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Supporting Information for

Closed-Loop Network of Skin-interfaced Wireless Devices for Quantifying Vocal Fatigue and Providing User Feedback

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19 **This PDF file includes:**

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21 Supporting text
22 Figures S1 to S17
23 Legends for Movies S1
24 Legends for Audio S1
25 SI Reference

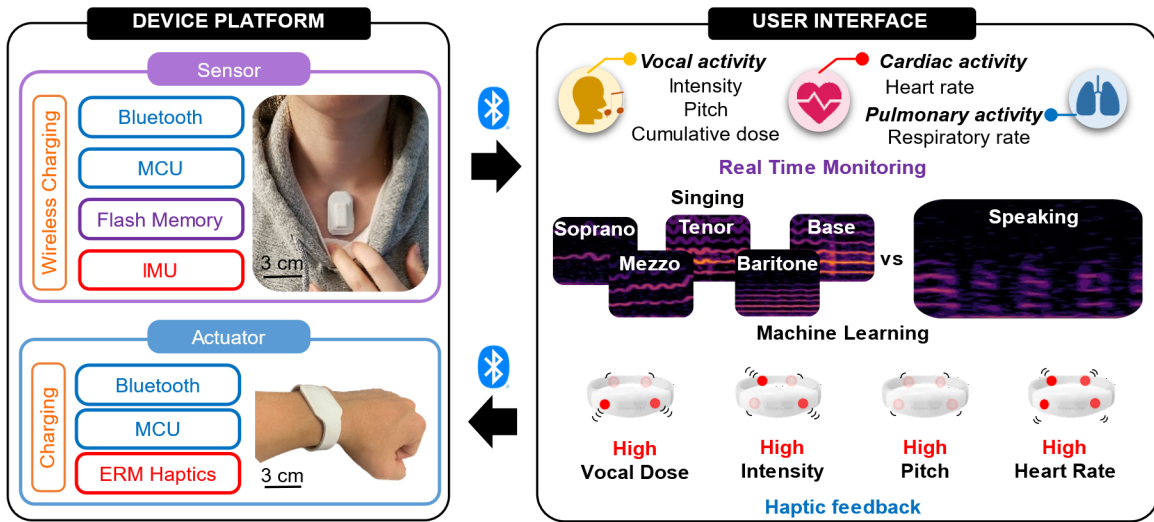
26 **Supporting Information Text**

27 **Quantifying the performance of the adhesives.** A Mark-10 Force Gauge enabled comparisons
28 of peel forces of four different adhesives on the skin. Similar measurements determined the forces
29 for removing the sensors from the magnetic adhesive when attached to the skin. *SI Appendix*, Fig.
30 S5 shows an image of the four adhesives tested: hydrogel, magnetic adhesive, fabric-based tape
31 with MED6500 adhesive, and the medical silicone adhesive. Peel forces are shown in *SI Appendix*,
32 Fig. S6. Removing the device from the magnetic adhesive and the hydrogel required the lowest
33 maximum peel force. The sharp steps in the force graph for removing the device from the magnetic
34 adhesives resulted from the device detaching sequentially from each magnet. The magnetic
35 adhesive with fabric-based tape for the skin interface demonstrated a moderate maximum peel
36 force and slope. The medical silicone adhesive (3M, 2477P) and the fabric-based tape coupled
37 with the MED6500SI adhesive had the highest maximum peel forces and greatest initial slopes. *SI*
38 *Appendix*, Fig. S7 shows images of a participant's skin at the suprasternal notch area to compare
39 redness at three time points (twenty minutes, forty minutes, and sixty minutes) after adhesive
40 removal. The images demonstrate decreasing redness of the skin from moderate at twenty minutes
41 to none at sixty minutes. The results of adhesive validation testing indicate that most users found
42 the adhesives easy to use and comfortable. Based on these results, the adhesive comprised of
43 fabric-based tape coupled with a MED6500 adhesive was chosen as the adhesive for continuous
44 monitoring for at least a day at a time.

45
46 **Verifying calculated vocal dose and perceived effort.** In studies to verify that the vocal dose
47 algorithm aligned with perceived effort, four singers (soprano, alto, tenor, and baritone) recorded a
48 set of known tasks. The tasks included normal speaking, speaking over 60 dB SPL of ambient
49 noise, whispered speaking, moderate singing (low to mid-range), moderately loud singing (mid to
50 high range), loud singing (low to mid-range), very loud singing (mid to high range), singing without
51 vibrato (low to mid-range), staccato arpeggios throughout the vocal range, and strained speaking.
52 Participants rated their perceived vocal effort on a scale from one to ten (1 = no effort, 10 =
53 maximum effort). Each task was recorded for one minute with one minute of rest in between.

54
55 **Aligning sensor data with self-reported activities.** Sixteen university singers, seven male and
56 eight female wore the MA device affixed just below the suprasternal notch and recorded all of their
57 daily vocal use. The singers wore the devices for between one day and five consecutive days,
58 removing them only to shower and sleep. Each simultaneously recorded their activities and self-
59 assessments on various surveys throughout each day. Each participant began and ended the day
60 with vocal fold "Swelling Check" exercises as a barometer of vocal function (1). The exercises
61 expose vocal fold mucosal swelling through singing quietly in the topmost part of their vocal range.
62 The highest pitch that the participant could no longer phonate clearly and softly defined the "ceiling
63 pitch". Participants reported the number of hours they slept along with their perceived quality of
64 sleep, and their morning ceiling pitch in a survey each day. Their morning swelling checks were
65 additionally recorded in a voice memo as a WAV file and uploaded to a cloud drive. Throughout
66 the day, participants completed multiple brief surveys, documenting the start and end time of each
67 unique vocal activity, as well as their perceived vocal effort for said activity. These activities
68 included speaking, solo singing, choral singing, quiet time, teaching, opera rehearsal and an option
69 to specify a type of activity that was not already listed. Every evening, at the completion of the
70 day's activities, participants performed both swelling checks and recorded the results in a voice
71 memo, reported their ceiling pitch in a survey, and then removed the device. The data was then
72 downloaded from the device to the smartphone.

