

RESEARCH ARTICLE

Why does noise reduce response to alarm calls? Experimental assessment of masking, distraction and greater vigilance in wild birds

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Abstract

1. Environmental noise from anthropogenic and other sources affects many aspects of animal ecology and behaviour, including acoustic communication. Acoustic masking is often assumed in field studies to be the cause of compromised communication in noise, but other mechanisms could have similar effects.
2. We tested experimentally how background noise disrupted the response to conspecific alarm calls in wild superb fairy-wrens, *Malurus cyaneus*, assessing the effects of acoustic masking, distraction and changes in vigilance. We first examined the birds' response to alarm-call playbacks accompanied by different amplitudes of background noise that overlapped the calls in acoustic frequency. We then scored and videoed their response to alarm calls in two types of background noise, that did or did not overlap call frequency, but were broadcast at a constant amplitude.
3. Birds were less likely to flee to alarm calls in higher amplitudes of overlapping noise, demonstrating that noise itself compromised communication independently of environmental correlates. Background noise affected the response only if it overlapped in frequency with the alarm calls, implying that the effect was not due to distraction. Further, birds were equally vigilant during background noise of overlapping or non-overlapping frequency, indicating that the lack of response to alarm calls in overlapping noise was not due to enhanced vigilance and awareness that there was no predator.
4. We conclude that alarm-call reception was compromised by masking, a mechanism that is often assumed but rarely tested in an ecological context. Masking compromised reception of high-frequency "aerial" alarm calls and so could reduce survival in background noise of similar frequency. While anthropogenic noise, which is often of lower frequency, is unlikely to affect communication with these calls, it could affect reception of acoustic cues of danger, or other conspecific or heterospecific alarm calls.

KEYWORDS

acoustic communication, alarm call, ambient noise, anthropogenic noise, anti-predator behaviour, birds

1 | INTRODUCTION

Environmental noise affects many aspects of animal behaviour, and is of conservation interest because of the growing global problem of anthropogenic noise. Anthropogenic noise from resource extraction, industry and transportation is now pervasive in both aquatic and terrestrial ecosystems around the world, and can disturb wildlife and affect the physiology, behaviour and fitness of species in diverse taxa (Barber, Crooks, & Fristrup, 2010; Cox, Brennan, Gerwing, Dudas, & Juanes, 2018; Kunc, McLaughlin, & Schmidt, 2016; Morley, Jones, & Radford, 2014; Shannon et al., 2015; Slabbekoorn et al., 2010). The most commonly studied behavioural impact of anthropogenic noise is that on acoustic communication, where the empirical focus has been on effects rather than mechanisms (Chan, Giraldo-Perez, Smith, & Blumstein, 2010; Cox et al., 2018; Francis & Barber, 2013; Morris-Drake, Bracken, Kern, & Radford, 2017; Radford, Kerridge, & Simpson, 2014; Shannon et al., 2015; Slabbekoorn & den Boer-Visser, 2006). However, understanding mechanisms is important because it can help predict effects of noise and suggest methods to ameliorate these effects (Francis & Barber, 2013). For example, remediation aimed at reducing acoustic masking may not solve problems of distraction or aversion (Luo, Siemers, & Koselj, 2015; Senzaki, Kadoya, Francis, Ishiyama, & Nakamura, 2018). Here, we focus on the mechanisms by which environmental noise can affect response to acoustic signals and thereby compromise communication.

Animals communicate acoustically for many reasons, and environmental noise—including anthropogenic noise—can affect both signaller and receiver behaviour. Animals produce a wide range of acoustic signals to communicate about, for instance, reproductive status, territory ownership, hunger, food and danger (Bradbury & Vehrencamp, 2011). However, most studies investigating impacts of noise have considered bird song or other signals used to attract mates and defend space (Aubin & Jouventin, 1998; Brumm, 2004; Brumm & Slabbekoorn, 2005; Read, Jones, & Radford, 2014; Shannon et al., 2015). Signallers may change their singing behaviour—for instance, the time of day that they vocalize and acoustic characteristics of songs—as a consequence of noise, and receiver responses to song can be compromised in noisy conditions (Brumm & Zollinger, 2013; Halfwerk et al., 2011; Moseley et al., 2018; Read et al., 2014; Slabbekoorn & den Boer-Visser, 2006). Far less work has considered how noise affects other types of acoustic communication, including anti-predator signalling (Kern & Radford, 2016; Lowry, Lill, & Wong, 2012; Morris-Drake et al., 2017; Potvin, Mulder, & Parris, 2014). Compromised response to signals about predators is likely to reduce survival, so it is important to understand constraints imposed by noise (Templeton, Zollinger, & Brumm, 2016).

Alarm calls are used to warn others of danger, but relatively little research has examined the impact of noise on the response by listeners to these acoustic signals. Many mammals and birds produce acoustic alarm signals in response to predators, using them to warn others of imminent danger or to recruit assistance when

mobbing threatening species (Hollén & Radford, 2009; Klump & Shalter, 1984; Zuberbühler, 2009). Conspecific receivers benefit from being warned of danger (Caro, 2005), and many animals also benefit by eavesdropping on heterospecific alarm calls (Magrath, Haff, Fallow, & Radford, 2015). The few experiments so far show that additional noise can compromise both conspecific and heterospecific receiver responses to alarm calls. For instance, nestling tree swallows (*Tachycineta bicolor*) crouched and stopped calling in response to parental alarm calls during quiet conditions, but failed to do so during playback of white noise (McIntyre, Leonard, & Horn, 2014). Similarly, great tits (*Parus major*) usually approached a loudspeaker playing conspecific mobbing calls during simultaneous playback of low-amplitude but not high-amplitude traffic noise (Templeton et al., 2016). Responses to heterospecific alarm calls can be similarly compromised: dwarf mongooses (*Helogale parvula*) were less likely to flee to tree squirrel (*Paraxerus cepapi*) alarm calls during traffic-noise playback compared to ambient-sound playback (Morris-Drake et al., 2017), and northern cardinals (*Cardinalis cardinalis*) failed to respond to tufted titmice (*Baeolophus bicolor*) alarm calls in noisier locations near roads (Grade & Sieving, 2016).

Auditory masking is the most commonly evoked mechanism to explain why noise compromises communication. Masking occurs when there is an increase in the threshold for detection or discrimination of an acoustic signal because of noise at similar acoustic frequencies (Moore, 2012). Laboratory studies find that the response to a test signal declines monotonically with increasing amplitude of noise (partial masking), until there is no longer any response (complete masking; Brumm & Slabbekoorn, 2005; Dooling & Blumenrath, 2013; Lohr, Wright, & Dooling, 2003). Partial masking implies that the signal is detectable but the detail is hard to discriminate, while complete masking implies that there is no detection of the signal (Clark et al., 2009; Kleist, Guralnick, Cruz, & Francis, 2016). Clearly, masking does occur in the laboratory with the appropriate frequency and amplitude of noise, but the ecological question is how masking or other mechanisms compromise communication in the wild. In general, acoustic communication becomes more difficult as the amplitude of background sound increases for birds, marine mammals, fish and amphibians, which is consistent with masking (Brumm & Slabbekoorn, 2005; Clark et al., 2009; Dooling, West, & Leek, 2009; Fay & Megela-Simmons, 1999), but does not exclude other mechanisms.

