

RESEARCH ARTICLE

Sensitive high-frequency hearing in earless and partially eared harlequin frogs (*Atelopus*)

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ABSTRACT

Harlequin frogs, genus *Atelopus*, communicate at high frequencies despite most species lacking a complete tympanic middle ear that facilitates high-frequency hearing in most anurans and other tetrapods. Here, we tested whether *Atelopus* are better at sensing high-frequency acoustic sound compared with other eared and earless species in the Bufonidae family, determined whether middle ear variation within *Atelopus* affects hearing sensitivity and tested potential hearing mechanisms in *Atelopus*. We determined that at high frequencies (2000–4000 Hz), *Atelopus* are 10–34 dB more sensitive than other earless bufonids but are relatively insensitive to mid-range frequencies (900–1500 Hz) compared with eared bufonids. Hearing among *Atelopus* species is fairly consistent, evidence that the partial middle ears present in a subset of *Atelopus* species do not convey a substantial hearing advantage. We further demonstrate that *Atelopus* hearing is probably not facilitated by vibration of the skin overlying the normal tympanic membrane region or the body lung wall, leaving the extratympanic hearing pathways in *Atelopus* enigmatic. Together, these results show *Atelopus* have sensitive high-frequency hearing without the aid of a tympanic middle ear and prompt further study of extratympanic hearing mechanisms in anurans.

KEY WORDS: Extratympanic hearing, Auditory brainstem recordings, Bufonidae, Lung hearing

INTRODUCTION

Most tetrapods have tympanic middle ears, which allow them to better sense their acoustic environment on land (Christensen-Dalsgaard and Carr, 2008; Manley, 2010; Manley and Sienknecht, 2013); however, a minority of tetrapods rely on alternative methods for sensing acoustic stimuli (Hartline, 1971; Christensen et al., 2012; Mason and Narins, 2002; Wever, 1975). Yet, these alternative hearing methods are only efficient at low frequencies (Hartline, 1971; Christensen et al., 2012; Mason and Narins, 2002; Wever, 1975), and tetrapods without a tympanic middle ear that both communicate and have effective hearing above 1000 Hz are uncommon (Boistel et al., 2011). In this study, we investigated the high-frequency hearing sensitivity and potential hearing mechanisms in harlequin frogs, genus *Atelopus*, which lack a

complete tympanic middle ear (Pereyra et al., 2016) but are known to communicate at high frequencies (1750–3780 Hz; Cocroft et al., 1990; Boistel et al., 2011).

Atelopus may be unique among bufonids in their ability to hear high frequencies (above 1500 Hz) without a middle ear. Earlessness, lack of all middle ear structures, has evolved at least 38 times in anurans (Pereyra et al., 2016), and is associated with a 16–25 dB decrease in hearing sensitivity above 1000 Hz in non-*Atelopus* bufonids (Womack et al., 2017). Likewise, two other anuran species show a 25 dB decrease in hearing sensitivity above 1000 Hz when the tympanic membrane is removed [*Hyliola regilla* (= *Hyla regilla*) and *Dryophytes versicolor* (= *Hyla versicolor*); Lombard and Straughan, 1974]. Yet, hearing tests on a limited number of *Atelopus* species show sensitivity to high-frequency sound above 1000 Hz. *Atelopus chiriquiensis* is only 5 dB less sensitive than the eared species *H. regilla* (Jaslow and Lombard, 1996) and three other *Atelopus* species [*Atelopus flavescens*, *Atelopus* sp. (Nusagandi) and *Atelopus lozanoi* [= *Atelopus* sp. (Chingaza)]]; have sensitive hearing well above 1000 Hz (Lindquist et al., 1998). However, *Atelopus* hearing has not been compared with the hearing of closely related eared and earless species and the extratympanic hearing pathways used by earless *Atelopus* lack experimental verification.

Researchers have proposed several anuran extratympanic hearing pathways, but only one has been experimentally verified and also has the potential to affect high-frequency hearing: the lung pathway. This pathway, which transfers airborne sound waves that vibrate the body lung wall to the inner ear (Narins et al., 1988), mediates hearing sensitivity at frequencies up to 1000 Hz (Ehret et al., 1990; Hetherington and Lindquist, 1999), and the body lung wall of three *Atelopus* species [*A. flavescens*, *Atelopus* sp. (Nusagandi) and *A. lozanoi*] were shown to vibrate at even higher frequencies (~2500 Hz) that relate to their species' dominant call frequency (Lindquist et al., 1998). Thus, the lung pathway is a strong candidate for a potential extratympanic hearing mechanism in *Atelopus* (Lindquist et al., 1998; Boistel et al., 2011). However, no one has experimentally tested this pathway in any species that has high-frequency hearing sensitivity but lacks a tympanic middle ear.

In addition to effective extratympanic hearing pathways, some *Atelopus* species have a partial middle ear that may provide a high-frequency hearing benefit. Although all *Atelopus* lack a complete middle ear, a small clade of *Atelopus* species have either retained or regained a partial middle ear that has the middle ear bone and cavity but lacks a tympanic membrane (Lindquist et al., 1998; Boistel et al., 2011; Pereyra et al., 2016). These middle ear components may function relatively normally, with the skin overlying the attachment to the middle ear bone (herein referred to as the otic epidermis) vibrating in response to airborne sound and transferring those vibrations through the middle ear bone to the inner ear. Comparison of a single partially eared *Atelopus* species (*A. flavescens*) and two earless *Atelopus* species [*Atelopus* sp. (Nusagandi) and *A. lozanoi*] found the partially eared species was 8–13 dB more sensitive to

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airborne sound from 2000 to 2500 Hz (Lindquist et al., 1998). High-frequency hearing in partially eared *Atelopus* may be mediated by this incomplete middle ear.

Here, we tested the hearing of three *Atelopus* species [*Atelopus elegans*, *Atelopus* sp. 1 (*spumarius* complex) and *Atelopus* sp. 2 (*spumarius* complex)] to assess their hearing sensitivity and to better understand mechanisms of hearing without a complete tympanic middle ear. First, we describe the ear structures of the three *Atelopus* species. Next, we assessed whether *Atelopus* hearing differs from hearing in other bufonids by comparing the hearing of each *Atelopus* species with previously reported hearing sensitivities of other eared and earless bufonids (Womack et al., 2017). We further assessed hearing differences within *Atelopus* associated with the presence of a partial middle ear by comparing hearing among two partially eared and one earless *Atelopus* species. Last, we aimed to identify the mechanisms of high-frequency hearing in *Atelopus* by manipulating two potential hearing pathways: the otic epidermis and the body lung wall. These studies provide broad hearing comparisons within the family Bufonidae, inform hypotheses of middle ear evolution within *Atelopus*, and test potential extratympanic hearing pathways in anurans.

MATERIALS AND METHODS

Animal collection

Adult animals were collected (earless *Atelopus elegans*, $n=6$; partially eared *Atelopus* sp. 1, $n=9$; partially eared *Atelopus* sp. 2, $n=4$; eared *Rhaebo haematiticus*, $n=4$; eared *Rhinella alata*, $n=4$; eared *Rhinella horribilis*, $n=4$; eared *Rhinella spinulosa*, $n=4$; eared *Rhinella tacana*, $n=2$) from field sites in Ecuador and Peru (Table 1), while an additional eight *Atelopus* sp. 2 individuals were bred in captivity at Centro Jambatu in Ecuador. We measured the mass of each individual animal to the nearest 0.01 g (individual masses available in Dataset 1) using a digital pocket scale (EHA701, Camry Industries Company Ltd, Guangdong, China) and measured the snout–vent length (SVL) of each animal to the nearest 0.1 mm (Table 1; individual SVLs available in Dataset 1) using a dial caliper (31–415–3, Swiss Precision Instruments Inc., Garden Grove, CA, USA). The Institutional Animal Care and Use Committee at Colorado State University approved all experiments (IACUC Protocol no. 12–3484A), and the Ministerio del Ambiente in Ecuador and the Servicio Nacional Forestal y de Fauna Silvestre in Peru approved collection, breeding, research and export permits (Table 1).

