveys. Studies
103 applying the playback method to survey pika populations are scarce, with only two small-
104 scale studies using it to assess the occupancy of the Northern Pika in Hokkaido (Ieiri and
105 Yanagawa 2009; Sato et al. 2009), although they did not detail their description of method
106 and efficacy, and did not objectively contrast results with the auditory observation approach.

107 To fill this research gap, we: (1) compare the efficacy of auditory observation and
108 playback to survey the occupancy of the Northern Pika; and (2) further investigate how
109 acoustic cues in playback promote species detection.

110

111

MATERIALS AND METHODS

112

113

114

115

116

117

118

119

Study area.— We conducted our study in Daisetsuzan National Park and the surrounding forests in the Tokachi region of Hokkaido, the northernmost island of Japan. We selected 18 rocky patches on talus slopes as study sites in June 2021 because rock interstices found within this landform are the primary microhabitats for Northern Pikas (Fig. 1). The study sites (elevation range 350–2200 m) were selected to cover to the greatest possible extent the known elevational distribution, ranging from 50 m (Kawabe 1990) to 2200 m (Onoyama and Miyazaki 1991). Vegetation changes across this wide elevational gradient, ranging from primarily forest vegetation at lower elevations to alpine vegetation at higher elevations. Based

120 on differences in the structure and amount of vegetation types, we classified the study sites
121 into higher ($n = 9$) and lower ($n = 9$) elevation sites (Fig. 1; Sakiyama et al. 2021).

122 All procedures in this study were conducted with the permission of the Hokkaido
123 Regional Forest Office and Daisetsuzan National Park (Permit numbers 2106071 and
124 2106072) and followed the American Society of Mammalogists guidelines (Sikes et al. 2016).

125 *Occupancy survey.*—Surveys were conducted at each site every month between July
126 and September 2021 because Northern Pikas in Hokkaido emit vocalizations frequently
127 during these months and the sites are not covered with snow. Surveys were conducted on both
128 clear and cloudy days, in the morning between 05:00 and 10:30 when pika activity is
129 reportedly highest (Kawamichi 1971).

130 To detect the presence of Northern Pikas, we conducted two point-count aural detection
131 surveys using both auditory observation and playback approaches. Auditory observation
132 consisted of listening to the natural vocalizations of the Northern Pika without intervention.
133 By contrast, playback involved the broadcast of a prerecorded pika vocalization as a cue to
134 elicit responding vocalizations. Upon arrival at each site, we remained silent and inactive for
135 10 min to reduce the possibility of human arrival stimulating pikas to vocalize. After this
136 initial waiting period, we conducted auditory observation first, followed by playback.
137 Nonetheless, pika vocalizations detected prior to the survey upon arrival at the site were also
138 recorded as a baseline to assess the efficacy of auditory observation and playback. For both
139 methods, we listened to vocalizations in a fixed facing direction (upslope or downslope) and
140 recorded the following information for each detected vocalization activity: (i) time from start
141 of the survey (time after the broadcast ended in the playback); (ii) vocalization type (see next
142 section); and (iii) direction at 10-degree intervals (all surveys were conducted by T.
143 Sakiyama). The survey timeframes were 50 min for auditory observation and 5 min for
144 playback. Auditory observation was longer because lack of intervention could delay the first

145 detection, whereas we expected a much quicker detection for playback. After the survey, site
146 occupancy was marked as present if any type of Northern Pika vocalization was detected and
147 absent if not.

148 For playback, we broadcast an audio file from a monophonic audio speaker (JBL
149 CHARGE4, JBL, California, USA; frequency response 60–20,000 Hz) as a cue to provoke
150 vocalizations from individuals. The audio file consisted of three repetitions of a Northern Pika
151 ‘song’ (also known as ‘long call’; Kawamichi 1981), each played with the speaker facing in
152 three directions perpendicularly (leftward, forward, and rightward relative to the facing
153 direction) to cover the entire study site. While the vocalization type used for playback was not
154 mentioned in previous studies (Ieiri and Yanagawa 2009; Sato et al. 2009), we used the song
155 as a cue because we successfully heard responding vocalizations in a preliminary trial when
156 using this type of call (T. Sakiyama, personal observation). We conducted surveys at a
157 monthly interval to prevent pika individuals from habituating to recorded vocalizations
158 (Trefry and Hik 2009). The song was prerecorded using a stereophonic audio recorder
159 (Olympus LS-P4 Linear PCM Recorder) located 5 m from a Northern Pika individual. After
160 successfully recording the song, we reduced external noise in the audio, modified the song
161 length from six elements (as defined in Kojima et al. 2006) to ten elements to enable longer
162 broadcast, and created an audio file comprising of three repeats of the song with an
163 approximate total duration of 30 s long. Both modification and creation procedures were
164 conducted using Audacity v. 3.0.0 (Audacity Team 2021). The spectrogram of a single song
165 and the audio file of the playback are available in Supplementary Data SD1 and SD2,
166 respectively.

167 In aural detection, including both auditory observation and playback, the auditory
168 capability of a surveyor may bias the quality of occupancy detection. Therefore, each
169 surveyor underwent preliminary training to locate pika vocalizations by ear, and we evaluated

170 the auditory capability of the surveyor in locating the sound source (distance and direction) in
171 a test experiment in order to confirm that a surveyor was efficient in detecting the Northern
172 Pika vocalizations at different directions and distances (see Supplementary Data SD3).
173 Moreover, Northern Pikas could have been more detectable from the front than from the back
174 during the actual surveys because the surveyor faced one fixed direction in the talus patch
175 during the surveys. To assess if such directional bias was introduced, we created histograms
176 of the direction from which vocalization activity was detected for both auditory observation
177 playback surveys. We counted the number of surveys that detected vocalization activities for
178 each direction over the three sessions across sites and used it as the frequency.

179 *Vocalization type classification.*—The Northern Pika is known to emit various types of
180 vocalizations (Kawamichi 1981; Lissovsky 2005), and therefore, we distinguished the
181 vocalization types by ear in the surveys. However, there has been high ambiguity in the
182 terminology of each vocalization type in previous literature. For example, Kojima et al.
183 (2006) followed terminologies in Kawamichi (1981), but Lissovsky et al. (2021) followed a
184 different terminological system (Lissovsky 2005; described in Russian). To minimize
185 potential ambiguities, we consider literature descriptions of qualitative and quantitative
186 characteristics of vocalization types as follows: the ‘long call’ described in Kawamichi (1981)
187 matches the ‘song’ described in Lissovsky (2005); the ‘short call’ in the former matches the
188 ‘alarm call’ in the latter; the ‘trill’ in the former equals the ‘chirring’ in the latter. Although
189 we could not obtain detailed information on function and spectrogram for the ‘submission
190 call’ in Kawamichi (1981) and the ‘trill’ in Lissovsky (2005), we did not detect these types in
191 our surveys. Therefore, we only considered the three aforementioned vocalization types in our
192 study, all being detectable by human ear (further detail in Supplementary Data SD4).

193 The detectable range of vocalization activities varied depending on the type of
194 vocalization—‘long call/song’ and ‘short call/alarm call’ were detectable from a distance of

195 100 m, whereas ‘trill/chirring’ was only detectable from a distance of 25 m or closer. Our
196 detection range is more conservative than that of Kawamichi (1969; i.e., 200 m) because those
197 ranges probably were attained under optimal ambient noise conditions. Given that our surveys
198 were occasionally conducted in windy conditions (≤ 6.0 m/s) and cicada singing background
199 noise (< 60 dB), we elected to use a conservative detection range of 100 m from the site
200 center in our surveys.

201 *Statistical analysis.*—To compare the efficacy of aural detection methods, we analyzed
202 the accuracy of each method with respect to time spent on the survey by creating confusion
203 matrices and computing the true positive rate and accuracy. True positive rate (also referred to
204 as sensitivity) indicates the probability of a method detecting presence at sites that are truly
205 occupied by the pika. Accuracy indicates the probability of a method detecting the true status
206 of occupancy (i.e., present or absent) at each site. The true status of occupancy at a site was
207 based on the overall combined results from all surveys over the season. Here, we assumed that
208 our overall surveys could detect the true status of occupancy at each site and that it did not
209 change during survey season (i.e., closed population setting). However, this assumption might
210 not hold true because, according to one study, the Northern Pika disperses from late July
211 through August (Kawamichi 1971), which overlaps with our survey season. Therefore, we
212 also examined how the number of sites detected changed over time. Since the timeframe
213 differed between auditory observation and playback (50 and 5 min, respectively), we divided
214 each auditory observation survey record into 10 survey periods of 5 min each and computed
215 true positive rate and accuracy of detection for each.