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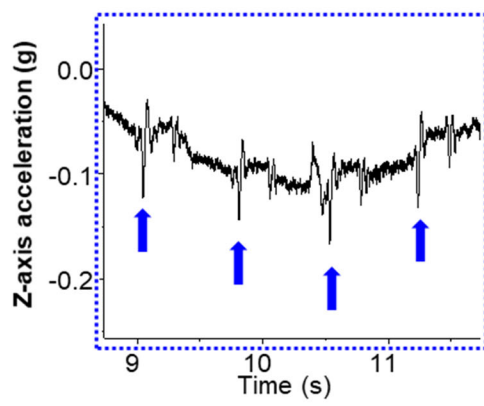
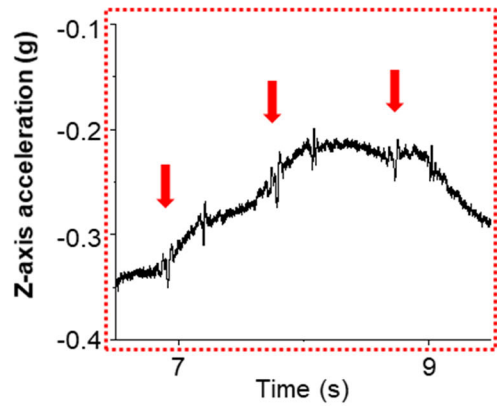
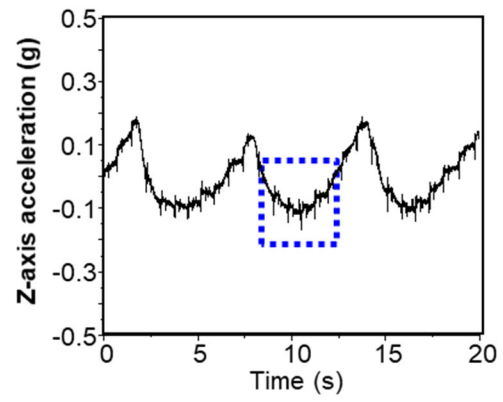
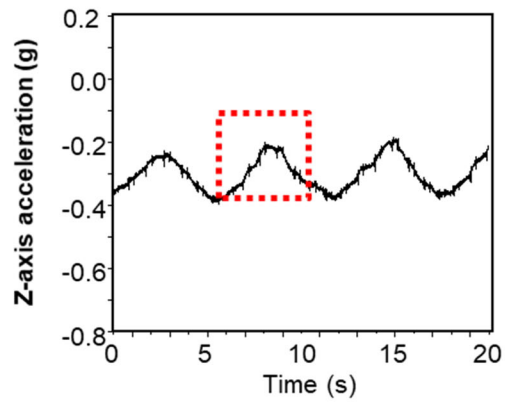
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Fig. S1. An illustration of the overall platform operation including sensor, user interface, and feedback actuator.



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79 **Fig. S2.** Devices packaged in various colors to promote user identification of devices on the mobile
80 application.



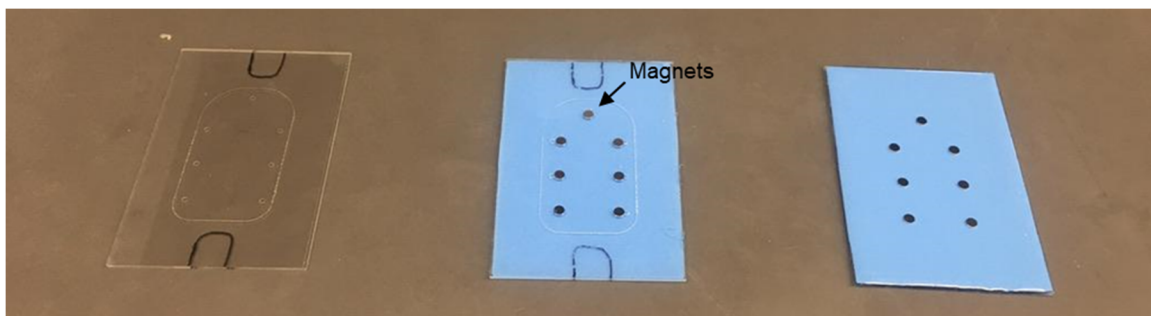
Double-sided medical silicone adhesive

Magnetic adhesive

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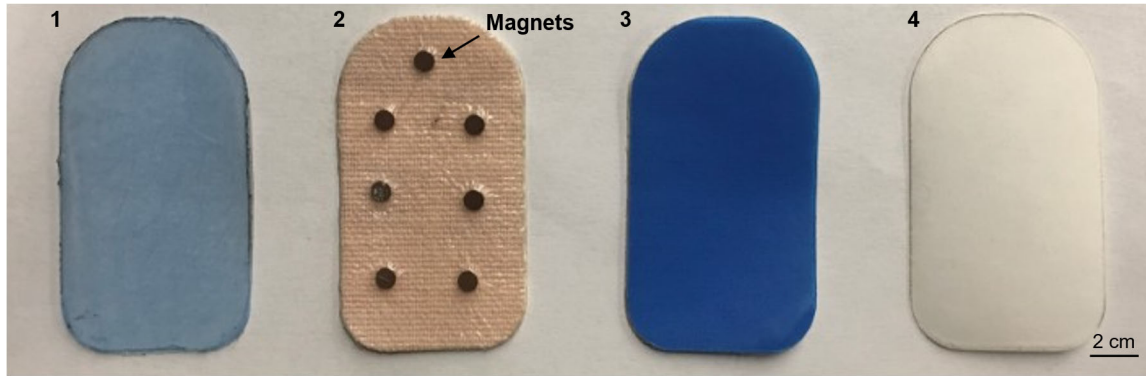
82 **Fig. S3.** Data collected using different adhesives during trials with subjects at rest. The bottom
 83 graphs are magnified sections of the top graphs to show that seismocardiogram signals are
 84 detectable using both adhesives.

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87 **Fig. S4.** Fabrication of the bottom encapsulation layer of the device with magnetic adhesives.