Auditory brainstem recordings (ABRs) to test *Atelopus* hearing

We tested the hearing ability of *Atelopus elegans* ($n=6$), *Atelopus* sp. 1 ($n=9$) and *Atelopus* sp. 2 ($n=4$) in Ecuador and Peru using the

same auditory brainstem recording methods detailed in Womack et al. (2016). Briefly, we paralyzed the bufonids with 0.05% succinylcholine chloride (Sigma-Aldrich, St Louis, MO, USA) at a dosage of $7.5 \mu\text{l g}^{-1}$ and then lightly anesthetized animals with a small topical application of 5% benzocaine at the sites of electrode placement. The topical application of 5% benzocaine should have brief, localized effects and wear off shortly after electrode placement. Most animals remained paralyzed throughout the 1–3 h of testing and only received one dose of succinylcholine chloride; however, individuals that showed slight movement (active breathing) during the testing period were given subsequent half-doses of succinylcholine chloride. We subdermally placed differential electrodes over the midbrain and VIIIth (auditory) nerve and placed a third ground electrode within the arm contralateral to the VIIIth nerve (Fig. S1) to measure the electrical signal generated by the VIIIth nerve. We linked the three electrodes to a pre-amplifier (RA4PA, Tucker-Davis Technologies, Alachua, FL, USA) connected to a mobile processor (RM2, Tucker-Davis Technologies) that relayed output and input signals from and to a laptop computer (Mini 210-2180, Hewlett Packard, Palo Alto, CA, USA). We placed bufonids on a wet paper towel and positioned them perpendicular to and 46 cm away from a suspended speaker. We calibrated speaker output with a $\frac{1}{2}$ inch free field microphone (46AE, G.R.A.S. Sound and Vibration A/S, Skovlytoften, Denmark).

We calibrated the experimental set-up using customized software (QuickABR_burst) that controlled stimulus presentation and data acquisition using a mobile processor (RM2 Info). We played 25 ms pure tones, ranging in frequency from 200 to 4000 Hz (Table 2) at 5 dB increments. We averaged response signals over 400 tone bursts and measured the response to a transient generated from a half-cycle 4000 Hz sinusoid at 105 dB sound pressure level (SPL) between every two frequencies to ensure that the auditory responsiveness remained stable throughout the testing session. If the transient response dropped below 75% of the original signal, we omitted all subsequent measurements of that individual from analyses. We visually determined thresholds for each frequency, using the minimum stimulus decibel level that evoked a response signal amplitude of 0.002 mV (roughly twice the noise level) or greater from the auditory nerve.

Vaseline manipulations to test hearing pathways in *Atelopus*

We performed additional ABRs on a subset of *Atelopus* sp. 2 individuals to test potential hearing pathways in a partially eared *Atelopus* species. All manipulation ABRs were recorded exactly as described above with the exception of four frequency omissions (200, 400, 1750 and 2250 Hz) to reduce test length. We chose not to

Table 1. Snout–vent length (SVL), collection country, sites and permit numbers for animals in the study

Species (taxonomic authority)	SVL (mm)	Country	Region	Permit no.
<i>Atelopus elegans</i> (Boulenger 1882)	38.2*	Ecuador	Ecuador, Provincia Esmeraldas, Río Durango, Durango	001-13 IC-FAU-DNB/MA
<i>Atelopus</i> sp. 1 (<i>spumarius</i> complex)	24.4–25*	Ecuador	Ecuador, Provincia Pastaza, Reserva Otoyacu, Río Pucayacu	001-13 IC-FAU-DNB/MA
<i>Atelopus</i> sp. 2 (<i>spumarius</i> complex)	26.6–35.7	Ecuador	Ecuador, Provincia Morona Santiago, San Carlos de Limón (Nueva Principal)	001-13 IC-FAU-DNB/MA
<i>Rhaebo haematiticus</i> Cope 1862	73.1–75.3	Ecuador	Reserva Otokiki, Río Baltazar, Esmeraldas Province	001-13 IC-FAU-DNB/MA
<i>Rhinella alata</i> (Thomiot 1884)	38.5–41.1	Ecuador	Playón de San Francisco (La Ceiba), Esmeraldas Province	001-13 IC-FAU-DNB/MA
<i>Rhinella horribilis</i> (Wiegmann 1833)	77.1–107.0	Ecuador	Unión del Toachi (Chorrera del Diablo), Cotopaxi Province; and in San Francisco (La Ceiba), Esmeraldas Province	001-13 IC-FAU-DNB/MA
<i>Rhinella spinulosa</i> (Wiegmann 1834)	68.1–77.6	Peru	K'iripampa Acopia in Acomayo, Departamento de Cusco	0071-2014-MINAGRI-DGFFS/DGEFFS
<i>Rhinella tacana</i> (Padial, Reichle, McDiarmid and De la Riva 2006)	28.4–30.4	Peru	Quincemil, Departamento de Cusco	0071-2014-MINAGRI-DGFFS/DGEFFS

Asterisks in the SVL column indicate species that have missing data for a subset of individuals.

randomize the order of non-treatment ABRs and Vaseline treatment ABRs because it was not possible to completely remove the Vaseline without stressing the animal between tests. Most animals remained paralyzed throughout the full 1–3 h of testing (including non-manipulation and manipulation ABRs) and only received one dose of succinylcholine chloride. We gave subsequent half-doses of succinylcholine chloride to animals that showed slight movement. However, the click response was monitored throughout both tests to ensure that overall response levels did not drop during the Vaseline ABR or change after a subsequent dose of succinylcholine chloride.

With four *Atelopus* sp. 2 individuals, we tested whether vibration of the otic epidermis was contributing to hearing by covering the otic epidermis on both sides of the head with a thick layer of Vaseline. If vibration of these regions is important to hearing, then the Vaseline applied to these surfaces should affect hearing by weighing the otic epidermis down and affecting its ability to vibrate in response to sound waves. For comparison, we covered the tympanic membranes of four individuals per species of other bufonids (*R. haematiticus*, *R. alata*, *R. horribilis* and *R. spinulosa*) with a thick layer of Vaseline to test how this affected hearing of eared bufonids that rely on vibration of the tympanic membrane. These additional four species were tested in either Ecuador or Peru with the same experimental set-up and protocol as for the *Atelopus* sp. 2 individuals and their unmanipulated ABR results were published in Womack et al. (2017).

With three other *Atelopus* sp. 2 individuals, we tested whether vibration of the body lung wall was involved in the hearing of *Atelopus* sp. 2 by wrapping the body lung wall along with the complete mid-region of the frog's body in a thick layer of Vaseline and cheesecloth. These manipulations are similar to those performed by Hetherington and Lindquist (1999), which resulted in decreased hearing sensitivity in *Bombina orientalis*.

Specimen fixation, histology and 3D reconstruction

After ABRs, we fixed two representatives from the three *Atelopus* species (*A. elegans*, *Atelopus* sp. 1 and *Atelopus* sp. 2) and compared their middle ear morphology with that of two representative individuals from a similarly sized bufonid with a complete tympanic middle ear, *R. tacana*. We killed two individuals of each species (total $n=8$) with 20% topical benzocaine, then decapitated the specimens, preserved the heads in 4% paraformaldehyde (diluted in phosphate-buffered saline from 16% paraformaldehyde solution; Electron Microscopy Sciences, Hatfield, PA, USA) for 24 h, performed three 15 min rinses in phosphate-buffered saline, and finally stored the cranial tissue in 70% ethanol.

We sliced the heads in half (sagittal) to isolate a single ear of each specimen and then decalcified the tissues in 10% EDTA (pH 7.4) for up to 1 week at room temperature. We then put the tissues through a graded ethanol series from 30% to 100%, and embedded them in hydroxypropyl methacrylate (HPMA) plastic (Electron Microscopy Sciences). We drilled holes of 1 mm diameter into the plastic around each tissue, sectioned through the ear structures at 5 μm thickness with a microtome (RM1265, Leica, Wetzlar, Germany), and mounted every other section onto Autofrost Adhesion Microscope Slides (Cancer Diagnostics, Inc., Durham, NC, USA). We then stained the tissue with Eosin and Toluidine Blue (Fisher Scientific, Pittsburgh, PA, USA) and photographed every third section with a dissection microscope (Olympus SZX10) and digital camera (Olympus DP71) for a final distance of 30 μm between imaged sections. We took images with a resolution of 1360 pixels \times 1024 pixels. We aligned the photographed sections using the drilled holes and then 3D modeled and measured ear structures within IMOD 3D (Kremer et al., 1996). For the 3D reconstruction figures (Fig. 1), we smoothed our reconstructions within IMOD 3D.