216 We compared the time until first detection between auditory observation and playback
217 methods to understand how intervention of the playback broadcast promoted their response.
218 We used the detection result from the first survey period for auditory observation to match the
219 timeframe of the playback survey, and used only those cases where both methods succeeded

220 in detecting the animal. We compared time until first detection using a generalized linear
221 mixed model with a negative binomial distribution. We selected this distribution because our
222 data included zero values (i.e., 0 s when detected) and because overdispersion was detected
223 when data were first analyzed using the Poisson distribution. We fitted time of first detection
224 as the response variable and methods (i.e., auditory observation and playback) as categorical
225 explanatory variables. The intercept in the linear predictor was omitted from the model and
226 site was included as a random intercept. We interpreted the difference between methods as
227 significant if the 95% confidence interval of the prediction did not overlap.

228 All statistical analyses were conducted using R 4.1.2 (R Core Team 2021). Confusion
229 matrix analysis was conducted using the *caret* package (Kuhn 2008) and mixed model
230 analyses were conducted using the *lme4* package (Bates et al. 2015).

231

232 RESULTS

233 We detected Northern Pika at 11 of the 18 sites from the occupancy surveys, with eight sites
234 corresponding to higher-elevation and three to lower-elevation sites (Fig. 1). The highest
235 elevation among the present sites was 2220 m at Mt. Hakuun, which corresponds with
236 previous reports (Onoyama and Miyazaki 1991). On the other hand, the lowest was 470 m
237 along the Horoka-Piribetsu river, which to our knowledge has previously not been reported as
238 Northern Pika habitat. The occurrence of Northern Pikas indicated a positive spatial
239 autocorrelation pattern, where present and absent sites were found relatively close to other
240 present and absent sites. We often detected presence upon arrival at the site in higher
241 elevations in early summer. In contrast, such cases were uncommon at lower elevations, with
242 lower-elevation sites necessitating the use of auditory observation or playback surveys to
243 detect presence most of the season (Supplementary Data SD5). Among the different
244 vocalization types, ‘long call/song’ and ‘short call/alarm call’ were more frequently heard

245 than ‘trill/chirring’ for all months at both higher and lower elevations (Supplementary Data
246 SD6).

247 Confusion matrix indices revealed that detection accuracy changed according to survey
248 duration for the auditory observation method. When duration increased from the shortest (5
249 min) to the longest (50 min) setting, the true positive rate increased from 0.606 to 0.909 and
250 accuracy increased from 0.759 to 0.944 (Fig. 2). Both indices initially changed drastically
251 over shorter survey durations, reaching a plateau around the sixth survey period,
252 approximately 30 min after the survey was started. For playback, which was conducted for
253 only 5 min, the true positive rate and accuracy were 0.879 and 0.926, respectively. We found
254 that values from playback are comparable to those of the longest duration setting of the
255 auditory observation method (Fig. 2).

256 The time at first detection of Northern Pika vocalizations was statistically significantly
257 shorter for playback than auditory observation (Table 1, Fig. 3). First vocalizations—recorded
258 using playback—were detected 23.8 s after the audio was played on average, whereas they
259 were detected after 127.6 s with auditory observation.

260 During auditory observation surveys, Northern Pika vocalizations were detected more
261 frequently from the front side than the back side for ‘short calls/alarm calls’. However, in
262 other cases, detections were oriented sideward (i.e., left and right sides), which differed from
263 our initial expectations. Detection of ‘trill/chirring’ was rare during our surveys (Fig. 4,
264 Supplementary Data SD6).

265 DISCUSSION

266 In this study, we compared the performance of two aural methods for detecting presence of
267 the Northern Pika by listening for their vocalizations. Our results showed that both auditory
268 observation and playback are highly and comparably effective in detecting presence. And
269 while presence was more frequently represented at higher-elevation than lower-elevation sites

270 (Fig. 1), this agrees with previous reports describing their distribution in Hokkaido (Onoyama
271 and Miyazaki 1991; Kawabe 2008), and modelled distribution of the species that reflects their
272 adaptation to cold environments (Sakiyama et al. 2021). These results indicate that
273 vocalization-based surveys have high potential for monitoring the Northern Pika in its natural
274 habitat and contributing to understanding the effects of environmental change on their
275 populations.

276 *Occupancy survey.*—The effectiveness of auditory observation increased significantly
277 as survey duration increased. This result is intuitive because a longer duration increases the
278 chance of recording vocalization activities. Pikas often vocalize for alarm and territorial
279 purposes (Somers 1973; Kawamichi 1981; Lissovsky 2005; Trefry and Hik 2009), as well as
280 for positioning and localizing other individuals (Conner 1982). Since these natural events are
281 likely to occur randomly over time, shorter survey durations are likely to include more false-
282 negative detections. Our initial expectation was that detection of Northern Pikas would
283 require survey durations longer than 5 min, a timeframe used in bird surveys (Ralph et al.
284 1995), because of their low conspicuity. Nonetheless, our results revealed that detection
285 efficacy reached a plateau when auditory observation was conducted for more than 30 min.
286 This suggests that continuing the survey after 30 min does not provide significant gains in
287 information relative to the effort invested—this saturation effect was also reported previously
288 in a study that used a similar point-count method (Hutschenreiter et al. 2021).

289 Notably, the five-minute playback survey reached a similar level of efficacy as the
290 longest auditory observation survey in detecting the presence of Northern Pikas. Moreover,
291 the playback method detected, on average, their presence approximately four times earlier
292 than the auditory observation method. Therefore, cues broadcast by the audio speaker are
293 highly effective in eliciting vocal responses. Although this phenomenon has been observed
294 and effectively used in previous studies to understand the behavioral nature of vocalization

295 (Somers 1973; Conner 1982; Trefry and Hik 2009), our results suggest that it can also be used
296 as a highly effective, fast, and affordable method for surveying site occupancy in this species.

297 Another significant advantage of the aural detection survey was that it enabled detection
298 of presence of Northern Pikas at lower elevations, including areas previously unreported as
299 pika habitat (i.e., along the Horoka-Piribetsu river). These landscapes are characterized by
300 denser vegetation structure and fewer rock interstices than higher elevations (Sakiyama et al.
301 2021), hampering traditional visual approaches for detection. Moreover, vocalization
302 activities were not easily detected upon arrival at their habitats at lower elevations, in contrast
303 to higher elevations (Supplementary Data SD5), making aural detection method the only
304 effective approach. Lower-elevation populations inhabit marginal regions of their distribution,
305 whereas core populations are located at higher elevations (Sakiyama et al. 2021). Therefore,
306 aural detection provides a promising tool for efficient monitoring of Northern Pikas that might
307 be more vulnerable to climate change and human land uses at lower elevations.

308 We found that the number of sites at which presences was detected increased gradually
309 during the summer (Supplementary Data SD6), suggesting that occupancy could be detected
310 by vocalization during and after the reproductive season, as expected from previous studies
311 (Kawamichi 1971). This trend was more apparent at lower elevations, and it could be a result
312 of false-negative observations, even though they did inhabit the site from early in the season.
313 This finding emphasizes the importance of repeating the survey throughout the summer to
314 improve chances of detecting presence. However, Northern Pikas have been reported to
315 disperse from late July to August at lower elevations (Kawamichi 1971), whereas American
316 Pikas disperse from mid-May to mid-August at high altitude (2000 m) habitats (Smith 1974).
317 Therefore, the increase in number of sites at which they are present over the summer could
318 instead be a result of dispersal events from adjacent sites. If this was the case, then the
319 assumption of a closed population for our survey would not have been met. Although we

320 primarily considered the timing of snowmelt to determine our survey season, a more precise
321 definition may be needed to better accommodate the population setting in future studies.
322 Fixing survey season prior to or after dispersal, for example, could enhance accuracy. To this
323 end, further studies into the dispersal ecology of the Northern Pika are urgently needed.