Hydrogel

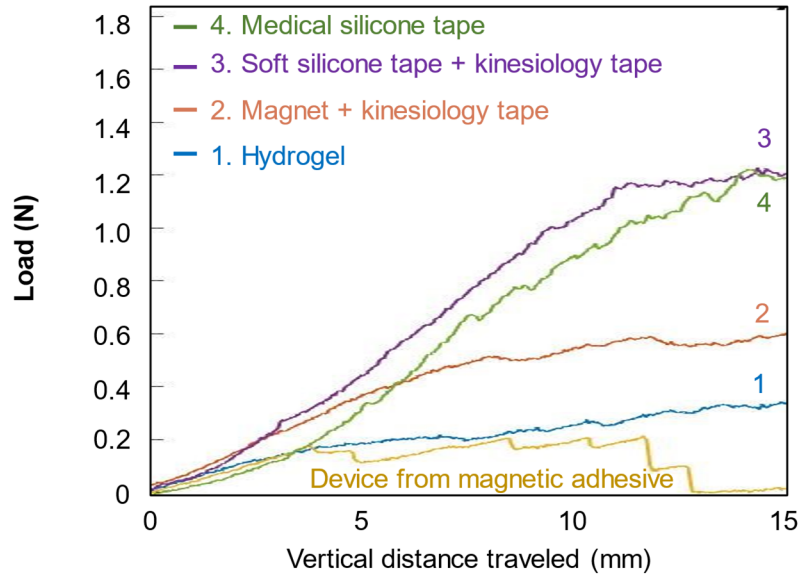
Kinesiology tape
+ Magnets

Kinesiology tape
+ soft silicone tape

Double sided medical
silicone tape

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89 **Fig. S5.** Various adhesives tested for skin comfort and redness after removal. The leftmost
90 adhesive is a hydrogel tape (KM 40A, KATECHO) alone. The second from the left is a combination
91 of kinesiology tape (SpiderTech, Nitto Denko) and magnets sealed with Tegaderm (3M). The
92 second from the right is a combination of kinesiology tape and a soft silicone adhesive (MED6500SI,
93 Avery Dennison). The rightmost adhesive is the medical silicone tape (2477P, 3M).



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95 **Fig. S6.** Peel forces for four adhesives on the skin (labeled 1 to 4) and for removal of a device from
96 the magnetic adhesive.