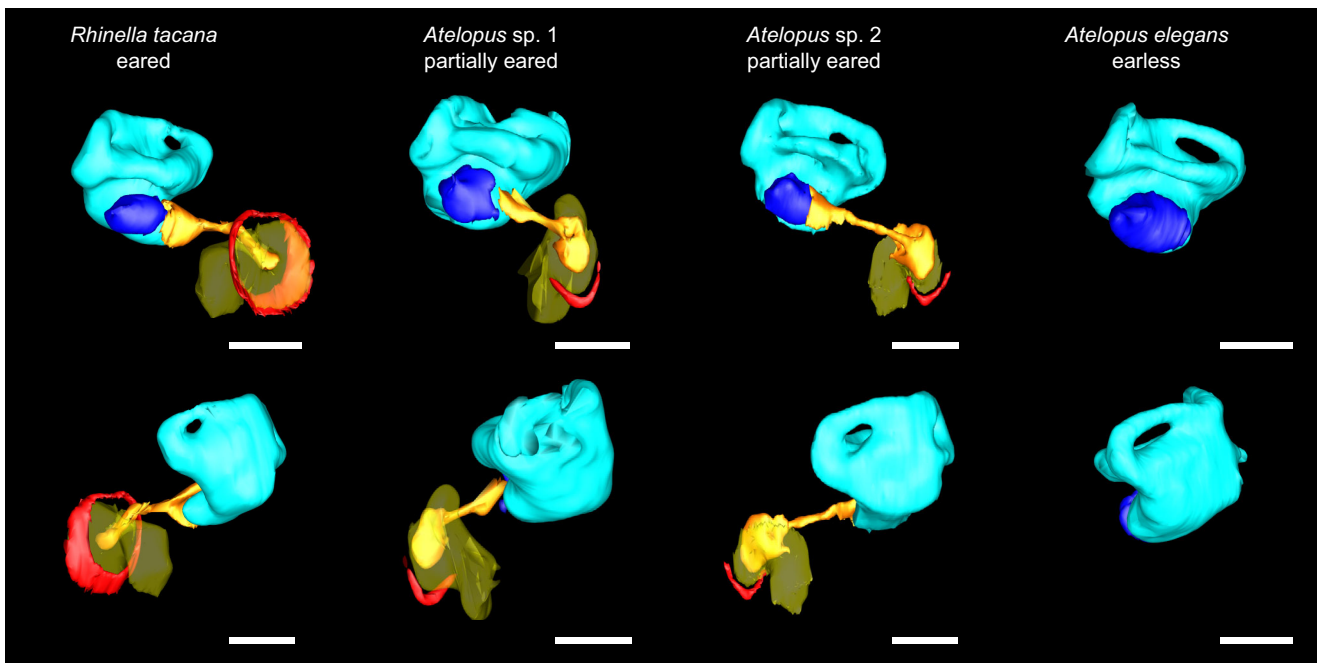


Fig. 1. Middle ear variation within *Atelopus*. 3D reconstructions from histology sections showing variation in middle ear structures among *Atelopus* species and an eared bufonid, *Rhinella tacana*. *Atelopus elegans* has no middle ear structures, while *Atelopus* sp. 1 and *Atelopus* sp. 2 both have all middle ear structures present (columella, Eustachian tube, middle ear cavity, tympanic annulus) except the tympanic membrane. The partially eared species, *Atelopus* sp. 1 and *Atelopus* sp. 2, have an incomplete tympanic annulus and an extended extracolumella, when compared with *R. tacana*. Inner ear, light blue; operculum, blue; columella, orange; Eustachian tube+middle ear cavity, yellow; tympanic annulus, red. Snout–vent length (SVL) of *R. tacana*, 28.4 mm; *Atelopus* sp. 1, 25.0 mm; *Atelopus* sp. 2, 32.0 mm; *A. elegans*, 26.0 mm. Scale bars: 1 mm.

Statistical analysis

We visualized audiograms representing hearing sensitivity of species by graphing the thresholds from the ABRs using the sme (smoothing-splines mixed-effects models) package (<http://CRAN.R-project.org/package=sme>) in R (<http://www.R-project.org/>). We used sme continuous graphs because comparing many species' audiograms was difficult with other graphic representations because of the large number of overlapping data points. We then tested for hearing differences between species and groups of species using a mixed model produced in the package lme4 (Bates et al., 2014) and *post hoc* analyses using the package lsmeans (Lenth, 2016) and lmerTest (<http://CRAN.R-project.org/package=lmerTest>) in R (<http://www.R-project.org/>). Because we wanted to test overall species differences in hearing sensitivity, we did not examine sex differences. Even though males and females may differ in hearing (see Boatright-Horowitz and Simmons, 1995; Miranda and Wilczynski, 2009; Shen et al., 2011), we combined data from the two sexes because of the limited sample size.

First, we tested hearing differences among *Atelopus* species and other earless and eared bufonids using the previously published hearing data for bufonid species measured with identical methods (Womack et al., 2017). We ran a model that had hearing thresholds of all species as the response variable, species, frequency (as a factor) and their interaction as fixed effects, and individual as a random effect. We ran *post hoc* contrasts to compare hearing of each *Atelopus* species with both the average hearing sensitivity of the other earless bufonids and the average hearing sensitivity of the eared bufonids. The least squares means gave us an estimate of the mean hearing threshold for each *Atelopus* species, all other earless bufonids and all eared bufonids at each frequency. We compared differences in those hearing thresholds between groups and adjusted *P*-values using Sidak's method.

Next, we tested whether *Atelopus* with a partial middle ear were more sensitive than earless *Atelopus* species by comparing hearing among our *Atelopus* species. Using the mixed model above, we estimated the least squares means hearing threshold for each of the three *Atelopus* species. We then ran pairwise comparisons of

those least squares means hearing thresholds for each *Atelopus* species at each frequency and adjusted *P*-values using Tukey's method.

Last, we determined the frequencies at which individuals with and without Vaseline treatment differed in sensitivity, using only hearing data from individuals that were tested both with and without the Vaseline treatment. We ran one model that had hearing threshold as the response variable, head Vaseline treatment (yes or no), frequency, species and their three-way interaction term as the fixed effects, and individual as a random effect. We then ran a second model that had hearing threshold as the response variable, body lung wall Vaseline treatment (yes or no), frequency, species and their three-way interaction term as the fixed effects, and individual as a random effect. For each model, we then calculated within-species differences in least squares means of individuals' hearing thresholds with and without Vaseline treatment at all frequencies. The least squares means gave us an estimate of the mean hearing threshold for each species with and without the Vaseline treatment and compared within-species differences in those hearing thresholds. All *post hoc* comparisons were adjusted using Tukey's method for multiple comparisons.

RESULTS

Description of *Atelopus* middle ear structures

We found no evidence of tympanic middle ear structures in *A. elegans*, while *Atelopus* sp. 1 and *Atelopus* sp. 2 both had all middle ear structures present (columella, Eustachian tube, middle ear cavity, tympanic annulus) except the tympanic membrane. However, both *Atelopus* sp. 1 and *Atelopus* sp. 2 had an incomplete tympanic annulus in which only the ventral half was present (Fig. 1). Additionally, both *Atelopus* sp. 1 and *Atelopus* sp. 2 had an extended extracolumella, the most distal portion of the columella that attaches to the tympanic membrane, when compared with a non-*Atelopus* bufonid with a complete middle ear, *R. tacana* (Fig. 1).