While auditory masking is often assumed to be the primary mechanism by which noise reduces the response to acoustic signals including alarm calls, other mechanisms, including receiver distraction or changes in vigilance, could have similar effects on response (Brumm & Slabbekoorn, 2005; Francis & Barber, 2013; Senzaki et al., 2018). Noise could distract receivers and so compromise responses to other sensory input (Chan & Blumstein, 2011; Chan et al., 2010). For instance, road-noise playback disrupted the adaptive responses of dwarf mongooses to predator faeces; since there was no auditory element to the predator cue, this cross-modal effect of noise implies distraction (Morris-Drake, Kern, & Radford, 2016). Similarly,

fathead minnows (*Pimephales promelas*) subjected to motorboat noise failed to respond to conspecific chemical alarm cues (Hasan, Crane, Ferrari, & Chivers, 2018), and crabs distracted by boat noise responded with a greater delay to a silent looming object (Chan et al., 2010). However, we know of no direct tests of whether acoustic distraction disrupts communication. Noise might also affect communication if it prompts animals to increase vigilance, as is found, for example, in chaffinches (*Fringilla coelebs*), house sparrows (*Passer domesticus*), California ground squirrels (*Otospermophilus beecheyi*) and dwarf mongooses (Kern & Radford, 2016; Meillère, Brischoux, & Angelier, 2015; Quinn, Whittingham, Butler, & Cresswell, 2006; Rabin, Coss, & Owings, 2006). This increased vigilance could then affect the response to alarm calls, because an animal has greater personal knowledge. Animals may therefore respond less to playback of alarm calls because they know that there is no predator nearby (Morris-Drake et al., 2017). Overall, discriminating the effects of masking, distraction and increased vigilance is difficult because all three can predict a decline in responses to alarm calls in the presence of noise.

In this study, we investigate experimentally how and why background environmental noise reduces the response of wild superb fairy-wrens (*Malurus cyaneus*) to conspecific alarm calls. Fairy-wrens give aerial alarm calls to predatory birds in flight and, under quiet conditions, almost always flee to cover after multi-element versions of these calls, which signal immediate danger (Fallow & Magrath, 2010). This conspicuous behaviour provides a clear and sensitive assay of response to an acoustic signal when birds are challenged with different amplitudes and types of noise. We therefore began with an experiment to quantify the reduction in response to aerial alarm calls under different amplitudes of background noise that overlapped calls in frequency. We then assessed in a second experiment why noise reduced the alarm-call response, broadcasting either overlapping or non-overlapping noise and quantifying vigilance. Together, these experiments allowed us to test the potential mechanisms of masking, distraction and greater vigilance.

2 | MATERIALS AND METHODS

2.1 | Study species and site

Superb fairy-wrens are small passerines (9–10 g) that forage primarily on the ground (Higgins, Peter, & Steele, 2001). They are territorial cooperative breeders that usually breed from September to January, although outside the breeding season they often join groups and move across adjacent territories (Rowley, 1965). The sexes can be distinguished by plumage and bill colour after gaining adult colours (Higgins et al., 2001). Fairy-wrens produce a variety of alarm calls, including aerial alarm calls to predatory birds in flight, mobbing calls to terrestrial and perched predators, distress calls when captured, and whining calls when mobbing cuckoos (Colombelli-Négrel, Robertson, Sulloway, & Kleindorfer, 2010; Feeney et al., 2013; Magrath, Pitcher, & Gardner, 2007; Rowley & Russell, 1997). Their aerial alarm calls are composed of repeated elements, each about 100 ms long and with a

mean peak frequency of about 9.1 kHz (Magrath, Pitcher, & Gardner, 2007). Fairy-wrens convey greater urgency of danger in their aerial alarm calls by including more elements; multi-element calls prompt immediate flight to cover, and birds remain in cover for longer when calls contain more elements (Fallow & Magrath, 2010).

We studied an individually colour-banded population of fairy-wrens in the Australian National Botanic Gardens in Canberra (−35.279°S, 149.109°E). The population has been studied for over 30 years, and the birds are accustomed to people (Cockburn, Brouwer, Margraf, Osmond, & Pol, 2016). Fairy-wrens occur throughout most of the 40 ha Gardens, which contain natural woodland, areas planted with Australian native plants, and lawn. Collared sparrowhawks (*Accipiter cirrhocephalus*) and pied currawongs (*Strepera graculina*) are locally common predators of fairy-wrens (Magrath, Pitcher, & Gardner, 2009).

2.2 | Overview of experimental design

We used playback experiments on fairy-wrens to investigate the effect of background noise on the response to conspecific aerial alarm calls. In Experiment 1, we examined how different amplitudes of background noise affected the response to alarm calls. The background noise in this experiment overlapped in frequency with the alarm calls, so all potential mechanisms affecting response—masking, distraction and greater vigilance—were possible. In Experiment 2, we tested which of these potential mechanisms could explain the noise effect found in Experiment 1 (Results). To do so, we assessed how background-noise playbacks with different frequencies, but a constant amplitude affected responses to alarm calls and vigilance. Birds were never exposed to real predators, individuals that fled to playbacks returned to feed within 1 min, and we observed no adverse effects on birds.

2.3 | Sound-file preparation

Ambient sound recorded at the study site was used to prepare 16 unique background-noise playbacks. We recorded 16 examples of ambient sound under relatively quiet conditions in the Gardens, using a Sennheiser ME62 omnidirectional microphone and a Marantz PMD670 recorder sampling at 44.1 kHz and 16 bits. We then excised any distinct foreground sounds, such as bird calls or nearby human noise, with remaining sections joined using the zero-crossing function in Adobe Audition CS6, to produce background noise that was 20 s long. This background noise was then filtered in Raven Pro 1.5 to produce two types of background noise that differed in frequency range (Figure 1). “Overlapping” noise ranged from 6 to 10 kHz, which covers the range in peak frequency of fairy-wren aerial alarm calls (8.6–9.5 kHz; Magrath, Pitcher, & Gardner, 2007). “Non-overlapping” noise ranged from 2 to 6 kHz, so it had the same bandwidth as “Overlapping” noise but did not overlap in frequency with the alarms. This Non-overlapping frequency range is well within typical passerine hearing (Dooling, 2004), and fairy-wrens respond to heterospecific alarm calls that fall completely within this range