20 min

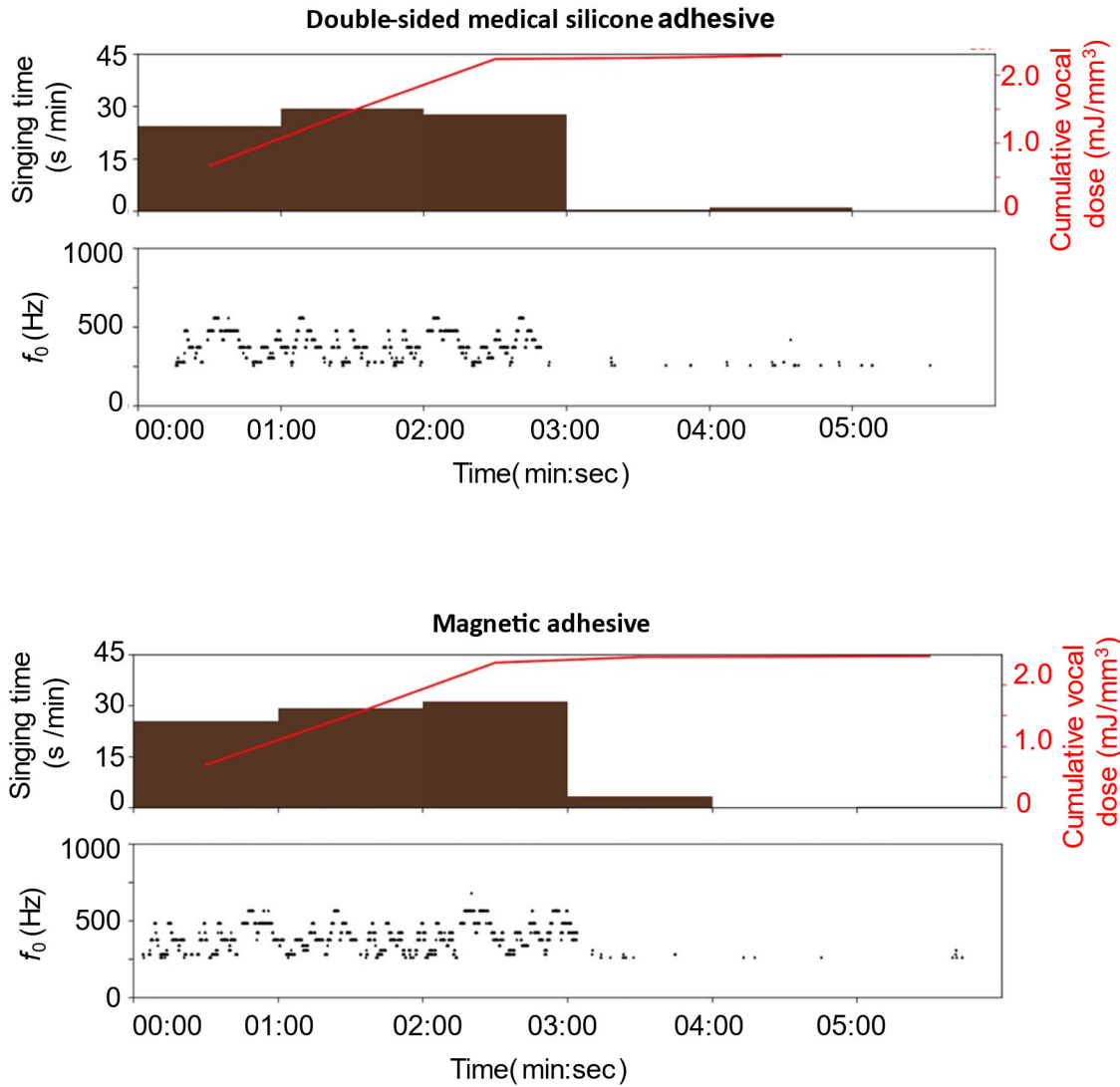
40 min

60 min

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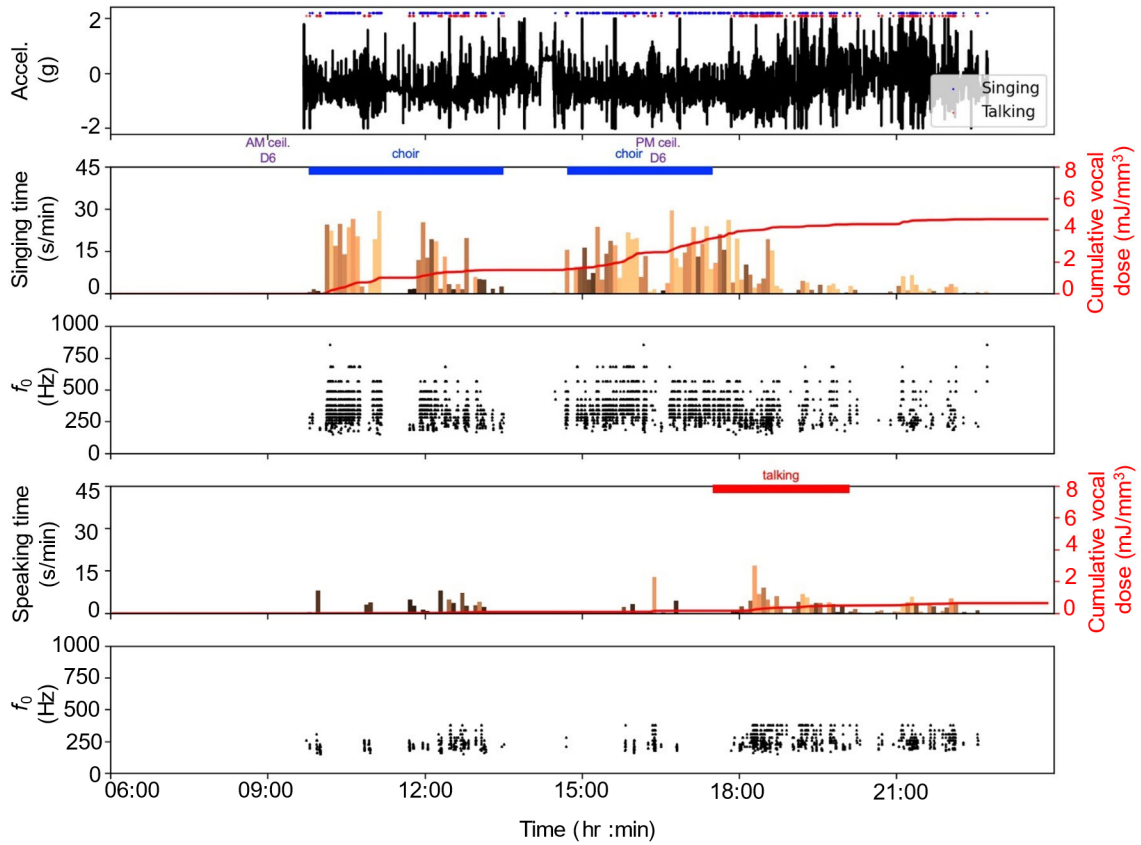
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Fig. S7. Images of skin redness at three time points after adhesive removal.



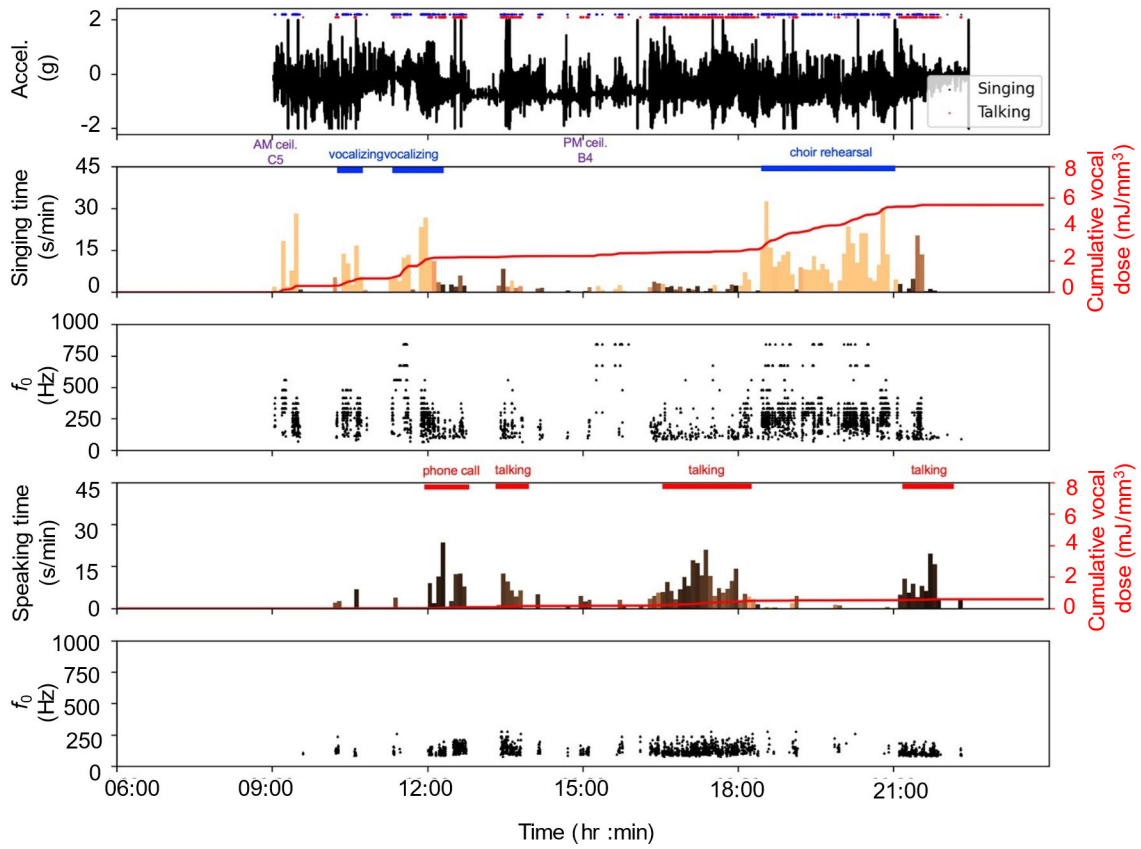
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100 **Fig. S8.** Comparison of data collected from devices coupled to the body using a magnetic adhesive
 101 and a non-magnetic adhesive. Data from the two devices yielded similar values for singing time,
 102 cumulative dose, and fundamental frequency during three minutes of speaking and three minutes
 103 of singing.