Atelopus hearing compared with that of other bufonids

When comparing bufonid hearing thresholds, we found that thresholds differed among species and varied by frequency

Table 2. Estimated least squares means differences between hearing sensitivity thresholds of *Atelopus* species and other earless and eared bufonids

Frequency (Hz)	<i>Atelopus</i> versus eared bufonids (dB)			<i>Atelopus</i> versus other earless bufonids (dB)		
	<i>A. elegans</i>	<i>Atelopus</i> sp. 1	<i>Atelopus</i> sp. 2	<i>A. elegans</i>	<i>Atelopus</i> sp. 1	<i>Atelopus</i> sp. 2
200	-11±4*	-8±4	11±3**	-13±4**	-10±4	9±3*
300	0±4	-1±4	10±3**	-1±4	-1±4	9±3*
400	8±4	9±4	16±3***	0±4	1±5	9±3*
500	4±4	10±4	15±3***	-3±4	4±5	9±3*
700	4±4	14±4**	21±3***	-4±4	6±5	13±3***
900	9±4*	16±4**	22±3***	-3±4	3±5	10±3*
1100	9±4*	15±4**	17±4***	-7±4	-1±5	1±4
1300	10±4*	23±4***	15±3***	-9±4	3±5	-4±4
1500	12±4**	19±4***	12±3***	-8±4	-1±5	-8±3
1750	8±4	16±4***	5±3	-11±4*	-3±5	-15±4***
2000	10±4*	7±4	12±3**	-13±4**	-15±5**	-11±3**
2250	10±4*	2±4	10±3**	-15±4***	-23±5***	-15±3***
2500	10±4*	2±4	4±3	-15±4***	-22±5***	-20±3***
3000	-6±4	-18±4***	-4±3	-22±4***	-34±5***	-21±3***
3500	NA	NA	NA	-22±4***	-30±5***	-16±4***
4000	-14±4***	-23±5***	-2±4	-22±4***	-31±5***	-10±4*

Estimated least squares means (\pm s.e.) rounded to the nearest dB are given for airborne sound sensitivity differences at each frequency, with significant differences between groups in bold (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$). A negative estimated difference indicates that the *Atelopus* species had a lower threshold (were more sensitive) at that frequency than the eared or earless bufonids, while a positive estimated difference indicates that the *Atelopus* species had a higher hearing threshold (were less sensitive) at that frequency than the eared or earless bufonids.

($F_{179,1119,09}=9.41$, $P<0.001$; Fig. 2). At high frequencies, all three *Atelopus* species were more sensitive than earless non-*Atelopus* bufonids (2000–4000 Hz; Fig. 2A, Table 2), and two *Atelopus* species (*A. elegans* and *Atelopus* sp. 1) were more sensitive than eared bufonids at select high frequencies (3000 and 4000 Hz; Fig. 2B, Table 2). Meanwhile, at low–mid-range frequencies, *Atelopus* were generally less sensitive than eared bufonids (Fig. 2B, Table 2). All three *Atelopus* species were less sensitive than eared species from 900 to 1500 Hz, and individual *Atelopus* species were less sensitive than eared species at a number of other frequencies below 2500 Hz (Table 2).

Testing effects of the partial middle ear on *Atelopus* hearing

Overall, the earless *A. elegans* was more sensitive than the two partially eared *Atelopus* species (Fig. 2, Table 3). The earless *A. elegans* was more sensitive than the partially eared *Atelopus* sp. 2 at a range of frequencies (200, 300, 500–900 and 4000 Hz) and was more sensitive than the partially eared *Atelopus* sp. 1 at 1300 Hz (Fig. 2, Table 3). Only at one frequency (3000 Hz) was a partially eared species (*Atelopus* sp. 1) more sensitive than the earless *A. elegans*.

Testing hearing mechanisms in *Atelopus*

Covering the tympanic region with Vaseline affected hearing in ways that varied by species and frequency ($F_{41,390,79}=2.39$, $P<0.001$; Fig. 3A). Covering the otic epidermis of *Atelopus* sp. 2 did not decrease hearing sensitivity at any frequency (Fig. 3A, Table 4). In contrast, covering the tympanic membrane of bufonid species with complete tympanic ears resulted in a 9–29 dB decrease in hearing sensitivity at frequencies ranging from 700 to 3500 Hz (Fig. 3A, Table 4).

Covering the body lung wall with Vaseline did not result in any overall difference in hearing sensitivity for the partially eared *Atelopus* sp. 2 ($F_{1,65}=1.15$, $P=0.288$; Fig. 3B, Table 4), nor did the effects of the Vaseline lung treatment vary by frequency ($F_{11,65}=1.43$, $P=0.183$).

DISCUSSION

Atelopus species are rare examples of tetrapods lacking tympanic middle ears yet sensing and communicating with high-frequency airborne sound. We showed that *Atelopus* have better hearing than other earless bufonids at high frequencies; however, *Atelopus* still have reduced sensitivity at mid-range frequencies in comparison to

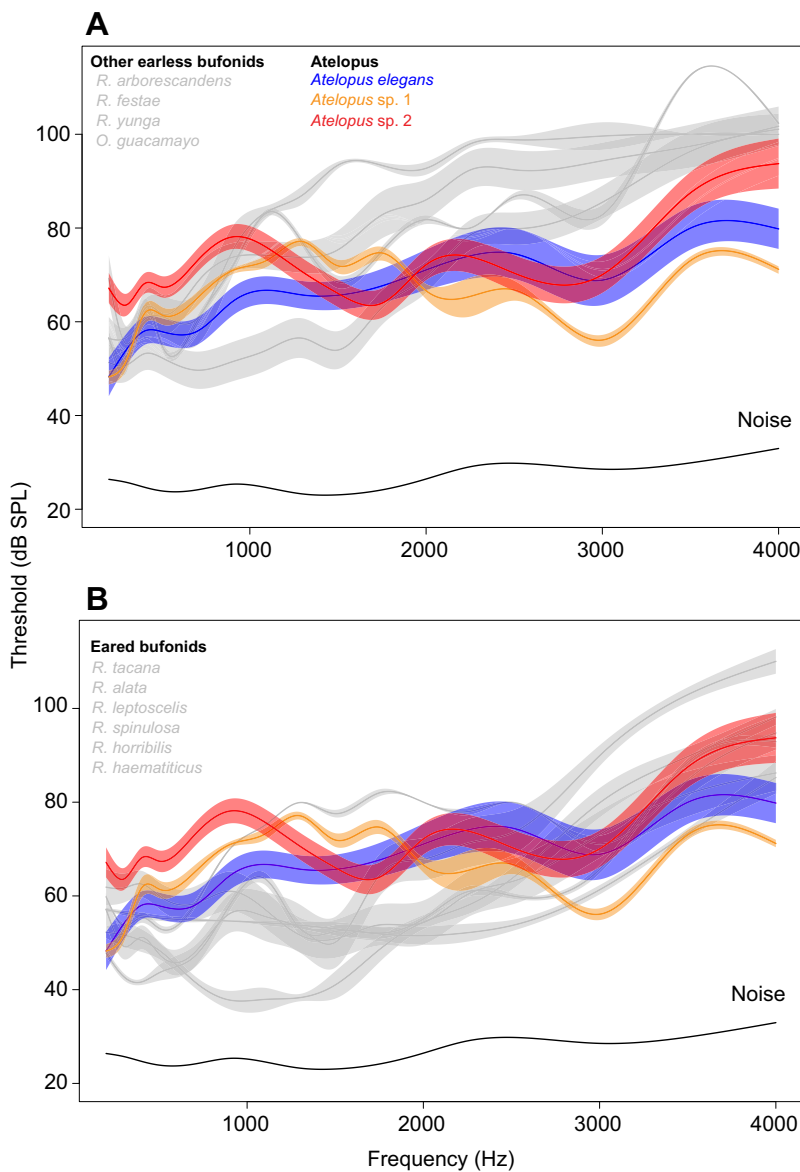


Fig. 2. Hearing differences between *Atelopus* and other eared and earless bufonids.

(A) Audiograms of the three *Atelopus* species (colors) and earless non-*Atelopus* bufonid species (gray). (B) Audiograms of the three *Atelopus* species (colors) and eared non-*Atelopus* bufonid species (gray). Within-chamber noise level is shown by the black line. Lower x-axis thresholds equate to higher hearing sensitivity. All gray species' data are from Womack et al. (2017). Species: *Atelopus elegans* ($N=6$), *Atelopus* sp. 1 ($N=9$) and sp. 2 ($N=4$), *Rhinella arborescandens* ($N=10$), *Rhinella festae* ($N=3$), *Rhinella yunga* ($N=9$), *Rhinella tacana* ($N=3$), *Rhinella alata* ($N=9$), *Rhinella leptoscelis* ($N=3$), *Rhinella spinulosa* ($N=5$), *Rhinella horribilis* ($N=5$), *Rhaebo haematiticus* ($N=5$) and *Osornophryne guacamayo* ($N=8$).