324 *Advantages and disadvantages of aural detection methods.*—Our results suggest that
325 both auditory observation and playback are effective in surveying the presence of the
326 Northern Pika, but each has advantages and disadvantages.

327 A common advantage of using point-count aural detection is that the surveyor need not
328 venture into densely vegetated and potentially dangerous habitats. Talus slopes consist of
329 piles of large, loose rocks with interspersed vegetation of various types and structures
330 including lichens, moss, herbaceous plants, shrubs, and trees (Sakiyama et al. 2021). Hence,
331 walking steadily within the site requires care, and the simultaneous search for signs of pikas is
332 laborious and time-consuming. Alternatively, when the surveyor does not move, vocalization
333 is less influenced by these factors, although the observational environment (e.g., strong wind)
334 needs to be considered in terms of its effect on auditory detection. Moreover, use of the point-
335 count survey is advantageous because it avoids trampling on and damaging vegetation, which
336 is the main dietary resource of Northern Pikas. Furthermore, while feces can represent
337 incomplete signs of recent activity (Nichols 2011), vocalization-based surveys only detect
338 active signs of occupancy.

339 Nevertheless, the use of vocalization-based methods has several disadvantages. First, it
340 is difficult to define the area where pika vocalizations can be detected by the human ear. We
341 conservatively assumed that the aurally-detectable range of the Northern Pika is 100 m,
342 although complex topography could alter this range. Our results show that detection of the
343 vocalization activities were more prevalent in frontward and sideward directions (Fig. 4),
344 suggesting that choice of surveying point and facing direction will influence results.

345 Moreover, if the study area is larger than the range of detection by ear (approx. 100–200 m;
346 Kawamichi 1969), it will be important to conduct the survey at multiple points. Second, aural
347 detection methods require training of the surveyor to differentiate Northern Pika vocalizations
348 from other animals. While no other small mammals cooccurring in rocky patches make
349 vocalizations identical to the Northern Pika, the Eurasian Woodcock (*Scolopax rusticola*)
350 makes a vocalization similar to the ‘short call/alarm call’ during their nocturnal flights (T.
351 Sakiyama personal observation). Therefore, we ensured no potential confusion by planning
352 our surveys during the morning, and likewise avoiding the active time of other animals are
353 likely to reduce false detections. Third, although we were cautious to minimize these factors,
354 we did not assess if and how the presence of the surveyor could affect results. For example,
355 although we stayed silent before and during the survey, animals could still have detected our
356 presence through olfaction. If this was the case and their vocalizing behavior was reduced as a
357 result, our estimation of presence frequency would likely have been underestimated. Fourth, it
358 remains unclear how behavioral differences at different life stages will affect aural detection
359 results. For example, younger individuals might vocalize at lower volume and less frequently
360 than older individuals, which could also lead to a reduction in detection probability.

361 The methodological differences between auditory observation and playback included
362 duration of the survey (survey durations were up to 50 min for auditory observation method
363 and 5 min for playback) and for the latter, broadcast of a prerecorded call to stimulate pika
364 vocalizations. However, detection results between playback and auditory observation were
365 comparable. Thus, playback has an advantage over auditory observation because it reduces
366 survey duration and, therefore, effort and cost. Resource allocation is key in planning field
367 surveys. With limited resources, researchers frequently face the challenging task of balancing
368 the number of sites and survey effort within each site. A reduction in survey time enables an
369 increase in the number of sites, which increases robustness of research results.

370 Playback was conducted by broadcasting prerecorded vocalizations to elicit responding
371 calls from inhabiting individuals. The implementation of playback requires important
372 considerations that are not required for auditory observation. First, pikas could become
373 habituated to playback audio if it were played frequently (Trefry and Hik 2009). Habituation
374 of target species should be avoided because it may increase false-negative observations and
375 preclude consistent surveys. This problem can be avoided by designing surveys such that they
376 are sufficiently spaced over time (i.e., once a month in this study)—broadcasting audios
377 recorded from different individuals could also reduce habituation. Second, the effect of the
378 prerecording source on response remains unclear. If the response behavior differs depending
379 on the spatial source of the broadcast (i.e., place of prerecording), as was reported for the
380 American Pika (Conner 1983), the efficacy of playback will likely be spatially biased.
381 However, this effect was likely minimal in our study because only a single acoustic dialect is
382 found in Northern Pika populations in Hokkaido (Lissovsky 2005; Lissovsky et al. 2021).
383 Conversely, the choice of prerecording source will require in-depth consideration in situations
384 where playback is conducted at a broader spatial extent, increasing the likelihood of including
385 multiple dialects—in these kinds of surveys, the simplicity and passivity of auditory
386 observation could be advantageous.
387

388

ACKNOWLEDGEMENTS

389 This study was financially supported by the JSPS KAKENHI Grant Number JP22J10191 and
390 grants-in-aid of The Inui Memorial Trust for Research on Animal Science, Japan and The
391 Zoshinkai Fund for Protection of Endangered Animals, Japan. We would like to thank the
392 Tokachi Shikaoi Geopark, Ministry of Environment, and Hokkaido Regional Forest Office for
393 helping us conduct the research. We are also grateful to Yu Takahata and Erika Yoshida for
394 assisting us in the field surveys.

395

396

CONFLICT OF INTEREST

397 The authors declare they have no conflict of interest.

398

399

AUTHOR CONTRIBUTIONS

400 TS contributed to conceptualization and design of methodology; TS contributed to fieldwork
401 for data collection, data analysis, funding acquisition, and writing the original draft; TS and
402 JGM contributed to discussion of results, writing the final manuscript and gave final approval
403 for publication.

404

405

DATA AVAILABILITY

406 The data that support the findings of this study are available from the corresponding author
407 (TS) upon reasonable request.

408

409

SUPPLEMENTARY DATA

410 **Supplementary Data SD1.**— Spectrographs of the audio file used in the playback
411 survey. The actual audio could be heard from the audio embedded in the file.

412

Supplementary Data SD2.— Audio file used in the playback survey.

413 **Supplementary Data SD3.**— A scatter plot showing the auditory capability of the
414 surveyor in estimating the location of the sound source in a talus patch.

415 **Supplementary Data SD4.**— Organization process of the terminologies used for the
416 Northern Pika vocalizations in previous literature.

417 **Supplementary Data SD5.**— Raw detection result of occupancy from the surveys.

418 **Supplementary Data SD6.**— Bar plots indicating the number of sites detected during
419 the surveys for each method and vocalization type.

420

421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445

LITERATURE CITED

Almond R.E.A., Grooten M., Peterson T. 2020. Living Planet Report 2020 – Bending the curve of biodiversity loss. World Wildlife Fund, Gland, Switzerland.

Audacity Team. 2021. Audacity(R): Free audio editor and recorder. Version 3.0.0 <https://audacityteam.org/>.

Bates D., Mächler M., Bolker B., Walker S. 2015. Fitting linear mixed-effects models using lme4. *Journal of Statistical Software* 67:1–48.

Beever E.A., Ray C., Mote P.W., Wilkening J.L. 2010. Testing alternative models of climate-mediated extirpations. *Ecological Applications* 20:164–178.

Beever E.A., Ray C., Wilkening J.L., Brussard P.F., Mote P.W. 2011. Contemporary climate change alters the pace and drivers of extinction. *Global Change Biology* 17:2054–2070.

Billman P.D., Beever E.A., McWethy D.B., Thurman L.L., Wilson K.C. 2021. Factors influencing distributional shifts and abundance at the range core of a climate-sensitive mammal. *Global Change Biology* 27:4498–4515.

Collen B., Loh J., Whitmee S., McRae L., Amin R., Baillie J.E.M. 2009. Monitoring change in vertebrate abundance: the Living Planet index. *Conservation Biology* 23:317–327.

Conner D.A. 1982. Geographic variation in short calls of pikas (*Ochotona princeps*). *Journal of Mammalogy* 63:48–52.