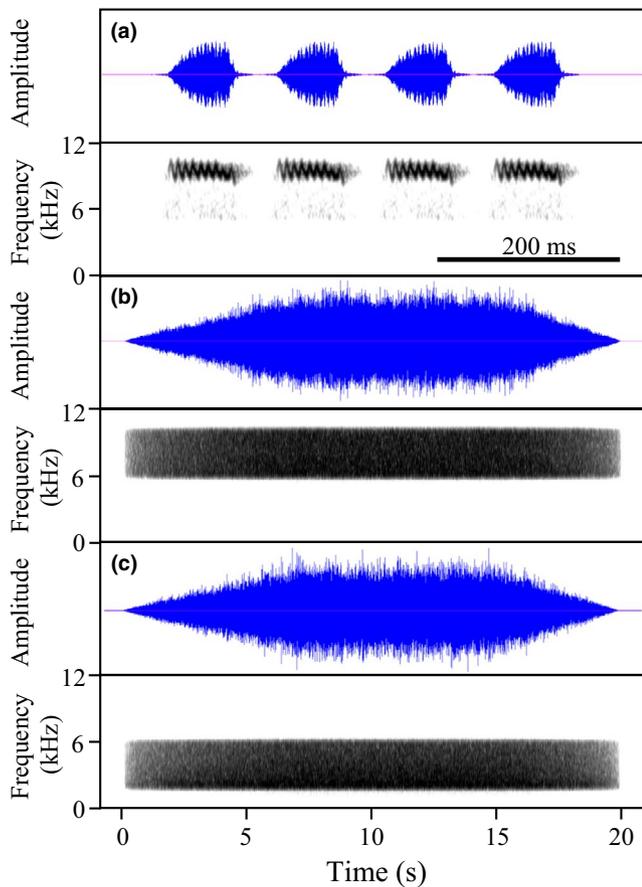


FIGURE 1 Sounds used during playback experiments: (a) fairy-wren aerial alarm call, (b) Overlapping noise (6–10 kHz), and (c) Non-overlapping noise (2–6 kHz), each showing the waveform (above) and spectrogram (below). Note the different timescale for the alarm call. Noise playbacks were calibrated as needed in experiments, and some treatments entailed alarm calls mixed with noise. Spectrograms were prepared in Raven Pro 1.5 and used a Blackman window type, 5.8 ms window size and 95% overlap

(Magrath & Bennett, 2012; Magrath et al., 2009). These 20 s noise tracks were faded in for 7 s and faded out for 5 s, to avoid abrupt changes in amplitude that might startle birds. The middle 8 s of each background-noise track was of relatively constant amplitude, with a standard deviation of average power over each second of less than 0.5 dB. Finally, we adjusted the amplitude of background-noise files to produce the required target broadcast amplitude (below), as measured with a Brüel & Kjær 2240 sound level meter at 10 m. All calibrations and field playbacks used the same set of equipment: a Roland Edirol R-05 HR digital recorder, connected to a custom amplifier, and a Peerless 810921 tweeter speaker (frequency response 2–11 kHz). Recordings of broadcast noise showed that it was roughly flat within the given range. All the equipment was mounted around the observer's waist as a mobile playback system, as used in previous experiments on fairy-wrens (e.g. Magrath, Haff, McLachlan, & Iqic, 2015).

We prepared 16 playbacks of fairy-wren aerial alarm calls, each recorded from a different individual (Figure 1). Calls were prompted

using gliding model pied currawongs or collared sparrowhawks (Fallow & Magrath, 2010; Magrath, Pitcher, & Gardner, 2007), and recorded using a Sennheiser ME66 or ME67 directional microphone and Marantz PMD670 recorder sampling at 44.1 kHz and 16 bits. A single element was pasted at a natural interval to form four-element alarm calls (Fallow & Magrath, 2010), which ranged from 0.43 to 0.97 s, depending on the duration of individual elements. We used four-element alarm calls because they indicate urgent danger and prompt immediate flight to cover (Fallow & Magrath, 2010). Sound below 5 kHz was filtered out using Raven Pro 1.5. We broadcast alarm calls so that elements had a mean amplitude of 52 dB SPL at 10 m, which is similar to the mean natural amplitude at that distance (mean 56.5 dB at 4–7 m, implying 48.5 to 53.4 at 10 m; Magrath, Pitcher, & Gardner, 2007). Alarm-call amplitude was determined by re-recording playbacks at 10 m, along with a calibration tone that had its amplitude measured with the Brüel & Kjær 2240 sound level meter. The alarm-call files were then adjusted iteratively to achieve the target amplitude.

As described in detail below, experiments used playbacks of alarm calls alone, background noise alone and alarm calls mixed with background noise. Each type of playback was broadcast from the single speaker; we used Adobe Audition CS6 to mix the calibrated alarm-call and noise files into mono files, such that alarm calls occurred during the period of stable, maximum noise amplitude. A single-speaker design has been used successfully in previous studies of the effects of noise on communication (e.g. Templeton et al., 2016), allowed use of our mobile playback system, and ensured that the signal and noise came from the same direction and distance, which prevented any spatial release of masking or unwanted variation in distance. The time when the alarm call started in a specific track (at 10, 11 or 12 s) was generated randomly by Excel, to reduce the possibility that birds would habituate to a predictable temporal pattern.

2.4 | Field methods

Playbacks used a matched design, with all treatments broadcast at 16 locations spread across the study site. As in several previous experiments on fairy-wrens (e.g. Magrath et al., 2009; Magrath & Bennett, 2012), we used a unique set of playbacks at each location, and so used location as the unit of replication. We matched by location, rather than specific individual, because in the non-breeding season individuals can wander over several territories, and because location controls for ecological variables such as predator activity and ambient sound. The matched design means any differences among locations will affect all treatments equally. It is possible that individual differences add variance to the results, but the randomised design ensures that this would not introduce bias. Playbacks at a given location always entailed more than one individual in Experiment 1 (non-breeding season; 4–7 individuals per location) and did so in 12/16 locations in Experiment 2 (breeding season; 1–4 individuals per location). Of the 60 individuals in Experiment 1, 12 received playbacks at two locations and one at three locations, but no bird received a specific treatment

more than once. All 33 individuals in Experiment 2 received playbacks at only one location, as the birds were then territorial.

To ensure treatment order did not confound the response, such as through habituation, playback order was generated randomly in R at each location, but constrained over each experiment so that any one treatment was presented roughly evenly at each order. In Experiment 1, with seven treatments, this meant that each treatment almost always occurred 2 or 3 times at each order, and in Experiment 2, with five treatments, each treatment almost always occurred 3 or 4 times at each order. To reduce the risk of habituation to alarm calls, we broadcast at most two playbacks at each location on a given day, with at least 1 hr between them. In practice, playbacks at a location were carried over a mean of 9.8 days (range 6–15) in Experiment 1 and 8.3 days (range 3–18) in Experiment 2, further reducing the risk of habituation. Response to the alarm-call playback was scored as fleeing to cover or not. We used the same score for playbacks to noise alone, taken from 10 to 12 s after the beginning of playback, so that it matched in time the playbacks that included alarm calls. All playbacks were carried out and scored by YZ, so there were no inter-observer issues to control.

Playbacks to wild individuals have the advantage of ecological validity, but the disadvantage that it is not possible to control fully the acoustic environment. We addressed this issue by using location as the unit of replication, restricting playbacks to relatively quiet periods and quantifying the ambient sound after every playback. We then calculated the contribution of ambient sound to the total amplitude experienced by the focal bird and found that it was negligible (Supporting Information Appendix S1).

2.5 | Experiment 1: effect of noise amplitude on the response to alarm-call playback

To investigate the effect of different amplitudes of noise on the response to aerial alarm calls, we broadcast alarm calls in combination with “Overlapping” noise (6–10 kHz). Birds received seven treatments: (a) aerial alarm call alone at 52 dB; aerial alarm call of 52 dB mixed with (b) 52 dB, (c) 55 dB, (d) 58 dB or (e) 61 dB Overlapping noise; and Overlapping noise alone at (f) 52 dB and (g) 61 dB. All dB levels refer to the amplitude at 10 m from the loudspeaker, which was similar to the distance to the focal bird during playbacks (mean 10.8 ± 1.0 SD, measured with a laser rangefinder). The aerial alarm call alone was the positive control, measuring response in the absence of experimental noise, and the two playbacks of noise alone tested whether the birds fled to the noise itself at the lowest and highest amplitudes used in the experiment. Playbacks were done in the non-breeding season, between 12 May and 27 June 2017.