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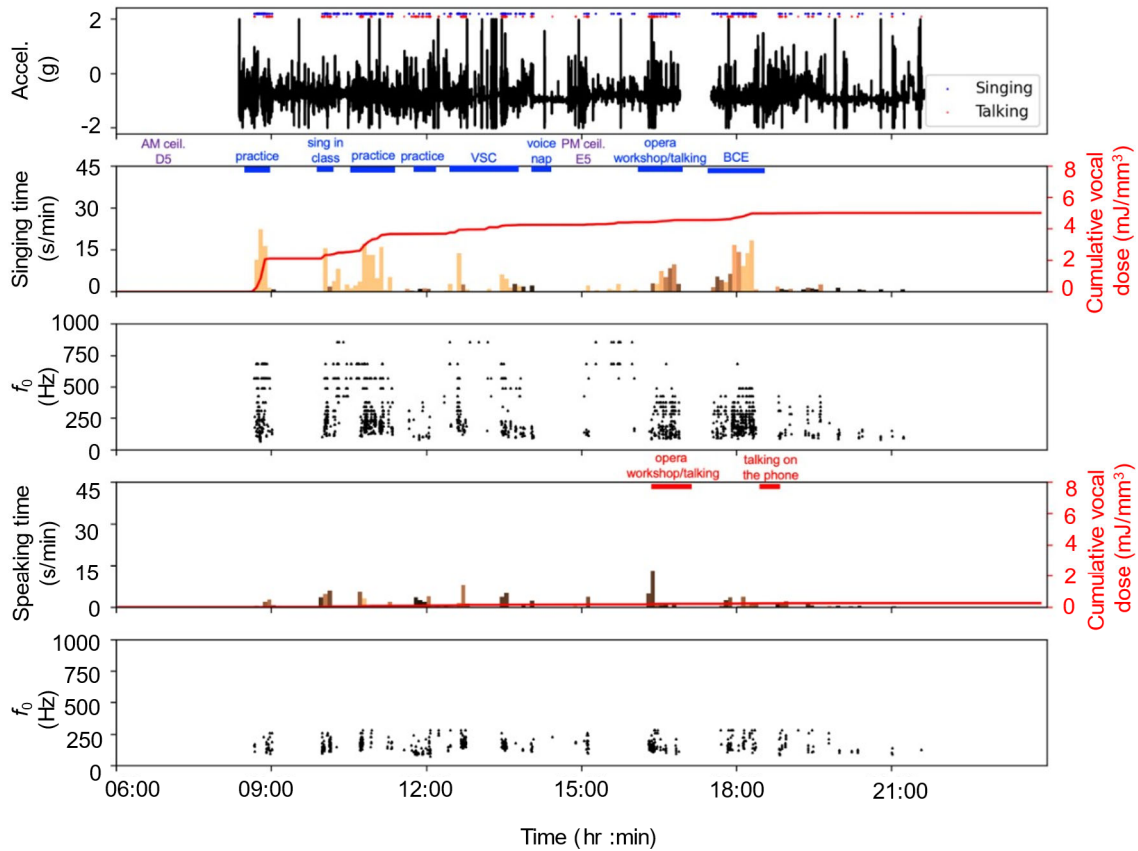
105 **Fig. S9.** Data from a mezzo-soprano with self-reported activities noted.



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Fig. S10. Data from a tenor with self-reported activities noted.

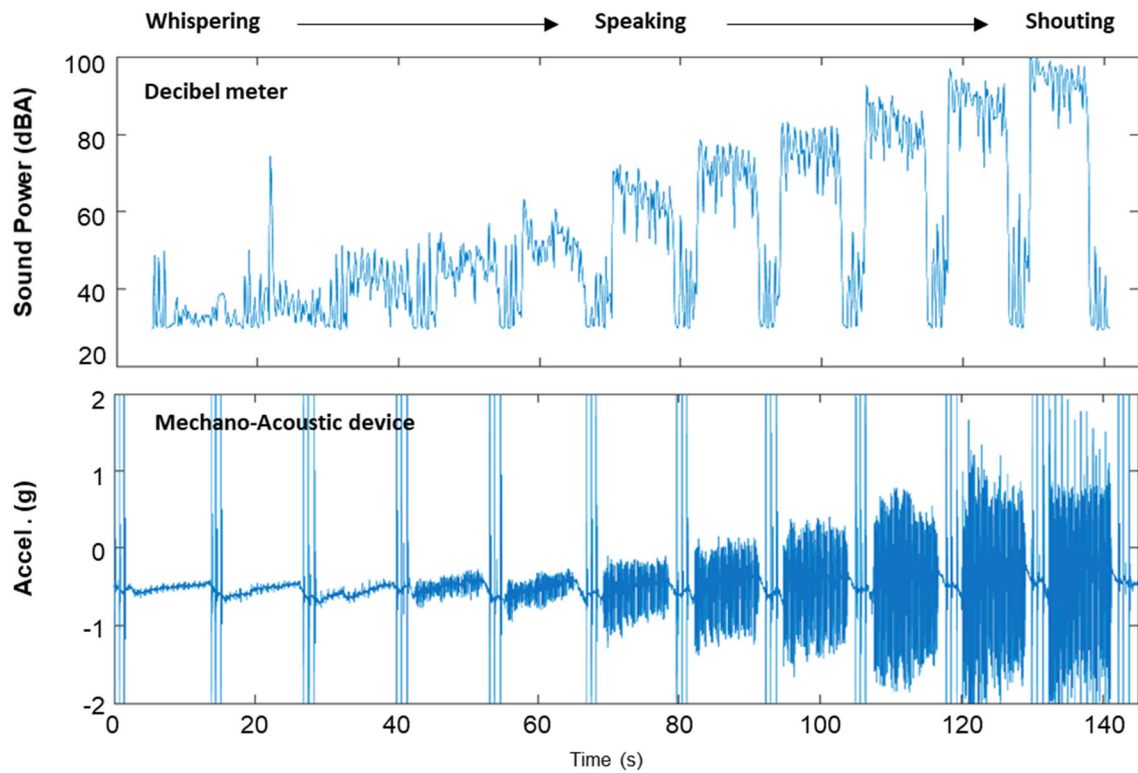


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Fig. S11. Data from a baritone with self-reported activities noted.