Conner D.A. 1983. Response of pikas to playback of geographic variations of the short call. *Journal of Mammalogy* 64:336–337.

Conner D.A. 1985a. The function of the pika short call in individual recognition. *Zeitschrift für Tierpsychologie* 67:131–143.

Conner D.A. 1985b. Analysis of the vocal repertoire of adult pikas: ecological and evolutionary perspectives. *Animal Behavior* 33:124–134.

- 446 Conway C.J., Gibbs J.P. 2005. Effectiveness of call-broadcast surveys for monitoring marsh
447 birds. *The Auk* 122:26–35.
- 448 Dacier A., de Luna A.G., Fernandez-Duque E., Di Fiore A. 2011. Estimating population
449 density of Amazonian Titi Monkeys (*Callicebus discolor*) via playback point counts.
450 *Biotropica* 43:135–140.
- 451 Di Fonzo M.M.I., Collen B., Chauvenet A.L.M., Mace G.M. 2016. Patterns of mammalian
452 population decline inform conservation action. *Journal of Applied Ecology* 53:1046–
453 1054.
- 454 Enari H., Enari H., Okuda K., Yoshita M., Kuno T., Okuda K. 2017. Feasibility assessment of
455 active and passive acoustic monitoring of sika deer populations. *Ecological Indicators*
456 79:155–162.
- 457 Erb L.P., Ray C., Guralnick R. 2011. On the generality of a climate-mediated shift in the
458 distribution of the American pika (*Ochotona princeps*). *Ecology* 92:1730–1735.
- 459 Gestich C.C., Caselli C.B., Nagy-Reis M.B., Setz E.Z.F., da Cunha R.G.T. 2017. Estimating
460 primate population densities: the systematic use of playbacks along transects in
461 population surveys. *American Journal of Primatology* 79:e22586.
- 462 Hall L.E., Chalfoun A.D., Beever E.A., Loosen A.E. 2016. Microrefuges and the occurrence
463 of thermal specialists: implications for wildlife persistence amidst changing
464 temperatures. *Climate Change Responses* 3:1–12.
- 465 Henry P., Russello M.A. 2011. Obtaining high-quality DNA from elusive small mammals
466 using low-tech hair snares. *European Journal of Wildlife Research* 57:429–435.
- 467 Hutschenreiter A., Ramos-Fernández G., Aureli F. 2021. Line-transect versus point-transect
468 sampling: the effects of survey area and survey effort on method efficiency for
469 Geoffroy's spider monkeys. *Wildlife Research* 48:590–597.

470 Ieiri A., Yanagawa H. 2009. Habitat of northern pikas (*Ochotona hyperborea yesoensis*) in
471 the low altitude area at the southern tip of the Hidaka Mountains. Research Bulletin of
472 Obihiro University 30:53–60.

473 Kawabe M. 1990. Northern pika *Ochotona hyperborea yesoensis* habitat at an altitude of 50m,
474 Hokkaido. Bulletin of Higashi Taisetsu Museum of Natural. History 12:89–92.

475 Kawabe M. 2008. The distribution of *Ochotona hyperborea yesoensis* in Hokkaido. Bulletin
476 of Higashi Taisetsu Museum of Natural. History 30:1–20.

477 Kawamichi T. 1969. Behaviour and daily activities of the Japanese Pika, *Ochotona*
478 *hyperborea yesoensis*. Journal of the Faculty of Science Hokkaido University Series
479 VI Zoology 17:127–151.

480 Kawamichi T. 1970. Social Pattern of the Japanese Pika, *Ochotona hyperborea yesoensis*.
481 Journal of the Faculty of Science Hokkaido University Series VI Zoology 17:462–473.

482 Kawamichi T. 1971. Annual cycle of behaviour and social pattern of the Japanese pika,
483 *Ochotona hyperborea yesoensis*. Journal of the Faculty of Science Hokkaido
484 University Series VI Zoology 18:173–185.

485 Kawamichi T. 1981. Vocalizations of *Ochotona* as a taxonomic character. In: Myers K.,
486 MacInnes C. D., editors. Proceedings of the World Lagomorph Conference. Guelph
487 University Press, Ontario, Canada; p. 324–339

488 Kawamura K., Yamaura Y., Senzaki M., Yabuhara Y., Akasaka T., Nakamura F. 2016.
489 Effects of land use and climate on the distribution of the Jungle Nightjar *Caprimulgus*
490 *indicus* in Hokkaido, Northern Japan. Ornithological Science 15:203–212.

491 Kojima N., Onoyama K., Kawamichi T. 2006. Year-long stability and individual differences
492 of male long calls in Japanese pikas, *Ochotona hyperborea yesoensis* (Lagomorpha).
493 Mammalia 70:80–85.

494 Korolev A.N. 2017. The technique of route census Northern Pika (*Ochotona (Pika)*
495 *Hyperborea* Pallas, 1811, Lagomorpha, Ochotonidae). ВЕСТНИК ИБ КОМИ НЦ
496 УрО РАН:44–48.

497 Kuhn M. 2008. Building predictive models in R using the caret package. Journal of Statistical
498 Software 28:1–26.

499 Lissovsky A.A. 2005. Comparative analysis of the vocalization of pikas (*Ochotona*,
500 Mammalia) from Alpina-hyperborea group. Byulleten’Moskovskogo Obshchestva
501 Ispytateley Prirody 110:12–26.

502 Lissovsky A.A., Obolenskaya E.V., Dokuchaev N.E., Okhlopkov I.M. 2021. Intraspecific
503 variation and taxonomy of northern pika *Ochotona hyperborea* (Mammalia,
504 Lagomorpha). Journal of Mammalogy 102:28–53.

505 Nichols L.B. 2011. Fecal pellets of American pikas (*Ochotona princeps*) provide a crude
506 chronometer for dating patch occupancy. Western North American Naturalist 70:500–
507 507.

508 Onoyama K., Miyazaki T. 1991. Distribution in Hokkaido. In: Hokkaido Government,
509 editors. Survey Report of Wildlife Distribution: Pika Ecology. Hokkaido Government,
510 Hokkaido, Japan; p. 25–55.

511 R Core Team. 2021. R: A language and environment for statistical computing. Vienna,
512 Austria. <https://www.R-project.org/>.

513 Ralph C.J., Droege S., Sauer J.R. 1995. Managing and monitoring birds using point counts:
514 standards and applications. In: Ralph C.J., Droege S., Sauer J.R., editors. Monitoring
515 Bird Populations by Point Counts. USDA Forest Service, Albany, USA; p. 161–169.

516 Sakiyama T., Morimoto J., Watanabe O., Watanabe N., Nakamura F. 2021. Occurrence of
517 favorable local habitat conditions in an atypical landscape: Evidence of Japanese pika
518 microrefugia. Global Ecology and Conservation 27:e01509.

519 Sato S., Yanagawa H., Ishiyama K., Yatsu K. 2009. Use of small rock slides by the northern
520 pika, *Ochotona hyperborea yesoensis* in low altitudinal areas of Taisetsu Mountains.
521 Journal of the Japanese Wildlife Research Society 34:31–36.

522 Sikes R.S., and the Animal Care and Use Committee of the American Society of
523 Mammalogists. 2016. 2016 Guidelines of the American Society of Mammalogists for
524 the use of wild mammals in research and education: Journal of Mammalogy 97:663–
525 688.

526 Smith A.T. 1974. The distribution and dispersal of pikas: influences of behavior and climate.
527 Ecology 55:1368–1376.

528 Smith A. T., Johnston C. H., Alves P. C., Hackländer K. 2018. Lagomorphs: pikas, rabbits,
529 and hares of the world. JHU Press, Baltimore, Maryland.

530 Somers P. 1973. Dialects in southern Rocky Mountain pikas, *Ochotona princeps*
531 (Lagomorpha). Animal Behavior 21:124–137.

532 Stewart J.A.E., Perrine J. D., Nichols L. B., Thorne J. H., Millar C. I., Goehring K. E.,
533 Massing C. P., Wright D. H. 2015. Revisiting the past to foretell the future: summer
534 temperature and habitat area predict pika extirpations in California. Journal of
535 Biogeography 42:880–890.