Table 3. Estimated least squares means differences between hearing sensitivity thresholds of partially eared and earless *Atelopus* species

Frequency (Hz)	Earless <i>A. elegans</i> versus partially eared <i>Atelopus</i> (dB)		<i>Atelopus</i> sp. 1 versus sp. 2 (dB)
	<i>Atelopus</i> sp. 1	<i>Atelopus</i> sp. 2	
200	3±5	22±4***	-19±5***
300	0±5	10±4*	-10±5
400	1±5	9±4	-8±5
500	6±5	11±4*	-5±5
700	10±5	18±4***	-8±5
900	6±5	13±4**	-6±5
1100	6±5	8±5	-2±5
1300	13±5*	5±5	8±5
1500	6±5	0±5	6±5
1750	8±5	-3±4	12±5
2000	-3±5	2±4	-4±5
2250	-8±5	0±4	-8±5
2500	-8±5	-5±4	-2±5
3000	-12±5*	1±4	-13±5*
3500	-7±5	6±5	-13±5*
4000	-9±5	13±4**	-21±5***

Estimated least squares means (\pm s.e.) rounded to the nearest dB are given for airborne sound sensitivity differences at each frequency, with significant differences between groups in bold (* $P<0.05$, ** $P<0.01$, *** $P<0.001$). A negative estimated difference indicates that the partially eared *Atelopus* species (*Atelopus* sp. 1 and 2) had a lower threshold (were more sensitive) at that frequency than the earless *Atelopus* species (*A. elegans*), while a positive estimated difference indicates that the partially eared *Atelopus* species had a higher hearing threshold (were less sensitive) at that frequency than the earless *Atelopus* species.

eared bufonids. We found no consistent differences in hearing between our partially eared and earless *Atelopus* species, indicating the partial middle ear found in a small clade of *Atelopus* does not

provide an advantage for airborne sound sensitivity. We also found no evidence that the otic epidermis and the body lung wall function as extratympanic pathways transmitting sound waves to the inner ears of *Atelopus* sp. 2. We discuss our hearing sensitivity data in relation to previous hearing studies on *Atelopus* and other anurans, middle ear lability in *Atelopus*, and hypotheses of extratympanic pathways in *Atelopus* and other anurans.

***Atelopus* hearing in comparison to that of other bufonids**

All *Atelopus* species were 10–31 dB more sensitive to high-frequency airborne sound than other earless bufonids. Even more interesting, this high-frequency hearing sensitivity matches the dominant call frequency of a partially eared species in this study (*Atelopus* sp. 2, d.f.=2250 Hz; Fig. S2), as well as other *Atelopus* species (Cocroft et al., 1990). Thus, even though all *Atelopus* species lack a tympanum and most completely lack a middle ear, they have maintained hearing sensitivity to high-frequency conspecific calls, despite reduced sensitivity at these high frequencies in other earless bufonids (Womack et al., 2017). This study is the first to test hearing differences between earless *Atelopus* and other earless anurans, but our results agree with previous studies on hearing using other *Atelopus* species, which found *Atelopus* were sensitive to high-frequency sound (Jaslow and Lombard, 1996; Lindquist et al., 1998; Boistel et al., 2011). These results all suggest that *Atelopus* have mechanisms for hearing high-frequency sound that other earless anurans lack.

However, these extratympanic hearing mechanisms in *Atelopus* do not seem to function very well at mid-range frequencies (900–1500 Hz). Despite the high-frequency hearing capabilities of *Atelopus*, their mid-range frequency hearing is consistently less sensitive compared with that of eared bufonids. Specializations for high-frequency hearing sensitivity in the extratympanic pathways or

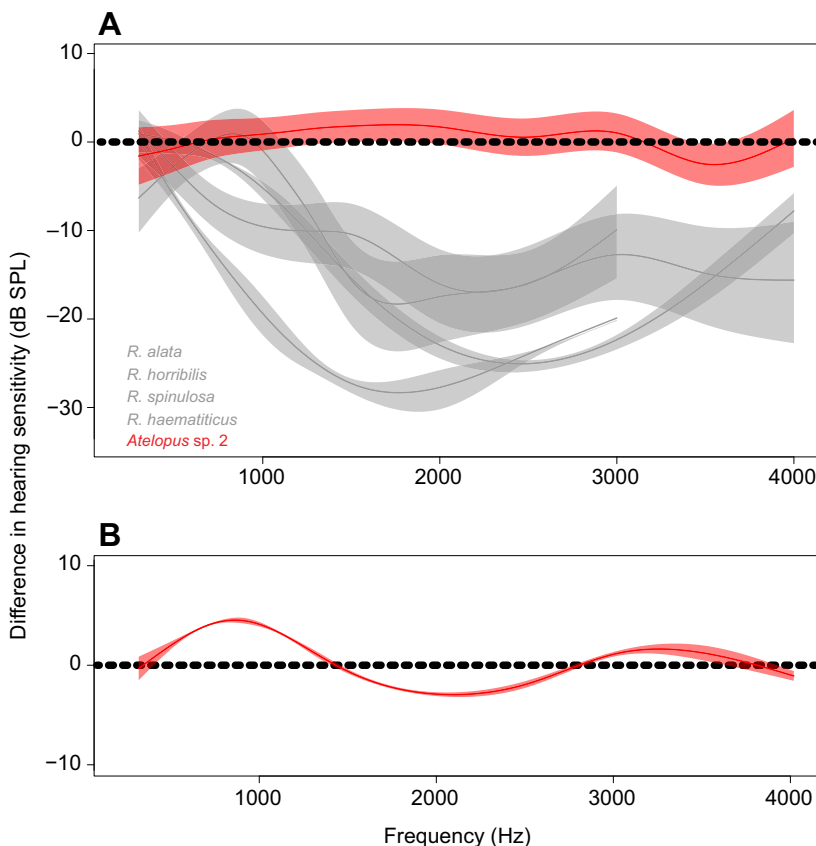


Fig. 3. Vaseline treatment effects on hearing of *Atelopus* sp. 2 and other eared bufonids. (A) Difference in hearing of individuals before and after application of Vaseline to cover the tympanic membrane in eared bufonid species (gray) and the otic epidermis in *Atelopus* sp. 2 (red). $N=4$ for all species (see Fig. 2 for species names). (B) Difference in hearing of *Atelopus* sp. 2 individuals ($N=3$) before and after application of Vaseline to cover the body wall overlying the lungs. Negative values indicate a decrease in hearing sensitivity following Vaseline manipulation.

Table 4. Estimated least squares means differences between hearing sensitivity thresholds of individuals with and without Vaseline treatment

Frequency (Hz)	<i>Atelopus</i> sp. 2		Tympanic membrane (dB)			
	Body lung wall (dB)	Tympanic area (dB)	<i>R. alata</i>	<i>R. haematiticus</i>	<i>R. horribilis</i>	<i>R. spinulosa</i>
300	-5±4	-1±4	1±4	1±4	1±4	-6±4
500	7±4	-5±5	-1±4	-4±4	-6±4	-3±4
700	5±4	0±5	-9±4*	-4±4	-14±4**	0±4
900	3±4	-3±5	-9±4*	1±4	-18±4***	1±4
1100	4±5	-2±5	-9±4*	-4±4	-20±4***	-4±4
1300	-1±4	2±5	-11±4*	-14±4***	-25±4***	-9±4*
1500	-4±4	2±5	-7±4	-16±4***	-29±4***	-16±4***
2000	-3±4	1±5	-16±4***	-24±4***	-28±4***	-18±4***
2500	-2±4	-4±5	-15±4***	-26±4***	-24±4***	-16±4***
3000	2±4	2±5	-11±4*	-22±4***	-13±5*	-19±7**
3500	2±4	-3±5	-13±5*	-17±4***	NA	NA
4000	0±4	3±5	-10±7	-10±4*	-7±7	NA