2.6 | Experiment 2: the mechanism of how noise disrupts alarm-call responses

To investigate the mechanism causing the reduced response to alarm calls in noise (Results, Experiment 1), we broadcast alarm calls mixed with noise with different frequency ranges, while simultaneously

videoing the birds' responses. Birds received five playback treatments: (a) aerial alarm call alone (peak frequency c. 9.1 kHz) at 52 dB; (b) aerial alarm call at 52 dB mixed with “Overlapping” noise (6–10 kHz) at 58 dB; (c) aerial alarm call at 52 dB mixed with “Non-overlapping” noise (2–6 kHz) at 58 dB; (d) “Overlapping” noise alone at 58 dB; and (e) “Non-overlapping” noise alone at 58 dB. All amplitudes were measured at 10 m, which was similar to the mean playback distance of $11.2 \text{ m} \pm 1.0$ SD, and again, the aerial alarm call and noise-alone playbacks were controls. Background noise was set to 58 dB because this was the lowest amplitude at which birds no longer responded to alarm calls of 52 dB in Experiment 1 (Results). As well as scoring in the field whether the focal bird fled or not, we videoed birds with a Panasonic HC-V770M video camera (50 frames per s, resolution 1920 × 1080p) mounted on the observer's shoulder and subsequently quantified the focal bird's vigilance in the 10 s of noise immediately before the alarm playback in the mixed tracks. The videos were scored blind, frame by frame by YZ, after the names of video files were re-assigned by others and the sound was muted. Vigilance was scored as the proportion of frames when the bird had its head up, rather than oriented towards the ground (where fairy-wrens glean food). Head position is an indirect measure of vigilance, but is a commonly used metric that correlates with exposure to greater risk in birds, such as being at the edge compared to the centre of a flock (Beauchamp, 2015). In at least some species, a raised head position has been shown experimentally to increase the probability of detecting danger (e.g. Tisdale & Fernández-Juricic, 2009). Playbacks were done in the breeding season, between 21 November 2017 and 1 January 2018.

The immediate response to playback and preceding vigilance allowed a test of the three mechanisms of how noise could affect the response to alarm calls. The flee response was used to discriminate acoustic masking from distraction. Acoustic masking predicts that birds will flee to alarm calls during Non-overlapping noise but not during Overlapping noise, while distraction implies a reduced response during either type of noise, compared to alarm calls alone, because both noise playbacks were of the same amplitude and lie well within avian hearing (Dooling, 2004). A similar logic has been used to discriminate the effects of masking and distraction on the foraging performance of bats in the presence of noise (Luo et al., 2015), and female frogs responding to male mating calls (Senzaki et al., 2018). A reduced response specifically during Overlapping noise could, however, also arise if birds were more vigilant during Overlapping noise, and so were aware that no predator was nearby and were therefore less likely to flee (Morris-Drake et al., 2017). We therefore quantified vigilance during the background noise immediately before the alarm playback to test whether differences in vigilance could explain observed responses.

2.7 | Statistical analysis

Each experiment entailed repeated measures at 16 locations, so we used matched statistical tests. Analysis of whether birds fled or not used Cochran Q tests for overall differences among treatments and

McNemar tests for paired comparisons, as both tests are designed for matched, dichotomous data (Siegel & Castellan, 1988). Vigilance was measured as a continuous variable, so we used a paired *t*-test to compare vigilance during Overlapping and Non-overlapping noise. We used the R statistical package (R Core Team, 2017), including the coin package for Cochran Q and McNemar tests (Hothorn, Hornik, Wiel, & Zeileis, 2008) and base package for *t*-tests.

3 | RESULTS

3.1 | Experiment 1: effect of noise amplitude on the response to alarm-call playback

None of the 16 fairy-wrens fled to cover during playbacks of Overlapping noise alone at either the low- or high-amplitude extremes (Figure 2). This means that any fleeing by birds in response to the mixed playbacks is due to the alarm calls and not noise.

Fairy-wrens were less likely to flee to cover to alarm calls as the amplitude of Overlapping noise increased (Figure 2). Almost all birds fled to playback of alarm calls alone, but this proportion was reduced with added noise, so that overall the response to alarm calls was affected by noise (all five treatments with alarm calls: Cochran's Q test, $Q = 38.4$, $df = 4$, $p < 0.001$). In addition, an increasing amplitude of noise in mixed treatments led to a monotonic reduction in the response to alarm calls (four mixed treatments with alarm calls and noise playback: $Q = 21.6$, $df = 3$, $p < 0.001$). Birds no longer fled to alarm calls mixed with 58 or 61 dB noise, and pairwise comparisons revealed a significant drop in response from 55 to 58 dB (Figure 2), consistent with a masking threshold between these values but potentially caused by any mechanism.

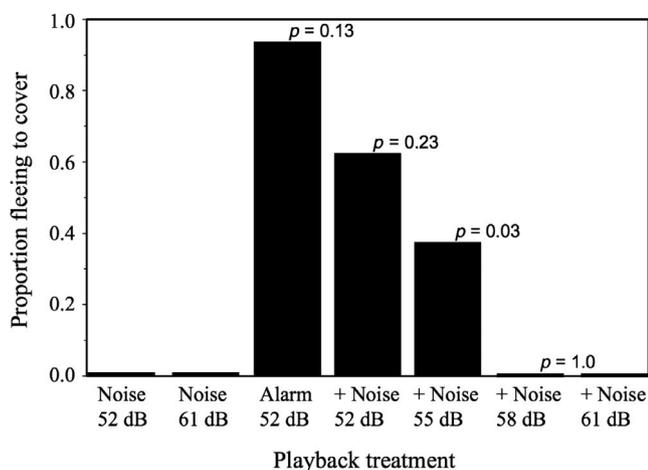


FIGURE 2 Proportion of fairy-wrens that fled to cover to Overlapping noise alone (Noise; 6–10 kHz), alarm call alone (Alarm; mean peak frequency 9.1 kHz) and a 52 dB alarm call mixed with different levels of Overlapping noise (+Noise). Probability values are from McNemar tests of differences between adjacent columns, with increasing amplitude of Overlapping noise. $N = 16$ locations for each treatment

3.2 | Experiment 2: the mechanism of how noise disrupts alarm-call responses

The effect of noise on the response to alarm calls was consistent with acoustic masking rather than distraction (Figure 3). While only 1 of 16 birds fled to cover in response to alarm calls mixed with Overlapping noise, all 16 birds fled in response to alarm calls mixed with Non-overlapping noise (McNemar test: exact 2-tailed, $p = 0.004$). Consistent with the previous experiment, all birds fled to alarm calls alone and none fled to Overlapping noise alone. In addition, no birds fled to Non-overlapping noise alone.

There was no significant difference in our measure of vigilance that could account for the different response of birds to alarm calls mixed with Overlapping and Non-overlapping noise (Figure 4). Birds were equally vigilant during Overlapping and Non-overlapping noise immediately before alarm calls (paired *t*-test, $t_{15} = 0.383$, $p = 0.96$).