536 Thompson W. 2004. Sampling rare or elusive species: concepts, designs, and techniques for
537 estimating population parameters. Island Press, Washington, D.C.

538 Trefry S.A., Hik D.S. 2009. Eavesdropping on the neighbourhood: Collared Pika (*Ochotona*
539 *collaris*) responses to playback calls of conspecifics and heterospecifics. Ethology
540 115:928–938.

541 Wang X., Liang D., Jin W., Tang M., Liu S., Zhang P. 2020. Out of Tibet: genomic
542 perspectives on the evolutionary history of extant pikas. Molecular Biology and
543 Evolution 37:1577–1592.

- 544 Yang J., Wang Z. L., Zhao X. Q., Wang D. P., Qi D. L., Xu B. H., Ren Y. H., and Tian H. F.
545 2008. Natural selection and adaptive evolution of leptin in the Ochotona Family driven
546 by the cold environmental stress. PLoS ONE 3:e1472.
- 547 Yoccoz N.G., Nichols J.D., Boulinier T. 2001. Monitoring of biological diversity in space and
548 time. Trends in Ecology & Evolution 16:446–453.
- 549
550
551

552 **Fig. 1.**—Map of the study area in Hokkaido, Japan (top-right panel) and the study sites within
553 to survey the occupancy of the Northern Pika (left panel), differentiated by occupancy status
554 and elevation class (black indicates presence, white indicates absence, circle indicates higher
555 elevations, and triangle indicates lower elevations). Aggregated sites are shown in higher
556 resolution (mid-right panel). Regional boundary is indicated in dotted lines and the area of
557 Daisetsuzan National Park is shown as a solid line.

558 **Fig. 2.**—Relationship between confusion matrix indices and duration of the survey to detect
559 presence of Northern Pikas. Note that auditory observation (circle) was conducted for ten
560 survey periods (five minutes \times ten periods), whereas playback (triangle) was conducted only
561 for one survey period (five minutes).

562 **Fig. 3.**—Difference between time of first detection of the animal using auditory observation
563 and playback method. Error bars indicate the 95% confidence interval around the mean.

564 **Fig. 4.**—Circular histograms showing frequency of the directions from which the
565 vocalization activities were detected, plotted separately for the survey method and detected
566 vocalization type.

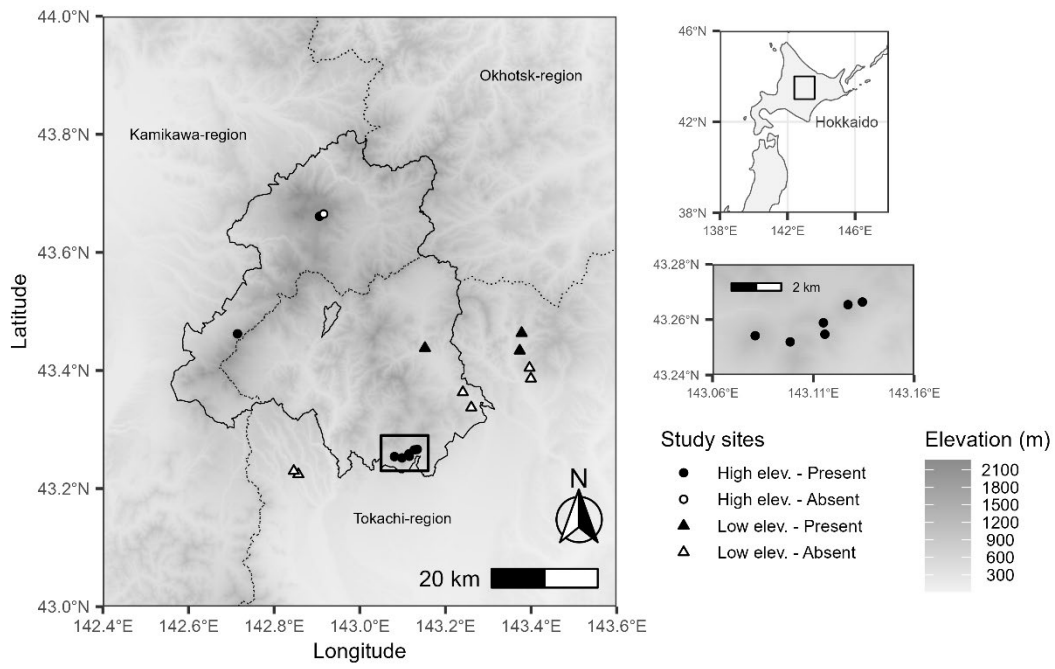
567

568

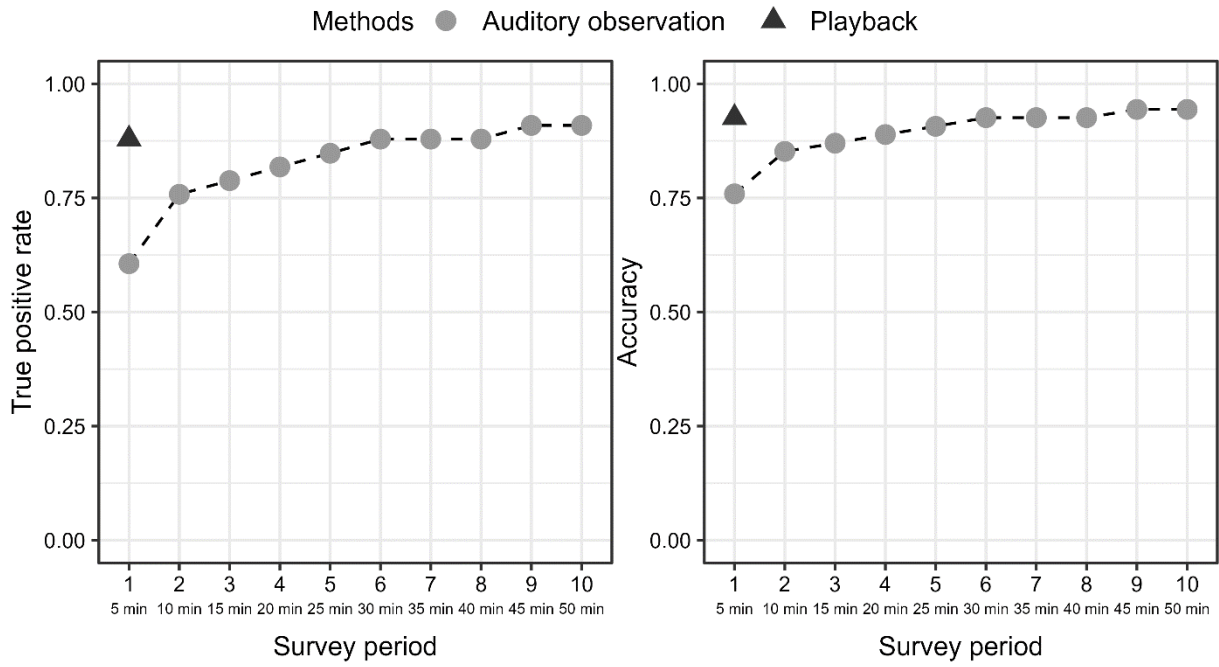
569 **Table 1.**—Parameters and performance of generalized linear mixed models to assess the
 570 difference of time until first detection of the Northern Pika between auditory observation and
 571 playback.