Estimated least squares means (\pm s.e.) rounded to the nearest dB are given for hearing sensitivity differences at each frequency, with significant differences between Vaseline treatments in bold (* P <0.05, ** P <0.01, *** P <0.001). A negative estimated difference indicates that the Vaseline treatment increased the hearing threshold (made the anuran less sensitive) at that frequency while a positive estimated difference indicates that the Vaseline treatment decreased the hearing threshold (made the anuran more sensitive) at that frequency.

inner ear may be ineffective at sensing mid-range frequencies. Alternatively, as *Atelopus* species often breed near streams (Savage, 1972; Cocroft et al., 1990; Hödl and Amézquita, 2001), which create high levels of ambient noise, reaching maximum levels at lower frequencies (below 900 Hz; Hödl and Amézquita, 2001; Brumm and Slabbekoorn, 2005), stream noise may relax selection for hearing sensitivity at mid-range frequencies. In general, the hearing differences between *Atelopus* and the eared bufonids in this study were larger than those found by Jaslow and Lombard (1996), who found only a 5 dB hearing difference above 1000 Hz between the earless *A. chiriquiensis* and the eared *H. regilla*. Given Jaslow and Lombard's (1996) results rely on the hearing sensitivity of a single *Atelopus* species and single eared non-bufonid, the discrepancy between our study and theirs could largely be due to their limited sampling. Thus, although *Atelopus* are able to hear high frequencies better than other earless bufonids, we found they have low sensitivity at mid-range frequencies, which could be due to the lack of a tympanic middle ear or to ambient stream noise and relaxed selection.

Hearing in earless and partially eared *Atelopus* and its implications for middle ear evolution within *Atelopus*

Despite containing almost all the functionally relevant pieces of a tympanic middle ear, the partial middle ear found in some *Atelopus* species does not consistently benefit hearing sensitivity. Our manipulation experiment provides evidence that the partial middle ear of *Atelopus* does not function similarly to the tympanic middle ear of other anurans. Putting Vaseline over the otic epidermis did not appear to decrease hearing sensitivity in *Atelopus* sp. 2. It is therefore unlikely that this *Atelopus* with a partial middle ear relies on vibration of the tympanic region for transferring sound waves to the inner ear. Thus, *Atelopus* are capable of hearing via some other extratympanic pathway that is probably functioning in both earless and partially eared species.

The lack of difference in hearing sensitivity between our earless and partially eared *Atelopus* species conflicts with the 8–13 dB hearing advantage from 2000 to 2500 Hz found by Lindquist et al. (1998). Given that Lindquist et al. (1998) only compared one species with a partial middle ear (*A. flavescens*) with two earless species [*Atelopus* sp. (Nusagandi) and *A. lozanoi*] and we only compared one earless species with two species with a partial middle ear, conflict could be attributed to taxon sampling. Within our study,

even partially eared species differed in hearing, so selecting small numbers of species in each study could lead to ambiguities in estimating any hearing advantages of partial middle ears. Furthermore, the ring of cartilage that normally surrounds the tympanic membrane (the tympanic annulus) was incomplete in both partially eared species in this study, and an incomplete tympanic annulus is associated with non-functional tympanic middle ears in developing bufonids (Womack et al., 2016). Whether it is due to the incomplete tympanic annulus or the lack of tympanic membrane, the partial middle ear in *Atelopus* appears to convey little to no hearing sensitivity benefits; thus, its presence is perplexing and requires further research.

To interpret the species differences in hearing within *Atelopus*, we need more extensive sampling among numerous species in a phylogenetic context to infer evolutionary shifts in acoustic sensitivity, vibrational sensitivity and sound localization. Partial middle ears may enhance sensitivity to substrate-borne vibration, or the coupling of the middle ears via the middle ear cavities and Eustachian tubes may provide sound localization benefits. Although sound localization was not explored in this study, sensitivity to vibration was not enhanced in our partially eared *Atelopus* species (M.C.W., unpublished data). To relate those patterns to selection, we need natural history data that characterize communication strategies across the same set of species.

Extratympanic hearing mechanisms for *Atelopus*

Our manipulative experiments suggest that vibration of the body lung wall is not an important extratympanic pathway in *Atelopus*. This is surprising given the body lung wall has been shown to vibrate in response to frequencies around 2500 Hz in other *Atelopus* species (Lindquist et al., 1998), and covering the body lung wall of *Bombina orientalis* with silicon grease resulted in a 20–25 dB decrease of hearing sensitivity at all frequencies tested in that study (100–1000 Hz; Hetherington and Lindquist, 1999). Despite being one of the most discussed and experimentally investigated extratympanic hearing pathways in anurans (Narins et al., 1988; Ehret et al., 1990, 1994; Hetherington, 1992; Hetherington and Lindquist, 1999; Mason, 2006; Boistel et al., 2013), the lung pathway does not appear to contribute to high-frequency hearing sensitivity in *Atelopus* via vibration of the body lung wall.

Other extratympanic pathways could contribute to the high-frequency hearing of *Atelopus*. Boistel et al. (2013) proposed bone

conduction enhanced by resonance of the oral cavity to explain high-frequency communication in the earless *Sechelophryne gardineri*. However, the effectiveness of this pathway at high frequencies has not yet been tested experimentally. Thus, the mechanisms of high-frequency hearing in *Atelopus* and other anurans without a middle ear remain unverified.

Concluding remarks

Atelopus species are sensitive to high-frequency airborne sound despite lacking a tympanic middle ear. The mechanisms of *Atelopus* high-frequency hearing remain unclear, but vibration of the body lung wall probably does not contribute. Additionally, the partial middle ear found in some *Atelopus* species does not convey a strong hearing advantage, making it unlikely that direct selection pressures for increased hearing sensitivity are acting to retain or regain middle ear structures within *Atelopus*. Future research into extratympanic hearing mechanisms in *Atelopus* is needed to fully understand those mechanisms within anurans more generally and their influence on middle ear evolution.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: M.C.W., K.L.H., J.C.-D.; Methodology: M.C.W., K.L.H., J.C.-D.; Software: J.C.-D.; Validation: M.C.W., J.C.-D.; Formal analysis: M.C.W., K.L.H.; Investigation: M.C.W., K.L.H., J.C.-D., L.A.C.; Resources: M.C.W., K.L.H., L.A.C.; Data curation: M.C.W., L.A.C.; Writing - original draft: M.C.W.; Writing - review & editing: M.C.W., K.L.H., J.C.-D., L.A.C.; Visualization: M.C.W.; Supervision: K.L.H., J.C.-D.; Project administration: M.C.W., K.L.H.; Funding acquisition: M.C.W., K.L.H.