4 | DISCUSSION

Background noise affected the response to alarm calls in wild fairy-wrens, probably only by acoustic masking rather than through distraction or changes in vigilance. In our first experiment, broadcasting background noise overlapping in frequency with alarm calls, birds were less likely to flee to the alarm calls in higher amplitudes of noise. This shows that noise affected the response to alarm calls, but does not establish the mechanism. In our second experiment, background noise affected the response to alarm calls only if it overlapped in frequency with alarm calls, and not if it was non-overlapping. This implies that the reduced response was not due to

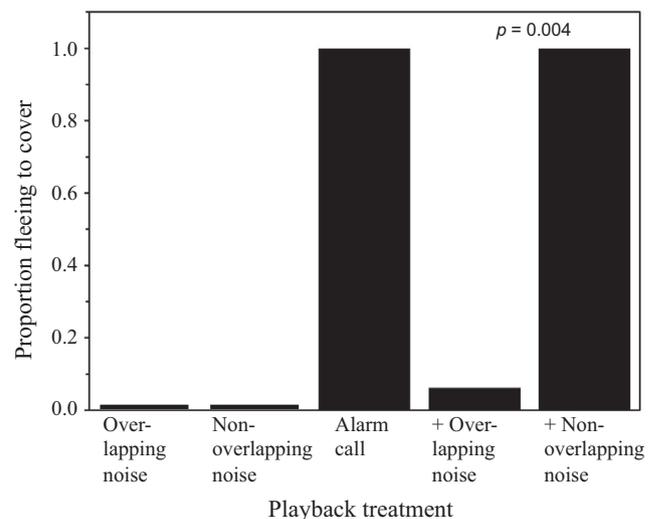


FIGURE 3 Proportion of fairy-wrens that fled to cover to Overlapping (6–10 kHz) and Non-overlapping (2–6 kHz) noise alone, alarm call alone (mean peak frequency 9.1 kHz) and alarm calls mixed with Overlapping or Non-overlapping noise. The probability value is from a McNemar test for a difference in response to alarm calls mixed with Overlapping or Non-overlapping noise. $N = 16$ locations for each treatment

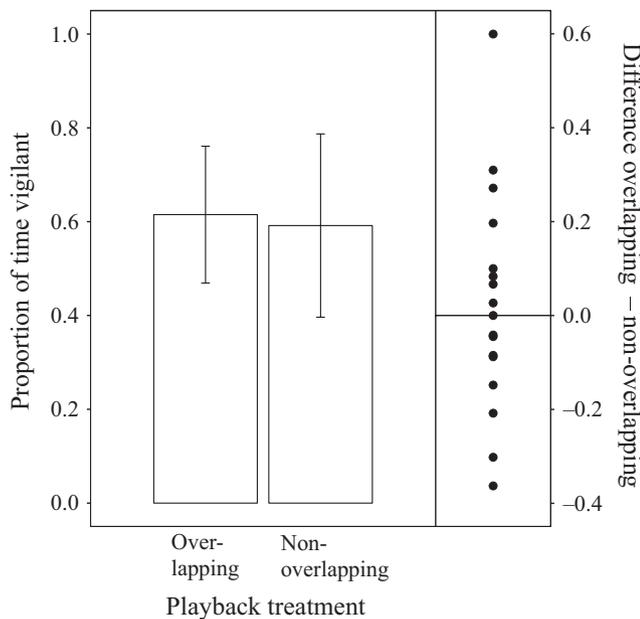


FIGURE 4 Proportion of time fairy-wrens spent vigilant (head up) during the 10 s of noise before alarm calls in playback treatments mixed with Overlapping or Non-overlapping noise. Columns show observed means \pm SD, and the scatterplot shows the difference in proportion of time vigilant during Overlapping minus Non-overlapping noise at each location. $N = 16$ locations for both treatments; 30/32 recordings had a full 10 s sample, while the video was accidentally cut short in two cases (8.3 and 9.4 s) but adjusted for the shorter duration

distraction and was consistent with masking. Finally, birds showed the same level of vigilance in response to Overlapping and Non-overlapping noise, indicating that the lack of response to alarm calls mixed with Overlapping noise was not due to enhanced vigilance and awareness that there was no predator. Together, our results indicate that alarm-call reception was disrupted by acoustic masking, which is something that is likely to be common but, as far as we are aware, not previously shown in experiments on wild birds.

Experiment 1 showed that increased levels of background noise led to a reduced probability of fleeing to cover after aerial alarm calls. Almost all birds fled to alarms when not mixed with noise, but response declined monotonically until there was no response when mixed with noise at or above 58 dB. This experiment demonstrates that background noise itself affected the response to alarm calls, because playbacks were matched by location, eliminating potential confounds such as location-specific variation in predator density or effects of noisy infrastructure unrelated to noise itself (Antze & Koper, 2018; Grade & Sieving, 2016). Furthermore, birds did not flee to noise-only playbacks, showing that fleeing was to alarm calls and not the accompanying noise. Short-term playbacks of noise potentially provoke startle responses (Francis & Barber, 2013), but we used locally recorded background sound that faded in over 7 s to a modest maximum amplitude, which may explain the lack of startle. Aerial alarm calls in fairy-wrens signal immediate threat from airborne predators (Magrath, Pitcher, & Gardner, 2007), so that any

noise-induced reduction in response to these alarm calls could prove fatal. More broadly, noise could affect the response to any acoustic sources of information about danger, including conspecific and heterospecific alarm calls, and acoustic cues from predators themselves (Barber et al., 2010; Corcoran, Barber, & Conner, 2009; Haff & Magrath, 2010; Magrath, Pitcher, & Dalziell, 2007).

The differences in immediate responses to alarm calls in Experiment 2 were consistent with acoustic masking and unlikely to be due to distraction. First, masking occurs primarily when noise is of a similar acoustic frequency to the signal (Brumm & Slabbekoorn, 2005), whereas distraction could occur regardless of the acoustic frequency of noise (Francis & Barber, 2013). Indeed, distraction can even be cross-modal, such as when noise affects the perception of visual and olfactory stimuli (Chan & Blumstein, 2011; Chan et al., 2010; Morris-Drake et al., 2016). Consistent with masking but not distraction, fairy-wrens almost never fled to alarm calls when the experimental noise overlapped the frequency of alarm calls, but always fled when the noise was of lower frequency, despite being the same amplitude and well within passerine hearing (Dooling, 2004). One caveat is that Overlapping noise might be more distracting than Non-overlapping noise, but this seems unlikely since birds showed no clear anti-predator response to either noise type alone and did not look up more to Overlapping noise. Second, our results on wild birds replicate patterns of masking in birds in the laboratory. Specific critical ratios for masking vary among species, frequencies, and type of noise and signal (Dooling, 2004), and so are difficult to compare, but our results parallel laboratory studies, which show a monotonic decline in response with increasing amplitude of overlapping white noise, eventually reaching a threshold beyond which there is no response to the signal (Dooling & Blumenrath, 2013; Lohr et al., 2003). By contrast, the degree of distraction is not necessarily related in a simple way to amplitude, and can be affected by the type of noise and the task being distracted (Banbury, Macken, Tremblay, & Jones, 2001; Naguib, 2013; Smith, 1989). For example, speech and intermittent noise, but not regular white noise, commonly affect human performance on cognitive tasks; in our experiment, both background-noise types had similar features aside from a difference in frequency range.