Parameter	β	SE	z	P
Auditory observation	4.849	0.267	18.142	< 0.001
Playback	3.171	0.343	9.246	< 0.001

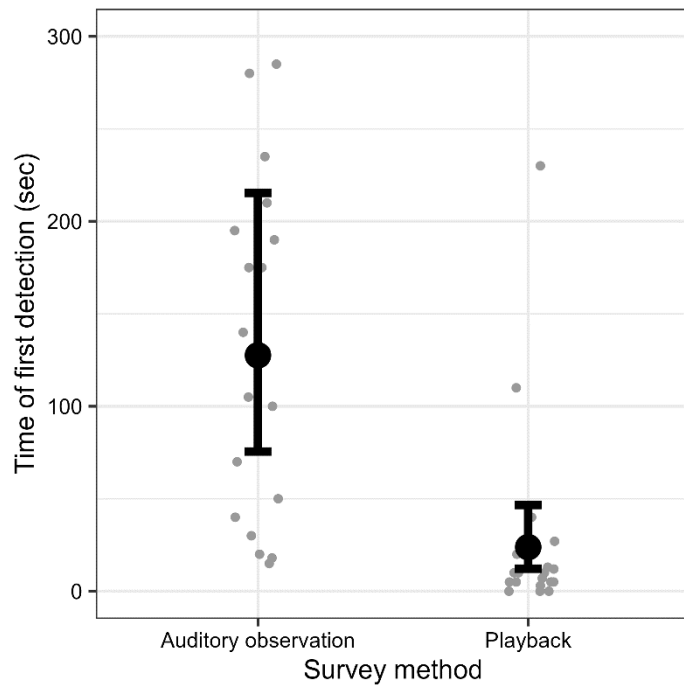
572
 573



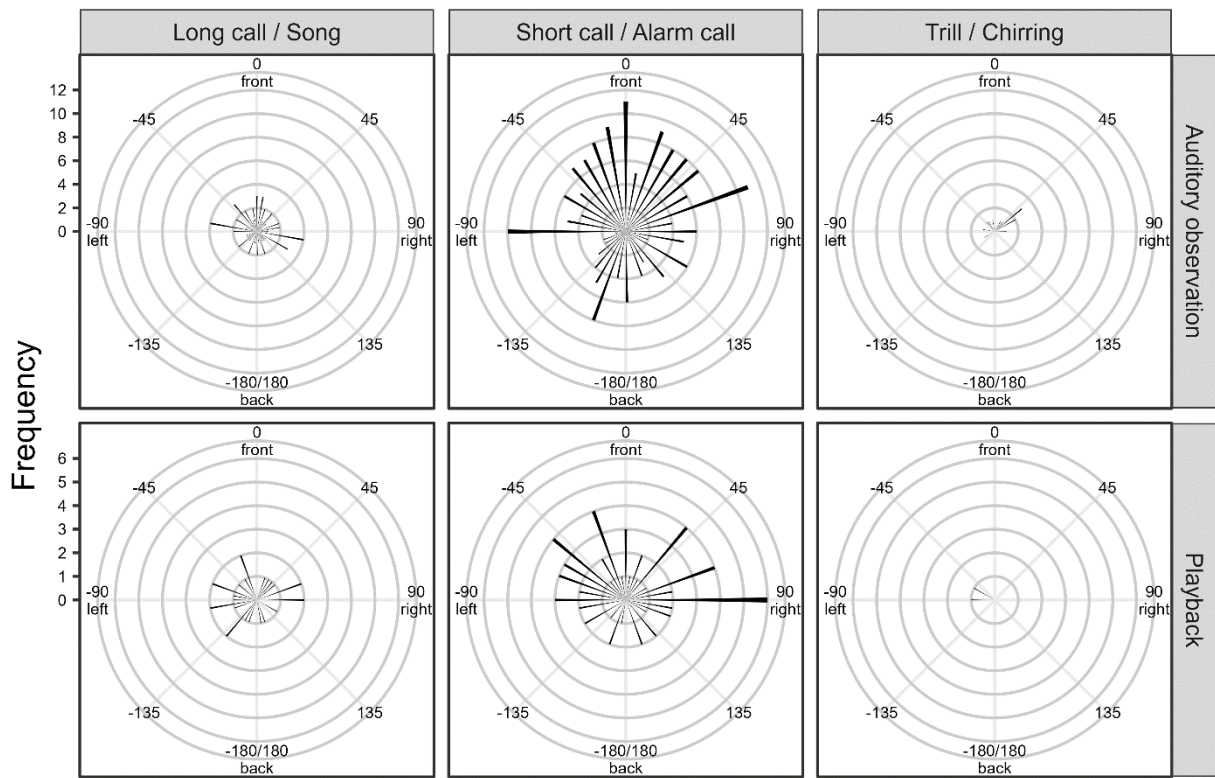
574
575 **Fig. 1**
576



577
578 **Fig. 2**
579



580
581 **Fig. 3**
582

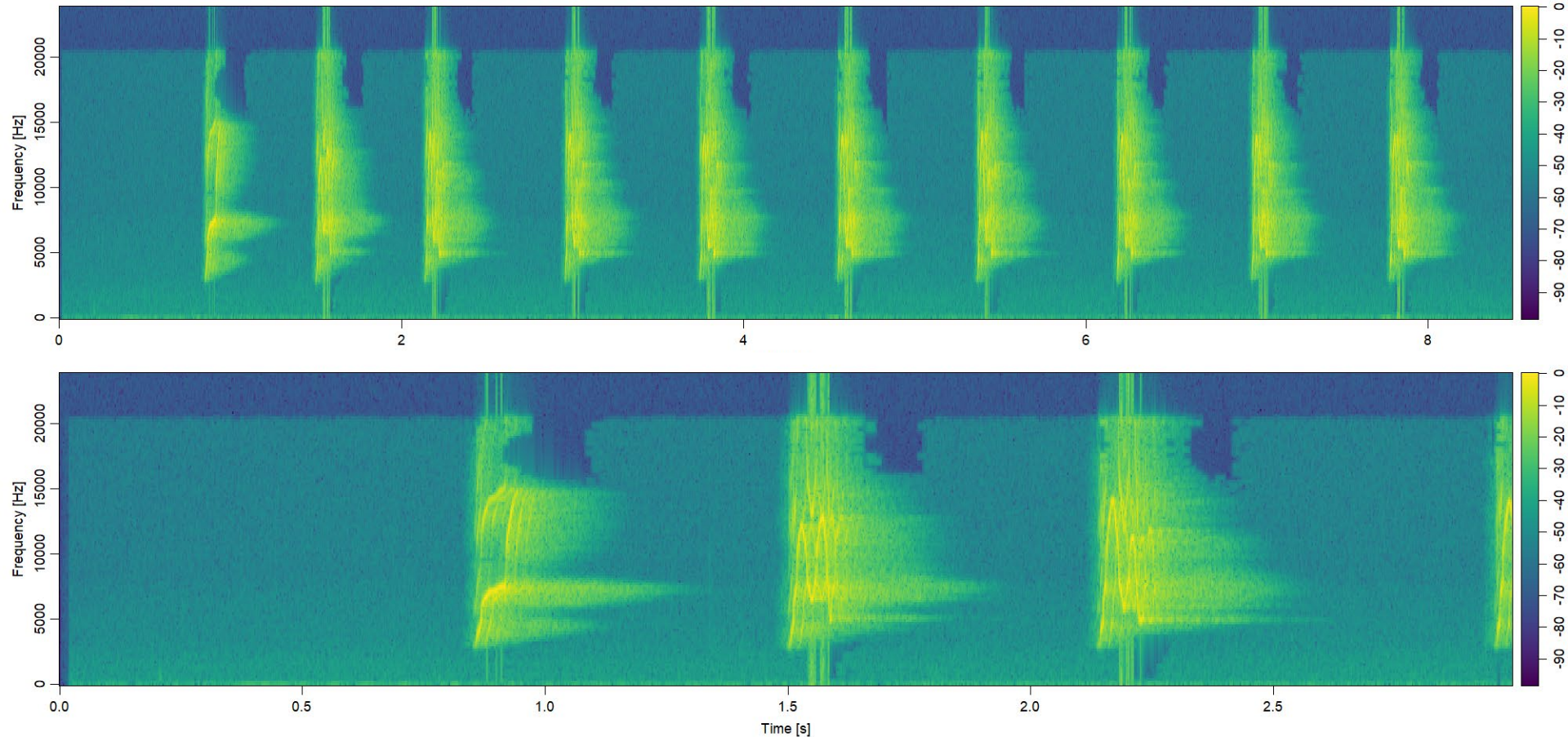


583

584 **Fig. 4**

Aural detection methods enable detection of northern pika occupancy in densely vegetated habitats