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Supplementary information

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References

- Bates, D., Maechler, M., Bolker, B. and Walker, S. (2015). Fitting linear mixed-effects models using lme4. *J. Stat. Softw.* **67**, 1-48.
- Boatright-Horowitz, S. S. and Simmons, A. M. (1995). Postmetamorphic changes in auditory sensitivity of the bullfrog midbrain. *J. Comp. Physiol. A Neuroethol. Sens. Neural Behav. Physiol.* **177**, 577-590.
- Boistel, R., Aubin, T., Cloetens, P., Langer, M., Gillet, B., Josset, P., Pollet, N. and Herrel, A. (2011). Whispering to the deaf: communication by a frog without external vocal sac or tympanum in noisy environments. *PLoS ONE* **6**, e22080.
- Boistel, R., Aubin, T., Cloetens, P., Peyrin, F., Scotti, T., Herzog, P., Gerlach, J., Pollet, N. and Aubry, J.-F. (2013). How minute sooglossid frogs hear without a middle ear. *Proc. Natl. Acad. Sci. USA* **110**, 15360-15364.
- Brumm, H. and Slabbekoorn, H. (2005). Acoustic communication in noise. *Adv. Study Behav.* **35**, 151-209.
- Christensen-Dalsgaard, J. and Carr, C. E. (2008). Evolution of a sensory novelty: tympanic ears and the associated neural processing. *Brain Res. Bull.* **75**, 365-370.
- Christensen, C. B., Christensen-Dalsgaard, J., Brandt, C. and Madsen, P. T. (2012). Hearing with an atympanic ear: good vibration and poor sound-pressure detection in the royal python, *Python regius*. *J. Exp. Biol.* **215**, 331-342.
- Cocroft, R. B., McDiarmid, R. W., Jaslow, A. P. and Ruiz-Carranza, P. M. (1990). Vocalizations of eight species of *Atelopus* (Anura: Bufonidae) with comments on communication in the genus. *Copeia* **1990**, 631-643.
- Ehret, G., Tautz, J., Schmitz, B. and Narins, P. M. (1990). Hearing through the lungs: lung-eardrum transmission of sound in the frog *Eleutherodactylus coqui*. *Naturwissenschaften* **77**, 192-194.
- Ehret, G., Keilwerth, E. L. and Kamada, T. S. (1994). The lung-eardrum pathway in three treefrog and four dendrobatid frog species: some properties of sound transmission. *J. Exp. Biol.* **195**, 329-343.
- Hartline, P. H. (1971). Physiological basis for detection of sound and vibration in snakes. *J. Exp. Biol.* **54**, 349-371.
- Hetherington, T. E. (1992). The effects of body size on functional properties of middle ear systems of anuran amphibians. *Brain Behav. Evol.* **39**, 133-142.
- Hetherington, T. E. and Lindquist, E. D. (1999). Lung-based hearing in an "earless" anuran amphibian. *J. Comp. Physiol. A* **184**, 395-401.
- Hödl, W. and Amézquita, A. (2001). Visual signaling in anuran amphibians. *Anuran Commun.* **18**, 121-141.
- Jaslow, A. P. and Lombard, R. E. (1996). Hearing in the neotropical frog, *Atelopus chiriquiensis*. *Copeia* **1996**, 428-432.
- Kremer, J. R., Mastrorarde, D. N. and McIntosh, J. R. (1996). Computer visualization of three-dimensional image data using IMOD. *J. Struct. Biol.* **116**, 71-76.
- Lenth, R. V. (2016). Least-squares means: the R package lsmeans. *J. Stat. Softw.* **69**, 1-33.
- Lindquist, E. D., Hetherington, T. E. and Volman, S. F. (1998). Biomechanical and neurophysiological studies on audition in eared and earless harlequin frogs (*Atelopus*). *J. Comp. Physiol. A* **183**, 265-271.
- Lombard, R. E. and Straughan, I. R. (1974). Functional aspects of anuran middle ear structures. *J. Exp. Biol.* **61**, 71-93.
- Manley, G. A. (2010). An evolutionary perspective on middle ears. *Hear. Res.* **263**, 3-8.
- Manley, G. A. and Sienknecht, U. J. (2013). The evolution and development of middle ears in land vertebrates. In *The Middle Ear* (ed. Puria, S., Fay, R. R. and Popper, A. N.), pp. 7-30. New York: Springer.
- Mason, M. J. (2006). Pathways for sound transmission to the inner ear in amphibians. In *Hearing and Sound Communication in Amphibians* (ed. P. Narins, A. S. Feng and R. R. Fay), pp. 147-183. New York: Springer.
- Mason, M. J. and Narins, P. M. (2002). Seismic sensitivity in the desert golden mole (*Eremitalpa granti*): a review. *J. Comp. Psychol.* **116**, 158.
- Miranda, J. A. and Wilczynski, W. (2009). Sex differences and androgen influences on midbrain auditory thresholds in the green treefrog, *Hyla cinerea*. *Hear. Res.* **252**, 79-88.
- Narins, P. M., Ehret, G. and Tautz, J. (1988). Accessory pathway for sound transfer in a neotropical frog. *Proc. Natl. Acad. Sci. USA* **85**, 1508-1512.
- Pereyra, M. O., Womack, M. C., Barrionuevo, J. S., Blotto, B. L., Baldo, D., Targino, M., Ospina-Sarria, J. J., Guayasamin, J. M., Coloma, L. A., Hoke, K. L. et al. (2016). The complex evolutionary history of the tympanic middle ear in frogs and toads (Anura). *Sci. Rep.* **6**, 34130.
- Savage, J. M. (1972). The harlequin frogs, genus *Atelopus*, of Costa Rica and western Panama. *Herpetologica* **28**, 77-94.
- Shen, J. X., Xu, Z. M., Yu, Z. L., Wang, S., Zheng, D. Z. and Fan, S. C. (2011). Ultrasonic frogs show extraordinary sex differences in auditory frequency sensitivity. *Nat. Commun.* **2**, 342.
- Sueur, J., Aubin, T. and Simonis, C. (2008). Seewave: a free modular tool for sound analysis and synthesis. *Bioacoustics* **18**, 213-226.
- Wever, E. G. (1975). The caecilian ear. *J. Exp. Zool. A Ecol. Genet. Physiol.* **191**, 63-71.
- Womack, M. C., Christensen-Dalsgaard, J. and Hoke, K. L. (2016). Better late than never: effective air-borne hearing of toads delayed by late maturation of the tympanic middle ear structures. *J. Exp. Biol.* **219**, 3246-3252.
- Womack, M. C., Christensen-Dalsgaard, J., Coloma, L. A., Chapparo, J. C. and Hoke, K. L. (2017). Earless toads sense low frequencies but miss the high notes. *Proc. R. Soc. B* **284**, 20171670.