Although the patterns of fleeing in Experiment 2 could in principle arise from increased vigilance during Overlapping noise, there was no evidence for this mechanism. Specifically, if playback of Overlapping noise prompted greater vigilance, then birds might not respond to the alarm calls because they already knew that there was no predator nearby, not because the calls were inaudible (Morris-Drake et al., 2017). Contrary to this explanation, blind-scoring of video revealed that there was no difference in the proportion of time birds had their head up during Overlapping and Non-overlapping noise immediately before the alarm call. The lack of a difference in our measure of vigilance is perhaps surprising, as noise that masks alarm signals might be expected to make animals warier and so increase vigilance, and thereby enhance detection of predators or acoustic signals or cues of danger (Beauchamp, 2015; Lynch et al., 2015). In support of these possibilities in other species, broadcast of traffic noise prompted black-tailed prairie

dogs (*Cynomys ludovicanus*) to increase vigilance, which led to an earlier response during human approach (Shannon, Angeloni, Wittemyer, Frstrup, & Crooks, 2014; Shannon, Crooks, Wittemyer, Frstrup, & Angeloni, 2016); and California ground squirrels (*Spermophilus beecheyi*) were more vigilant when near noisy wind turbines, and in those locations also responded more strongly to audible alarm playbacks (Rabin et al., 2006). However, in our study vigilance was unaffected by the acoustic frequency of noise, and therefore did not account for the pattern of fleeing to alarm calls, arguing against this alternative to masking.

Our study was designed to examine specifically the mechanisms by which noise reduced the response of fairy-wrens to their aerial alarm calls, but there are many opportunities for research on how noise could affect alarm communication in this and other species. First, masking alone appeared to be the critical mechanism reducing response to alarm calls in our experiments, which means that low-frequency urban noise should not affect reception of these high-frequency alarm calls. However, mobbing, distress and whining alarm calls include lower frequencies (Colombelli-Négre et al., 2010; Feeney et al., 2013) and so could be partially masked by urban noise. Furthermore, low-frequency urban noise could mask the sound of predators themselves, and so eliminate alternative information on danger when high-frequency natural noises, such as cicada choruses, mask aerial alarm calls. The effect of anthropogenic noise could therefore depend on the type of alarm call and concurrent sources of noise. Second, we examined only the response to alarm calls, but it is also relevant to consider alarm-call production (Brumm & Zollinger, 2013). While there is limited opportunity to increase the effectiveness of urgent alarm calls by adjusting the location or timing of calling in noisier conditions, birds might modify alarm calls to make them more audible. For example, callers can increase the amplitude of alarm calls (Templeton et al., 2016), modify acoustic structure to increase their audible range (Potvin et al., 2014), or beam calls to specific receivers (Yorzinski & Patricelli, 2010). Nonetheless, regardless of sender adaptations, receivers gain information from both conspecific and heterospecific alarm calls from varying distances, so that noise rather than sensory acuity will often limit the distance over which calls can be received (Klump, 1996).

We conclude that masking, rather than distraction or changes in vigilance, accounted for the noise-induced disruption of alarm communication in wild fairy-wrens. It would be valuable to test explicitly for masking in other species, including both alarm calls and other cues of danger. More broadly, we advocate studies on how noise affects animal behaviour, and suggest that multiple mechanisms, including distraction and vigilance effects, will affect many species and are not mutually exclusive (e.g. Luo et al., 2015; Senzaki et al., 2018). Evaluating mechanisms is important in understanding the evolution of communication, predicting the effects of anthropogenic noise, and suggesting ways to ameliorate the consequences of this global pollutant (Francis & Barber, 2013).

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AUTHORS' CONTRIBUTIONS

R.D.M. and A.N.R. conceived the study, all authors designed the experiments and contributed to writing, and Y.Z. carried out fieldwork and analyses.

DATA ACCESSIBILITY

Data deposited in the Dryad Digital Repository: <https://doi.org/10.5061/dryad.b4f2m1t> (Zhou, Radford, & Magrath, 2019).

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REFERENCES

- Antze, B., & Koper, N. (2018). Noisy anthropogenic infrastructure interferes with alarm responses in Savannah sparrows (*Passerculus sandwichensis*). *Royal Society Open Science*, *5*, 172168.
- Aubin, T., & Jouventin, P. (1998). Cocktail-party effect in king penguin colonies. *Proceedings of the Royal Society of London B: Biological Sciences*, *265*, 1665–1673. <https://doi.org/10.1098/rspb.1998.0486>
- Banbury, S. P., Macken, W. J., Tremblay, S., & Jones, D. M. (2001). Auditory distraction and short-term memory: Phenomena and practical implications. *Humans Factors*, *43*, 12–29. <https://doi.org/10.1518/001872001775992462>
- Barber, J. R., Crooks, K. R., & Frstrup, K. M. (2010). The costs of chronic noise exposure for terrestrial organisms. *Trends in Ecology & Evolution*, *25*, 180–189. <https://doi.org/10.1016/j.tree.2009.08.002>
- Beauchamp, G. (2015). *Animal vigilance: Monitoring predators and competitors*. Amsterdam, the Netherlands: Elsevier.
- Bradbury, J. W., & Vehrencamp, S. L. (2011). *Principles of animal communication*. Sunderland, MA: Sinauer Associates.
- Brumm, H. (2004). The impact of environmental noise on song amplitude in a territorial bird. *Journal of Animal Ecology*, *73*, 434–440. <https://doi.org/10.1111/j.0021-8790.2004.00814.x>
- Brumm, H., & Slabbekoorn, H. (2005). Acoustic communication in noise. *Advances in the Study of Behavior*, *35*, 151–209.
- Brumm, H., & Zollinger, S. A. (2013). Avian vocal production in noise. In H. Brumm (Ed.), *Animal communication and noise* (pp. 187–227). Berlin, Germany: Springer.
- Caro, T. (2005). *Antipredator defenses in birds and mammals*. Chicago, IL: University of Chicago Press.
- Chan, A.-A.-Y.-H., & Blumstein, D. T. (2011). Attention, noise, and implications for wildlife conservation and management. *Applied Animal Behaviour Science*, *131*, 1–7. <https://doi.org/10.1016/j.applanim.2011.01.007>
- Chan, A.-A.-Y.-H., Giraldo-Perez, P., Smith, S., & Blumstein, D. T. (2010). Anthropogenic noise affects risk assessment and attention: The distracted prey hypothesis. *Biology Letters*, *6*, 458–461. <https://doi.org/10.1098/rsbl.2009.1081>
- Clark, C. W., Ellison, W. T., Southall, B. L., Hatch, L., Van Parijs, S. M., Frankel, A., & Ponirakis, D. (2009). Acoustic masking in marine ecosystems: Intuitions, analysis, and implication. *Marine*