Supplementary Data SD1

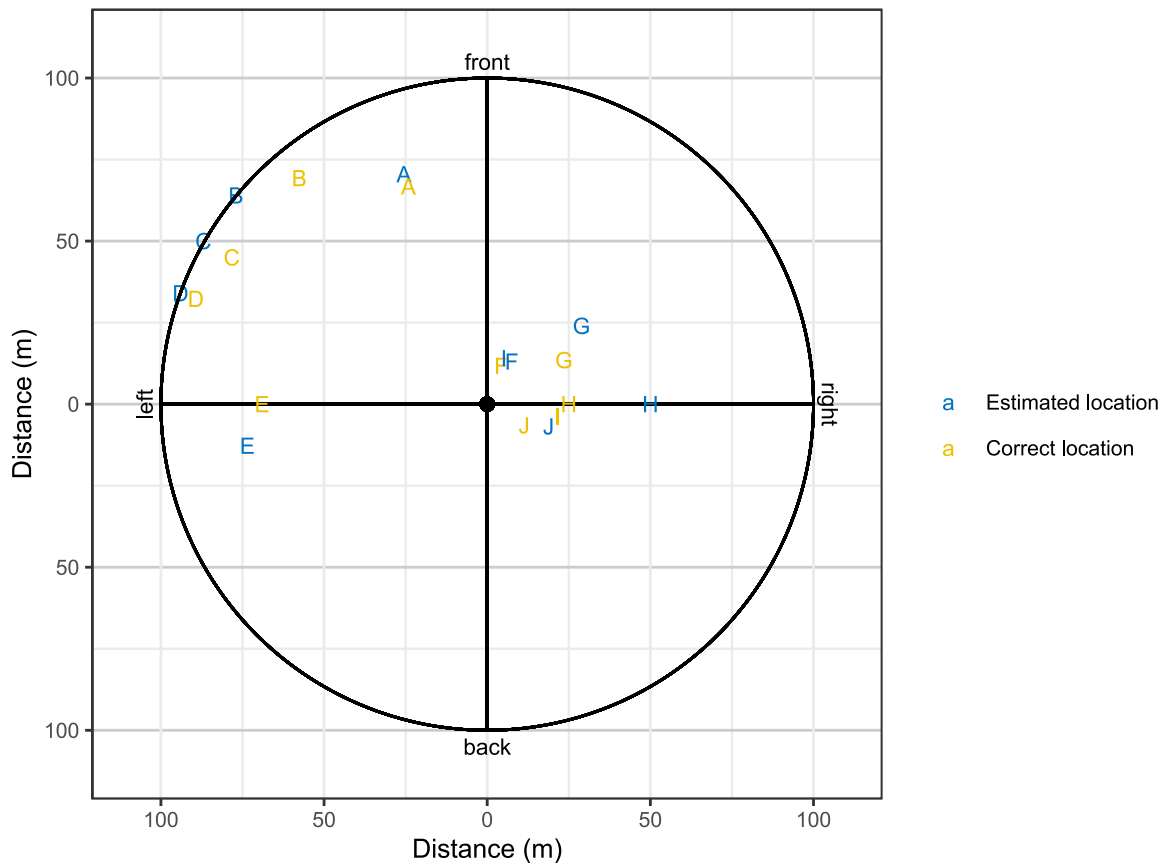


Spectrographs of the audio file used in the playback survey, showing the whole length (top) and the first three seconds in zoom (bottom) of one “long call” or the “song” of the northern pika. Three repeats of this long call was broadcasted from an audio speaker in the actual survey. Audio could be heard from the input audio on the right. Actual audio file used for the playback is archived in Supplementary Data SD2.



Aural detection methods enable detection of northern pika occupancy in densely vegetated habitats

Supplementary Data SD3



We evaluated the auditory capability of the surveyor to detect the presence of the northern pika. Given that controlling the location of the animal vocalization is impossible, a survey assistant instead broadcasted the northern pika vocalization from an audio speaker hidden at a random location in the talus patch and the surveyor estimated the distance and direction to the sound source from the site center. These records were then compared with the correct location of the sound source. The plot shows the estimated and correct location of the sound source, indicating that the surveyor was efficient in detecting the northern pika vocalization heard from various directions and distances.

Aural detection methods enable detection of northern pika occupancy in densely vegetated habitats

Supplementary Data SD4

Various types of vocalizations are known to be used for specific functions in northern pikas, as described in detail in Kawamichi (1981) and Lissovsky (2005) (described in Russian). However, different terminologies were used between these literatures to describe the vocalization system of northern pikas. Moreover, since such terminological ambiguity remained untouched for unification, successive literature often refer to either terminological system for citation, which hamper comparison between studies. For example, Kojima et al. (2006) followed terminologies in Kawamichi (1981), whereas Lissovsky et al. (2021) followed Lissovsky (2005).

To reach a unified understanding of the vocalization system of the northern pika between Kawamichi (1981) and Lissovsky (2005), we referred to the characteristics and functions of each vocalization type and situations in which they are used for qualitative comparison. We also considered the spectrographs (= sonograms) of the vocalizations to enable quantitative comparison. Since large geographical variation are known to exist even within a same vocalization type in northern pikas (Lissovsky 2005; Lissovsky et al. 2021), we referred to geographically close population pairs between the literature for spectrograph comparison (i.e., Hokkaido, Japan in Kawamichi (1981) and Sakhalin and Sikhote-Alin, Russia in Lissovsky (2005)).

Various terms such as *long call*, *short call*, *trill*, and *submission call* have been used in Kawamichi (1981) with reference to American pika literature. And *alarm call* (Предупреждающий об опасности сигнал), *song* (Песня), and *chirring* (Стрекотание)

have been used in Russian terms in Lissovsky (2005), and *song* of the northern pika consist of vocalization elements such as the *trill* (Трель) and *modulated call* (модулированный сигнал) (Note that these Russian-English translations follow the English abstract of Lissovsky 2005). As we compared the qualitative and quantitative characteristics between the literature, we organized the vocalization terminology as follows (Fig. SD4-1, SD4-2): *long call* in Kawamichi (1981) matched with the *song* in Lissovsky (2005) because elements of the *long call* showed similar spectrographic shape of modulation with the *modulation call* of a *song* (W-shaped waves) and both are explained as being made by males only and for territorial demonstration; *short call* in the former matched with *alarm call* in the latter because both showed similar spectrographic shape (upward stroke) and are described as being emitted in single or multiple vocalizations with irregular intervals in response to an approaching danger or without any reason; *trill* in the former matched with *chirring* in the latter because both showed similar spectrographic shape (diminishing upward strokes) and are explained as being heard when the animal escapes into the rock under fright conditions. We could not obtain detailed information of function and spectrogram for *submission call* in Kawamichi (1981) and *trill* in Lissovsky (2005). The term *trill* was used differently between the literature, but we classified them as different vocalization types since the spectrographic shape and situation of their use were not resembling, suggesting that their use requires care (Lissovsky 2005).

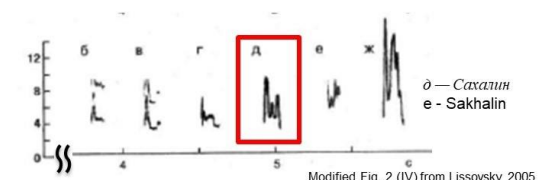
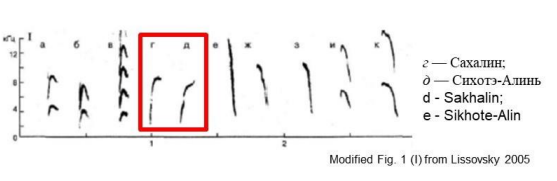
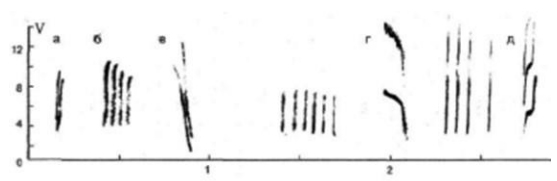
Kawamichi 1981 Study region: Hokkaido, Japan	Lissovsky 2005 Study region: Russia
<p>Long call Explanation: Emitted in series, consisting of 4 -16 cries. Given only by males in both reproductive and non - reproductive season. May function as territorial advertisement.</p>	<p>Песня English translation: song Explanation: Emitted in series, consisting of 1-23 modulated calls (модулированный сигнал). Used only by males for roll-calls and territory demonstrations.</p>
<p>Shown as A-D in Fig.1 in Kawamichi 1981</p>	
<p>Short call Explanation: Emitted singly or in series at irregular intervals. Produced by all individuals, irrespective of sex and age. Increase in the non-reproductive season and may be related with food storage.</p>	<p>Предупреждающий об опасности сигнал English translation: Alarm call Explanation: Emitted in series at irregular intervals, consisting of 1-20 cries. Produced both in response to an approaching danger: a person, a bird, or a rock fallen from a rock, and without any visible reason.</p>
<p>Shown as E-J in Fig.1 in Kawamichi 1981</p>	
<p>Trill Explanation: Produced by all individuals, irrespective of sex and age. Emitted singly when pikas quickly enter rock crevices.</p>	<p>Стрекотание English translation: Chirring, chirping Explanation: Consist of short chirp or a series of squeaks. Produced during a strong fright, when the animal unexpectedly collides with a human and rushes under rocks.</p>
<p>Shown as M and N in Fig.1 in Kawamichi 1981</p>	