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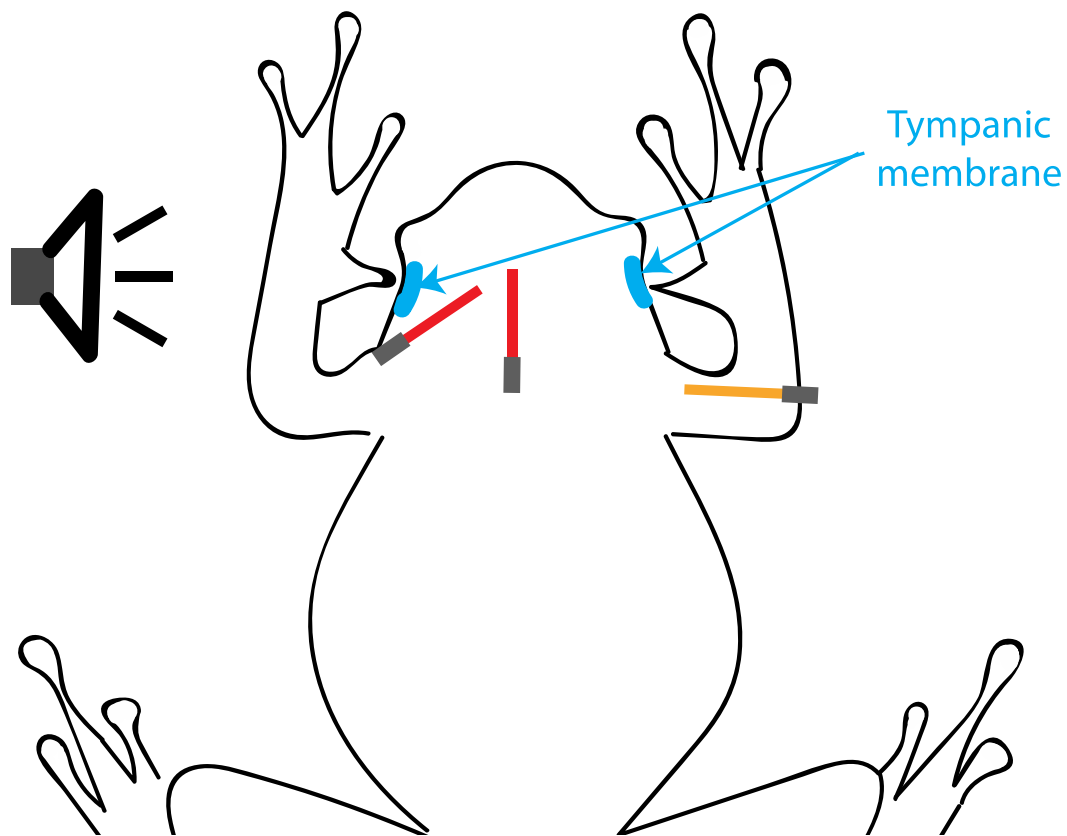
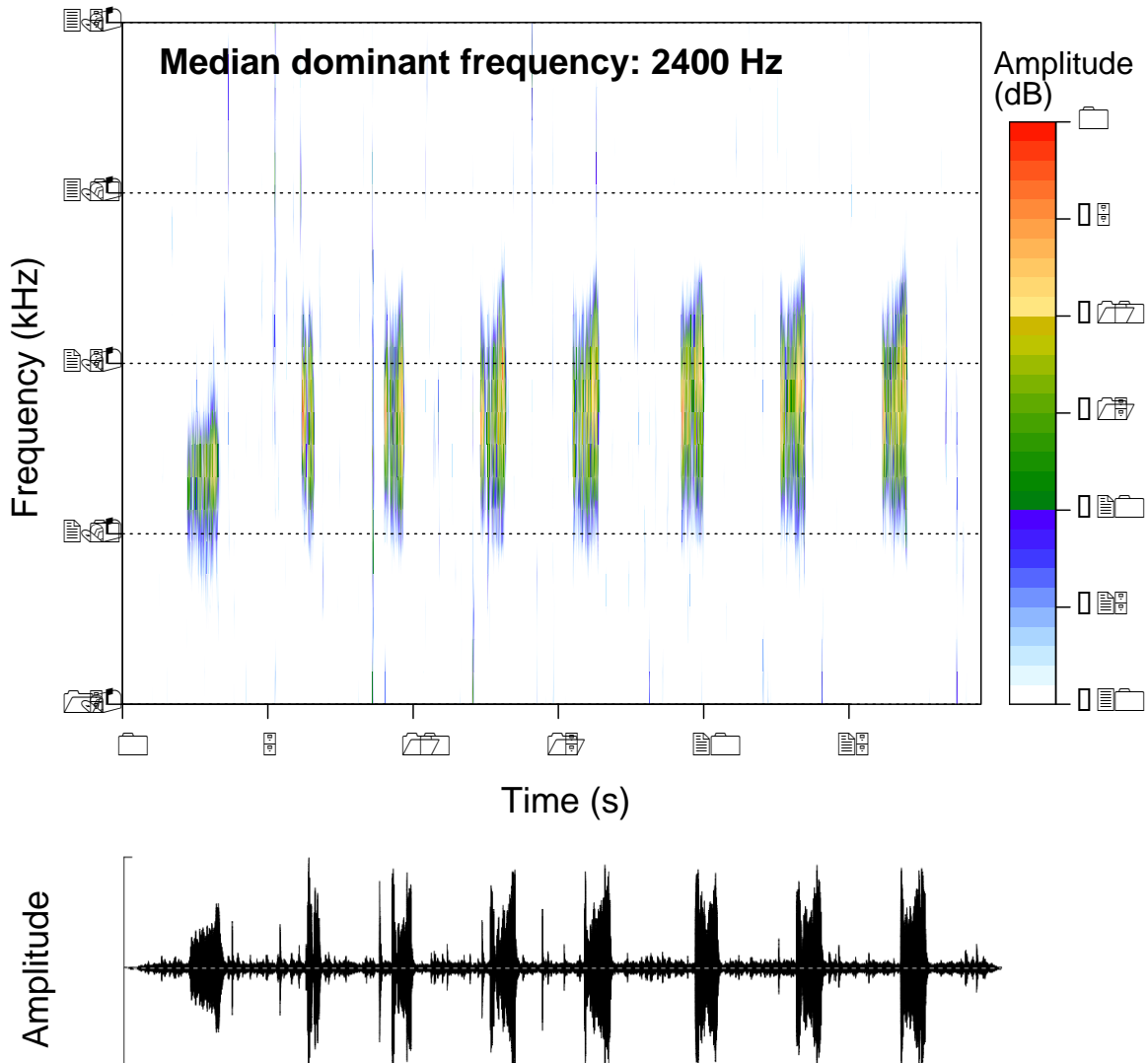


Fig. S1. A diagram of the electrode (red) placement for auditory brainstem recordings (ABRs). We subdermally placed differential electrodes (red) over the midbrain and VIIIth (auditory) nerve and placed a third ground electrode (orange) within the arm contralateral to the VIIIth nerve being measured. Tympanic membranes are shown in blue.

A



B

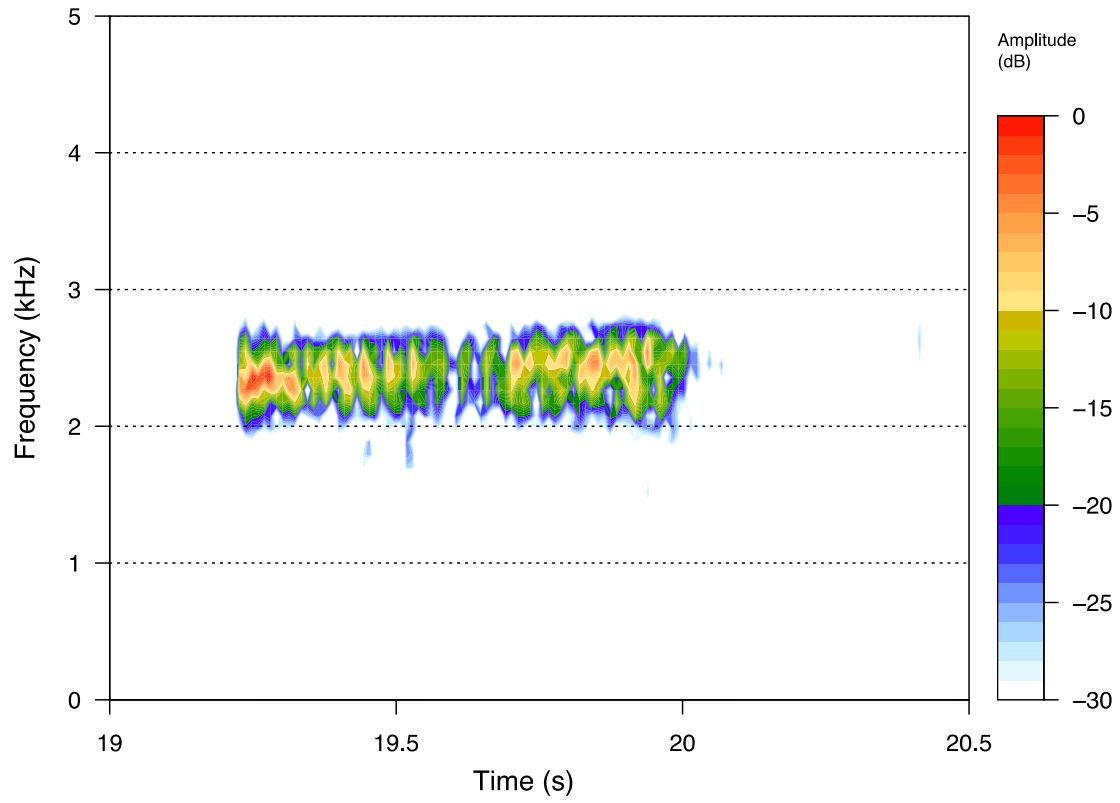


Fig. S2. *Atelopus* sp. 2 (*spumarius* complex) call collection & analysis.

The call of *Atelopus* sp. 2 was analyzed for this study. Elicio E. Tapia recorded the call on July 6th 2016 at 9:00 H and 10:00 H with a digital recorder (Olympus Linear PCM Recorder LS 10S) and microphone (Sennheisser). These recording was done (under lab conditions) of a male born in captivity from parents from San Carlos de Limón, Morona Santiago Province, Ecuador. The frog was inside a terrarium 40x40 cm partially opened in the upper cover with other males in the terrarium. The microphone was at about 5 cm from the frog. Atmospheric pressure was about 730.78 mb, the altitude of Centro Jambatu lab is 2700 m asl, and the frog's temperature was about 20 °C.

The call was analyzed using the package seewave (Sueur et al., 2008) in R (R Core Team, 2015). The call was visualized with spectrograms and then trimmed to the time frame of the call. We eliminated background noise at frequencies above and below the call using a single band pass frequency filter (window length = 1024, window = "hanning", overlap = 75). From these trimmed call files, we calculated the median dominant frequency by performing an instantaneous frequency extraction by zero crossing.

(A) A spectrogram (top) and oscillogram (bottom) of the advertisement call of *Atelopus* sp. 2 with median dominant frequency given. For spectrogram - window length = 512, window = "hanning", overlap = 0.

(B) A spectrogram of one note within the advertisement call of *Atelopus* sp. 2 shown in A. For spectrogram - window length = 512, window = "hanning", overlap = 0.