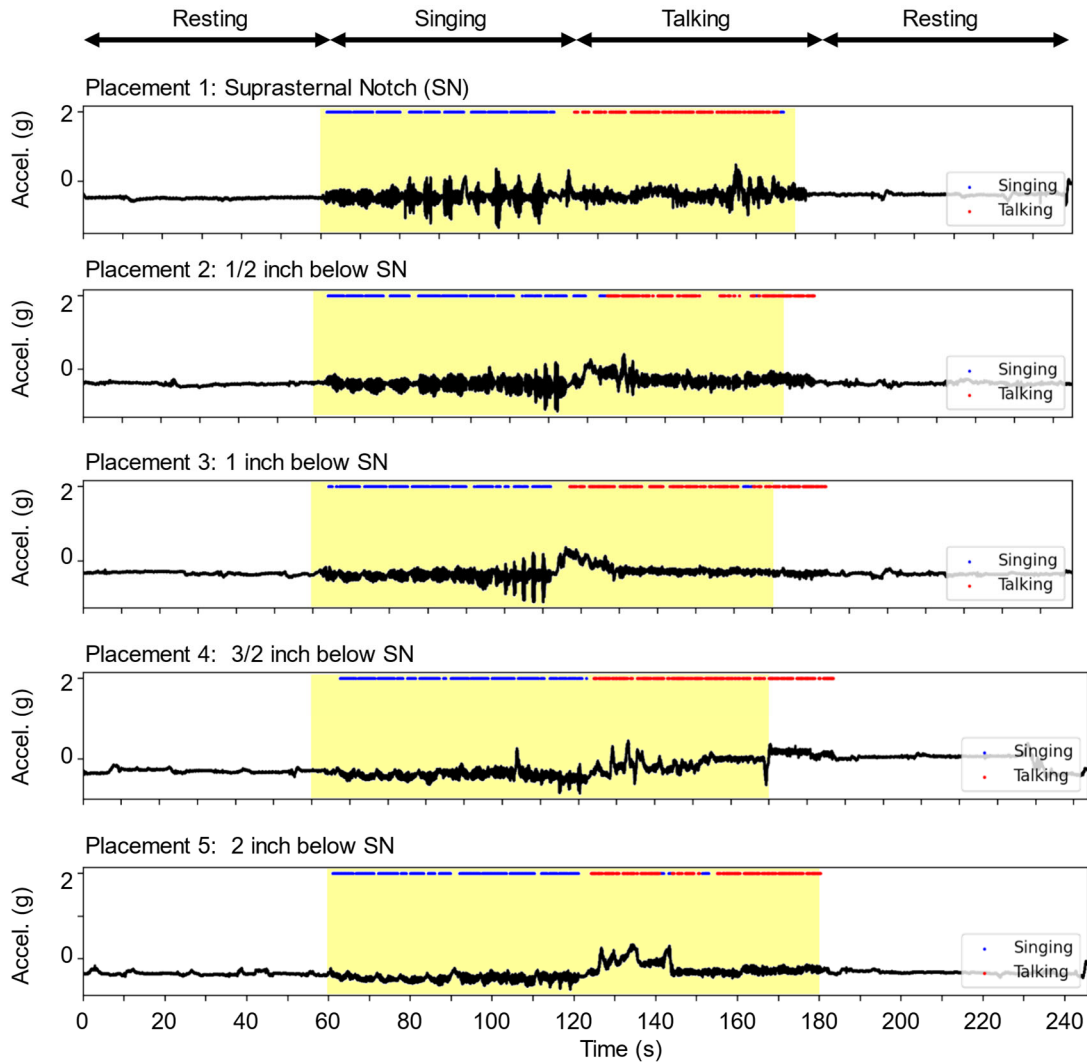
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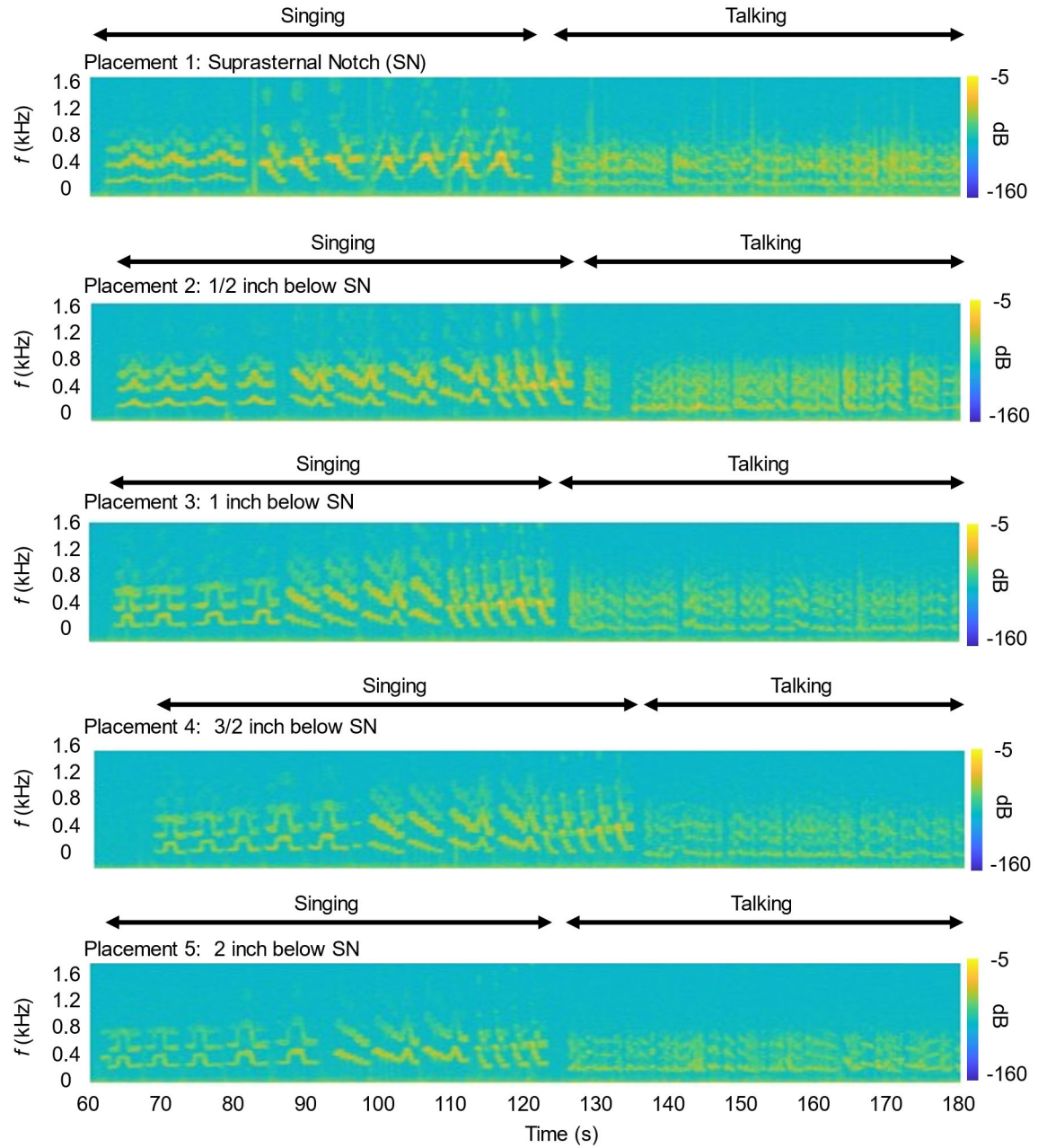
112 **Fig. S12.** Raw data for calibration of mechano-acoustic power and acoustic power. Note that the
 113 tapping signals between transitions causes saturated acceleration values.