Fig. SD4-1 Matching vocalization types between Kawamichi (1981) and Lissovsky (2005). Spectrograms with a red box in Lissovsky (2005) are from northern pikas in Sakhalin or Sikhote-Alin, which are the geographically closest populations from that in Kawamichi (1981).

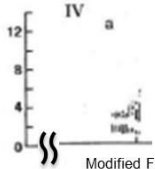
Kawamichi 1981 Study region: Hokkaido, Japan	Lisovsky 2005 Study region: Russia
No matching vocalization type.	<p data-bbox="794 309 1347 488"> Трель English translation: Trill Explanation: Produced both independently and in combination with other vocalization types. Used for territorial and mating demonstrations. Given only by males. </p>  <p data-bbox="970 678 1241 701">Modified Fig. 2 (IV) from Lisovsky 2005</p>
<p data-bbox="209 741 762 857"> Submission call Explanation: Whining sound. Repeated several times. Used when an individual was approached or chased by another. </p>	No matching vocalization type.
No spectrogram provided.	

Fig. SD4-2 Unmatching vocalization types between Kawamichi (1981) and Lisovsky (2005).

Aural detection methods enable detection of northern pika occupancy in densely vegetated habitats

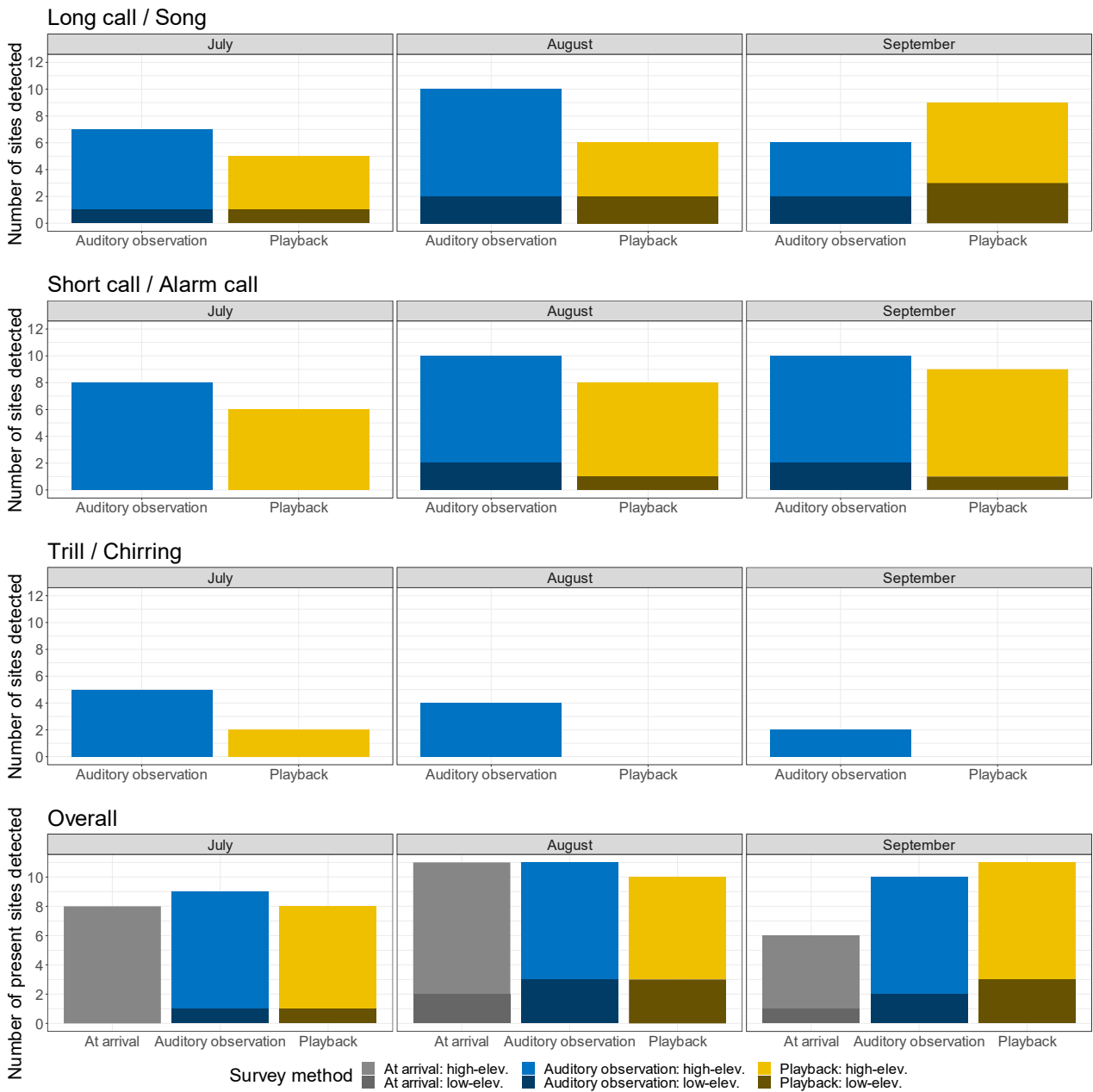
Supplementary Data SD5

Detection result of occupancy upon arrival at the site, after auditory observation, and after playback. “0” and “1” indicate absence and presence, respectively.

Site	Elevation (m)	Class	Upon arrival			Auditory observation			Playback			True occupancy
			Jul	Aug	Sep	Jul	Aug	Sep	Jul	Aug	Sep	
1	350	lower	0	0	0	0	0	0	0	0	0	0
2	350	lower	0	0	0	0	0	0	0	0	0	0
3	450	lower	0	0	0	0	0	0	0	0	0	0
4	450	lower	0	0	0	0	0	0	0	0	0	0
5	470	lower	0	0	0	0	1	0	0	1	1	1
6	510	lower	0	0	0	0	0	0	0	0	0	0
7	560	lower	0	1	0	1	1	1	1	1	1	1
8	590	lower	0	1	1	0	1	1	0	1	1	1
9	650	lower	0	0	0	0	0	0	0	0	0	0
10	900	higher	1	1	0	1	1	1	1	1	1	1
11	1010	higher	1	1	1	1	1	1	1	1	1	1
12	1010	higher	1	1	0	1	1	1	1	1	1	1
13	1150	higher	0	1	1	1	1	1	1	1	1	1
14	1160	higher	1	1	1	1	1	1	1	1	1	1
15	1190	higher	1	1	0	1	1	1	1	0	1	1
16	1420	higher	1	1	0	1	1	1	1	1	1	1
17	2100	higher	0	0	0	0	0	0	0	0	0	0
18	2220	higher	1	1	1	1	1	1	0	1	1	1

Aural detection methods enable detection of northern pika occupancy in densely vegetated habitats

Supplementary Data SD6



Bar plots indicating the number of sites where each vocalization type was heard by using auditory observation (blue) and playback (yellow) methods. Overall result shows the number of present sites including detection at arrival (gray). Darker and lighter colors indicate lower- and higher-elevation sites, respectively.