- Ecology Progress Series*, 395, 201–222. <https://doi.org/10.3354/meps08402>
- Cockburn, A., Brouwer, L., Margraf, N., Osmond, H. L., & van de Pol, M. (2016). Superb fairy-wrens: Making the worst of a good job. In W. D. Koenig & J. L. Dickinson (Eds.), *Cooperative breeding in vertebrates: Studies of ecology, evolution, and behavior* (pp. 133–149). Cambridge, UK: Cambridge University Press.
- Colombelli-Négrel, D., Robertson, J., Sulloway, F. J., & Kleindorfer, S. J. (2010). Extended parental care of fledglings: Parent birds adjust anti-predator response according to predator type and distance. *Behaviour*, 147, 853–870. <https://doi.org/10.1163/000579510X495771>
- Corcoran, A. J., Barber, J. R., & Conner, W. E. (2009). Tiger moth jams bat sonar. *Science*, 325, 325–327. <https://doi.org/10.1126/science.1174096>
- Cox, K., Brennan, L. P., Gerwing, T. G., Dudas, S. E., & Juanes, F. (2018). Sound the alarm: A meta-analysis on the effect of aquatic noise on fish behavior and physiology. *Global Change Biology*, 24, 3105–3116. <https://doi.org/10.1111/gcb.14106>
- Dooling, R. J. (2004). Audition: Can birds hear everything they sing? In P. Marler & H. Slabbekoorn (Eds.), *Nature's music: The science of birdsong* (pp. 206–225). San Diego, CA: Elsevier.
- Dooling, R. J., & Blumenrath, S. H. (2013). Avian sound perception in noise. In H. Brumm (Ed.), *Animal communication and noise* (pp. 229–250). Berlin, Germany: Springer.
- Dooling, R. J., West, E. W., & Leek, M. R. (2009). Conceptual and computational models of the effects of anthropogenic noise on birds. *5th International Conference on Bioacoustics 2009, Proceedings of the Institute of Acoustics*, pp. 99–106. Holywell Park.
- Fallow, P. M., & Magrath, R. D. (2010). Eavesdropping on other species: Mutual interspecific understanding of urgency information in avian alarm calls. *Animal Behaviour*, 79, 411–417. <https://doi.org/10.1016/j.anbehav.2009.11.018>
- Fay, R. R., & Megela-Simmons, A. (1999). The sense of hearing in fishes and amphibians. In R. R. Fay (Ed.), *Comparative hearing: Fish and amphibians* (pp. 269–318). Berlin, Germany: Springer.
- Feeney, W. E., Medina, I., Somveille, M., Heinsohn, R., Hall, M. L., Mulder, R. A., ... Langmore, N. E. (2013). Brood parasitism and the evolution of cooperative breeding in birds. *Science*, 342, 1506–1508. <https://doi.org/10.1126/science.1240039>
- Francis, C. D., & Barber, J. R. (2013). A framework for understanding noise impacts on wildlife: An urgent conservation priority. *Frontiers in Ecology & the Environment*, 11, 305–313. <https://doi.org/10.1890/120183>
- Grade, A. M., & Sieving, K. E. (2016). When the birds go unheard: Highway noise disrupts information transfer between bird species. *Biology Letters*, 12, 20160113. <https://doi.org/10.1098/rsbl.2016.0113>
- Haff, T. M., & Magrath, R. D. (2010). Vulnerable but not helpless: Nestlings are fine-tuned to cues of approaching danger. *Animal Behaviour*, 79, 487–496. <https://doi.org/10.1016/j.anbehav.2009.11.036>
- Halfwerk, W., Bot, S., Buix, J., van der Velde, M., Komdeur, J., ten Cate, C., & Slabbekoorn, H. (2011). Low-frequency songs lose their potency in noisy urban conditions. *Proceedings of the National Academy of Sciences of the United States of America*, 108, 14549–14554. <https://doi.org/10.1073/pnas.1109091108>
- Hasan, M. R., Crane, A. L., Ferrari, M. C., & Chivers, D. P. (2018). A cross-modal effect of noise: The disappearance of the alarm reaction of a freshwater fish. *Animal Cognition*, 21, 419–424. <https://doi.org/10.1007/s10071-018-1179-x>
- Higgins, P. J., Peter, J. M., & Steele, W. K. (2001). *Handbook of Australian, New Zealand and Antarctic birds. Volume 5: Tyrant-flycatchers to chats*. Melbourne, Vic.: Oxford University Press.
- Hollén, L. I., & Radford, A. N. (2009). The development of alarm call behaviour in mammals and birds. *Animal Behaviour*, 78, 791–800. <https://doi.org/10.1016/j.anbehav.2009.07.021>
- Hothorn, T., Hornik, K., van de Wiel, M. A., & Zeileis, A. (2008). Implementing a class of permutation tests: The Coin package. *Journal of Statistical Software*, 28, 1–23.
- Kern, J. M., & Radford, A. N. (2016). Anthropogenic noise disrupts use of vocal information about predation risk. *Environmental Pollution*, 218, 988–995. <https://doi.org/10.1016/j.envpol.2016.08.049>
- Kleist, N. J., Guralnick, R. P., Cruz, A., & Francis, C. D. (2016). Anthropogenic noise weakens territorial response to intruder's songs. *Ecosphere*, 7, e01259. <https://doi.org/10.1002/ecs2.1259>
- Klump, G. M. (1996). Bird communication in the noisy world. In D. E. Kroodsma & E. H. Miller (Eds.), *Ecology and evolution of acoustic communication in birds* (pp. 321–338). Ithaca, NY: Cornell University Press.
- Klump, G., & Shalter, M. (1984). Acoustic behaviour of birds and mammals in the predator context. I. Factors affecting the structure of alarm signals. II. The functional significance and evolution of alarm signals. *Zeitschrift Für Tierpsychologie*, 66, 189–226. <https://doi.org/10.1111/j.1439-0310.1984.tb01365.x>
- Kunc, H. P., McLaughlin, K. E., & Schmidt, R. (2016). Aquatic noise pollution: implications for individuals, populations, and ecosystems. *Proceedings of the Royal Society of London B: Biological Sciences*, 283, 20160839. <https://doi.org/10.1098/rspb.2016.0839>
- Lohr, B., Wright, T. F., & Dooling, R. J. (2003). Detection and discrimination of natural calls in masking noise by birds: Estimating the active space of a signal. *Animal Behaviour*, 65, 763–777. <https://doi.org/10.1006/anbe.2003.2093>
- Lowry, H., Lill, A., & Wong, B. B. (2012). How noisy does a noisy miner have to be? Amplitude adjustments of alarm calls in an avian urban 'adapter'. *PLoS One*, 7, e29960. <https://doi.org/10.1371/journal.pone.0029960>
- Luo, J., Siemers, B. M., & Koselj, K. (2015). How anthropogenic noise affects foraging. *Global Change Biology*, 21, 3278–3289. <https://doi.org/10.1111/gcb.12997>
- Lynch, E., Northrup, J. M., McKenna, M. F., Anderson, C. R., Angeloni, L., & Wittemyer, G. (2015). Landscape and anthropogenic features influence the use of auditory vigilance by mule deer. *Behavioral Ecology*, 26, 75–82. <https://doi.org/10.1093/beheco/aru158>
- Magrath, R. D., & Bennett, T. H. (2012). A micro-geography of fear: Learning to eavesdrop on alarm calls of neighbouring heterospecifics. *Proceedings of the Royal Society B: Biological Sciences*, 279, 902–909. <https://doi.org/10.1098/rspb.2011.1362>
- Magrath, R. D., Haff, T. M., Fallow, P. M., & Radford, A. N. (2015). Eavesdropping on heterospecific alarm calls: From mechanisms to consequences. *Biological Reviews*, 90, 560–586.
- Magrath, R. D., Haff, T. M., McLachlan, J. R., & Igic, B. (2015). Wild birds learn to eavesdrop on heterospecific alarm calls. *Current Biology*, 25, 1–4.
- Magrath, R. D., Pitcher, B. J., & Dalziel, A. H. (2007). How to be fed but not eaten: Nestling responses to parental food calls and the sound of a predator's footsteps. *Animal Behaviour*, 74, 1117–1129. <https://doi.org/10.1016/j.anbehav.2007.01.025>
- Magrath, R. D., Pitcher, B. J., & Gardner, J. L. (2007). A mutual understanding? Interspecific responses by birds to each other's aerial alarm calls. *Behavioral Ecology*, 18, 944–951. <https://doi.org/10.1093/beheco/arm063>
- Magrath, R. D., Pitcher, B. J., & Gardner, J. L. (2009). An avian eavesdropping network: Alarm signal reliability and heterospecific response. *Behavioral Ecology*, 20, 745–752. <https://doi.org/10.1093/beheco/arp055>
- McIntyre, E., Leonard, M. L., & Horn, A. G. (2014). Ambient noise and parental communication of predation risk in tree swallows, *Tachycineta bicolor*. *Animal Behaviour*, 87, 85–89. <https://doi.org/10.1016/j.anbehav.2013.10.013>
- Meillère, A., Brischoux, F., & Angelier, F. (2015). Impact of chronic noise exposure on antipredator behavior: An experiment in breeding house sparrows. *Behavioral Ecology*, 26, 569–577. <https://doi.org/10.1093/beheco/aru232>
- Moore, B. C. J. (2012). *An introduction to the psychology of hearing*. Leiden, the Netherlands: Brill.