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116 **Fig. S13.** Comparison of singing and speaking classification by the CNN model from the data
 117 collected with five different placements: suprasternal notch (SN), 0.5 inches below SN, 1 inch below
 118 SN, 1.5 inches below SN, and 2 inches below SN. Blue and red dot denotes classified singing and
 119 talking events.

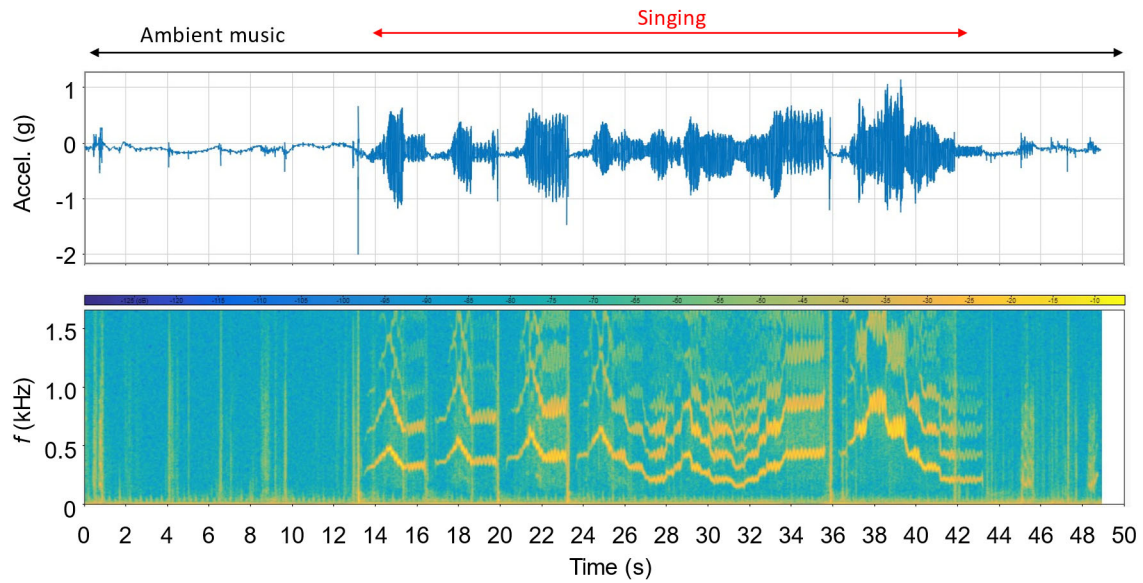


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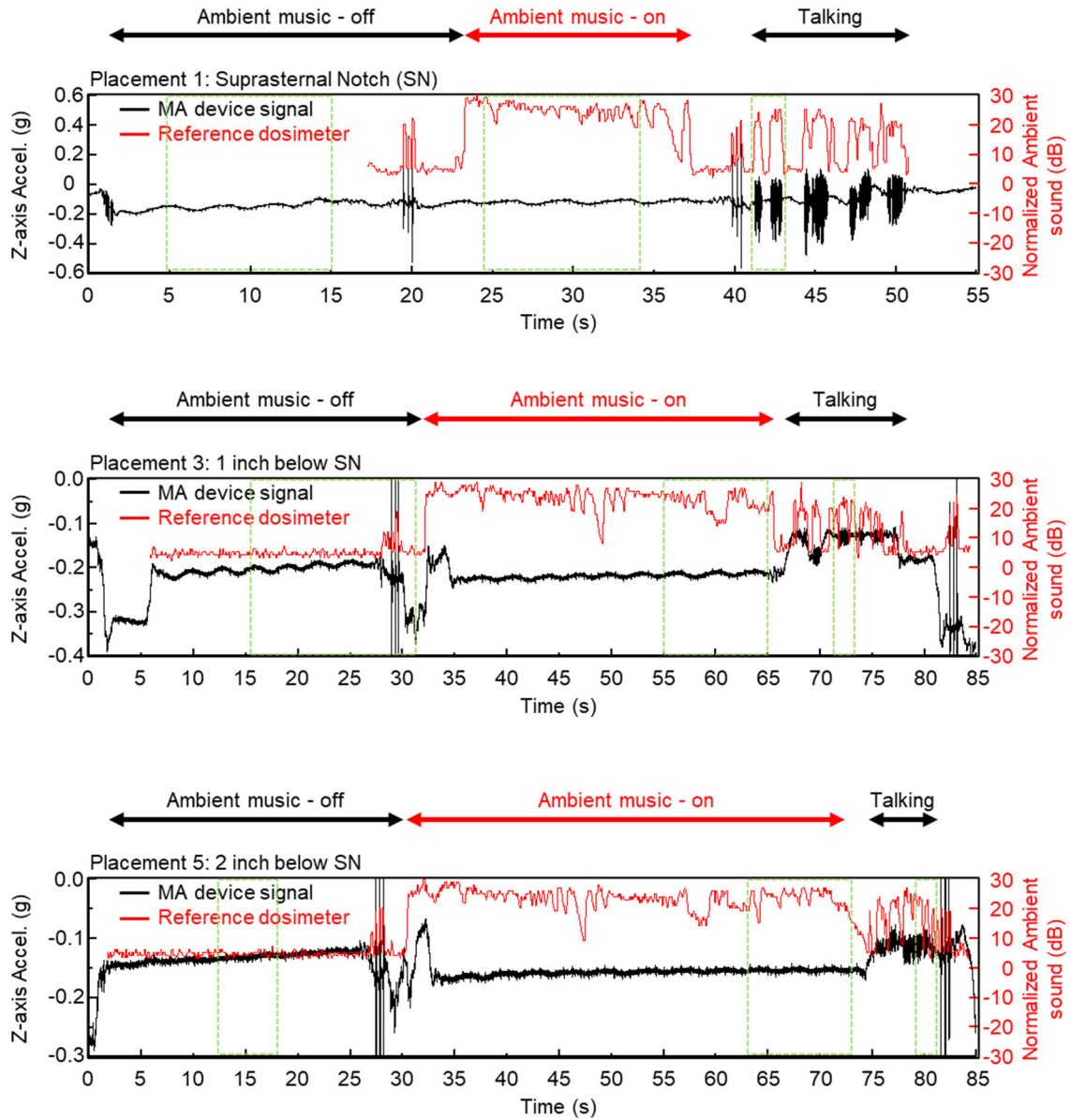
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Fig. S14. Comparison of spectrograms for singing and speaking data highlighted in the data from Fig. S13