- Morley, E. L., Jones, G., & Radford, A. N. (2014). The importance of invertebrates when considering the impacts of anthropogenic noise. *Proceedings of the Royal Society of London B: Biological Sciences*, 281, 20132683. <https://doi.org/10.1098/rspb.2013.2683>
- Morris-Drake, A., Bracken, A. M., Kern, J. M., & Radford, A. N. (2017). Anthropogenic noise alters dwarf mongoose responses to hetero-specific alarm calls. *Environmental Pollution*, 223, 476–483. <https://doi.org/10.1016/j.envpol.2017.01.049>
- Morris-Drake, A., Kern, J. M., & Radford, A. N. (2016). Cross-modal impacts of anthropogenic noise on information use. *Current Biology*, 26, R911–R912. <https://doi.org/10.1016/j.cub.2016.08.064>
- Moseley, D. L., Derryberry, G. E., Phillips, J. N., Danner, J. E., Danner, R. M., Luther, D. A., & Derryberry, E. P. (2018). Acoustic adaptation to city noise through vocal learning by a songbird. *Proceedings of the Royal Society of London B: Biological Sciences*, 285, 20181356.
- Naguib, M. (2013). Living in a noisy world: Indirect effects of noise on animal communication. *Behaviour*, 150, 1069–1084. <https://doi.org/10.1163/1568539X-00003058>
- Potvin, D. A., Mulder, R. A., & Parris, K. M. (2014). Silvereyes decrease acoustic frequency but increase efficacy of alarm calls in urban noise. *Animal Behaviour*, 98, 27–33. <https://doi.org/10.1016/j.anbehav.2014.09.026>
- Quinn, J., Whittingham, M., Butler, S., & Cresswell, W. (2006). Noise, predation risk compensation and vigilance in the chaffinch *Fringilla coelebs*. *Journal of Avian Biology*, 37, 601–608. <https://doi.org/10.1111/j.2006.0908-8857.03781.x>
- R Core Team (2017). *R: A language and environment for statistical computing*. Vienna, Austria: R Core Team. R Foundation for Statistical Computing.
- Rabin, L. A., Coss, R. G., & Owings, D. H. (2006). The effects of wind turbines on antipredator behavior in California ground squirrels (*Spermophilus beecheyi*). *Biological Conservation*, 131, 410–420. <https://doi.org/10.1016/j.biocon.2006.02.016>
- Radford, A. N., Kerridge, E., & Simpson, S. D. (2014). Acoustic communication in a noisy world: Can fish compete with anthropogenic noise? *Behavioral Ecology*, 25, 1022–1030.
- Read, J., Jones, G., & Radford, A. N. (2014). Fitness costs as well as benefits are important when considering responses to anthropogenic noise. *Behavioral Ecology*, 25, 4–7. <https://doi.org/10.1093/beheco/art102>
- Rowley, I. (1965). The life history of the superb blue wren *Malurus cyaneus*. *Emu*, 64, 251–297.
- Rowley, I., & Russell, E. M. (1997). *Fairy-wrens and grasswrens: Maluridae*. Oxford, UK: Oxford University Press.
- Senzaki, M., Kadoya, T., Francis, C. D., Ishiyama, N., & Nakamura, F. (2018). Suffering in receivers: Negative effects of noise persist regardless of experience in female anurans. *Functional Ecology*, 32, 2054–2064. <https://doi.org/10.1111/1365-2435.13130>
- Shannon, G., Angeloni, L. M., Wittemyer, G., Fristrup, K. M., & Crooks, K. R. (2014). Road traffic noise modifies behaviour of a keystone species. *Animal Behaviour*, 94, 135–141. <https://doi.org/10.1016/j.anbehav.2014.06.004>
- Shannon, G., Crooks, K. R., Wittemyer, G., Fristrup, K. M., & Angeloni, L. M. (2016). Road noise causes earlier predator detection and flight response in a free-ranging mammal. *Behavioral Ecology*, 27, 1370–1375. <https://doi.org/10.1093/beheco/aww058>
- Shannon, G., McKenna, M. F., Angeloni, L. M., Crooks, K. R., Fristrup, K. M., Brown, E., ... Wittemyer, G. (2015). A synthesis of two decades of research documenting the effects of noise on wildlife. *Biological Reviews*, 91, 982–1005. <https://doi.org/10.1111/brv.12207>
- Siegel, S., & Castellan, N. J. (1988). *Nonparametric statistics for the behavioral sciences*. New York, NY: McGraw Hill.
- Slabbekoorn, H., Bouton, N., van Opzeeland, I., Coers, A., ten Cate, C., & Popper, A. N. (2010). A noisy spring: The impact of globally rising underwater sound levels on fish. *Trends in Ecology & Evolution*, 25, 419–427. <https://doi.org/10.1016/j.tree.2010.04.005>
- Slabbekoorn, H., & den Boer-Visser, A. (2006). Cities change the songs of birds. *Current Biology*, 16, 2326–2331. <https://doi.org/10.1016/j.cub.2006.10.008>
- Smith, A. (1989). A review of the effects of noise on human performance. *Scandinavian Journal of Psychology*, 30, 185–206. <https://doi.org/10.1111/j.1467-9450.1989.tb01082.x>
- Templeton, C. N., Zollinger, S. A., & Brumm, H. (2016). Traffic noise drowns out great tit alarm calls. *Current Biology*, 26, R1173–R1174. <https://doi.org/10.1016/j.cub.2016.09.058>
- Tisdale, V., & Fernández-Juricic, E. (2009). Vigilance and predator detection vary between avian species with different visual acuity and coverage. *Behavioral Ecology*, 20, 936–945. <https://doi.org/10.1093/beheco/arp080>
- Yorzinski, J. L., & Patricelli, G. L. (2010). Birds adjust acoustic directionality to beam their antipredator calls to predators and conspecifics. *Proceedings of the Royal Society of London B: Biological Sciences*, 277, 923–932.
- Zhou, Y., Radford, A. N., & Magrath, R. D. (2019). Data from: Why does noise reduce response to alarm calls? Experimental assessment of masking, distraction and greater vigilance in wild birds. *Dryad Digital Repository*, <https://doi.org/10.5061/dryad.b4f2m1t>
- Zuberbühler, K. (2009). Survivor signals: The biology and psychology of animal alarm calling. *Advances in the Study of Behavior*, 40, 277–322.

SUPPORTING INFORMATION

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