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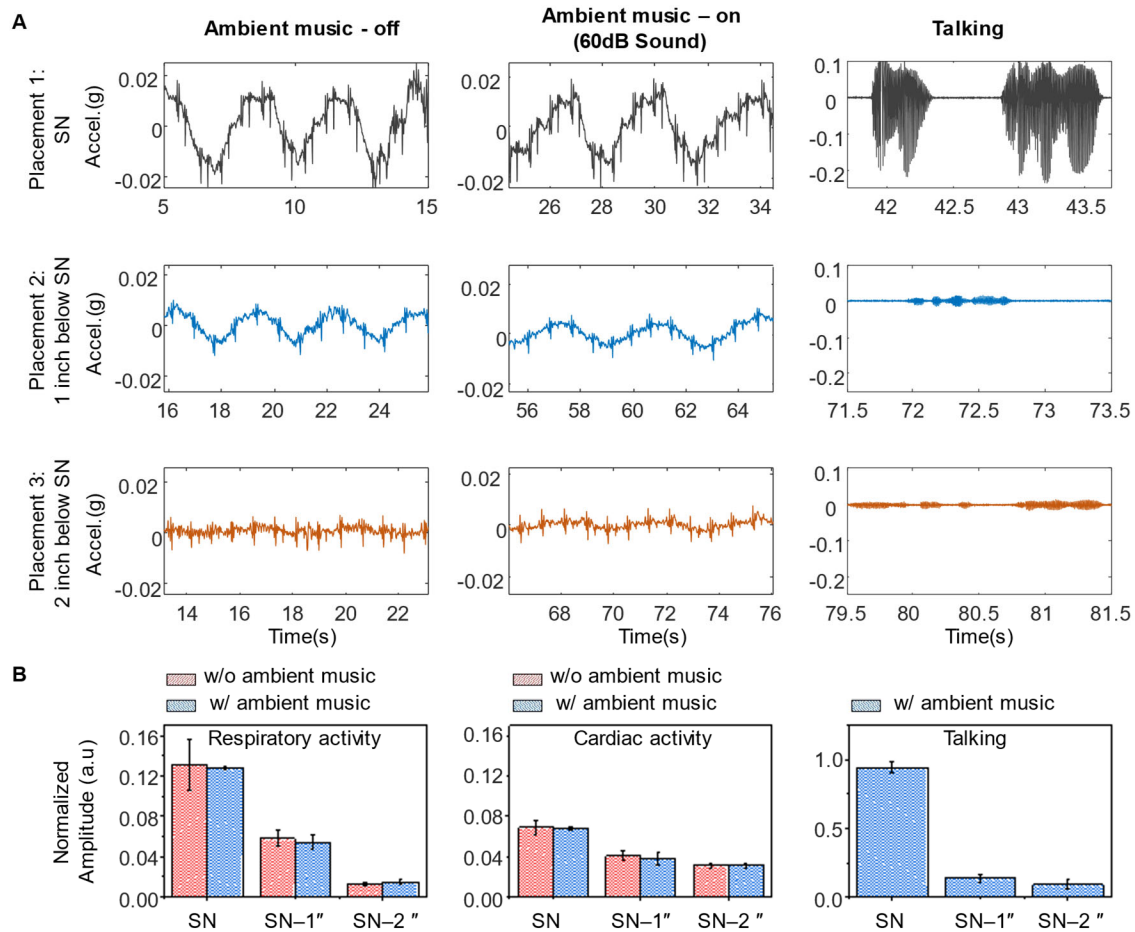
124 **Fig. S15.** Validation of interference by ambient noise. The subject wore an MA device on the upper
 125 chest. Data were collected with ambient loud music (Beethoven Symphony No. 9, 62 - 79 dBA
 126 range) while singing with different ranges of pitch.



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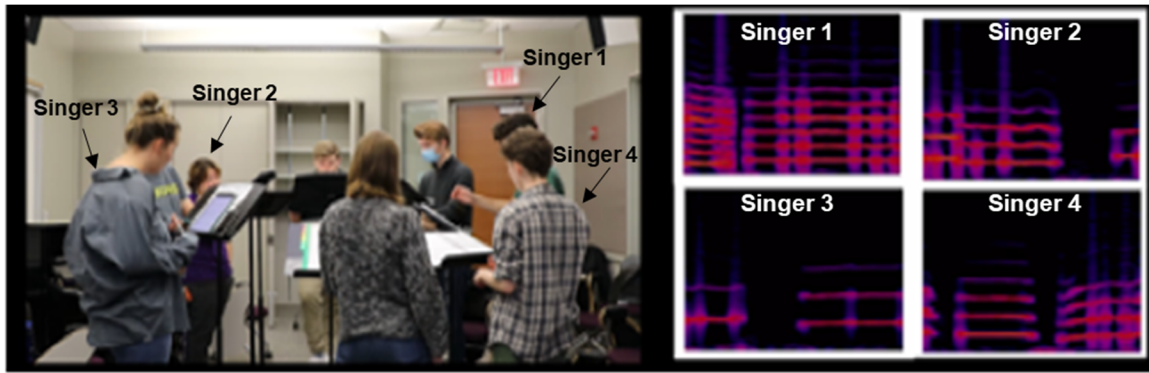
128 **Fig. S16.** Comparison of data collected in a noisy environment at three different locations including

129 suprasternal notch (SN), 1 inch below SN, and 2 inches below SN.



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131 **Fig. S17.** Quantitative comparison of signal-to-noise ratio (SNR) collected in a noisy environment
 132 at three different locations including suprasternal notch (SN), 1 inch below SN, and 2 inches below
 133 SN. (A) Zoom-in signals from green region in Fig. S16. (B) Normalized signal amplitude from three
 134 different placements in terms of respiratory activity, cardiac activity, and talking signal after data
 135 processing.



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Movie 1. Devices used in a choir rehearsal setting demonstrate the capacity to capture data from an individual singer without influence from vocalization by other singers. The studies involve four singers (one soprano, one alto, one tenor, and one bass) during a rehearsal, with devices paired with smartphones or tablets for real-time dosimetry calculations.



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Audio 1. Validation of interference by ambient noise with converted audio. Converting the MA data (Fig. S15) to audio files confirmed that the influence of ambient music is almost negligible.

146 **SI Reference**

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- 148 1. R. W. Bastian, A. Keidar, K. Verdolini-Marston, Simple vocal tasks for detecting vocal fold
149 swelling. *J. Voice* **4**, 172–183 (1990).