

# Masculine voices signal men's threat potential in forager and industrial societies

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Humans and many non-human primates exhibit large sexual dimorphisms in vocalizations and vocal anatomy. In humans, same-sex competitors and potential mates attend to acoustic features of male vocalizations, but vocal masculinity especially increases perceptions of physical prowess. Yet, the information content of male vocalizations remains obscure. We therefore examined relationships between sexually dimorphic acoustic properties and men's threat potential. We first introduce a new measure of the structure of vocal formant frequencies, 'formant position' ( $P_f$ ), which we show is more sexually dimorphic and more strongly related to height than is the most widely used measure of formant structure, 'formant dispersion', in both a US sample and a sample of Hadza foragers from Tanzania. We also show large sexual dimorphisms in the mean fundamental frequency ( $F_0$ ) and the within-utterance standard deviation in  $F_0$  ( $F_0 - \text{s.d.}$ ) in both samples. We then explore relationships between these acoustic parameters and men's body size, strength, testosterone and physical aggressiveness. Each acoustic parameter was related to at least one measure of male threat potential. The most dimorphic parameters,  $F_0$  and  $P_f$ , were most strongly related to body size in both samples. In the US sample,  $F_0$  predicted testosterone levels,  $P_f$  predicted upper body strength and  $F_0 - \text{s.d.}$  predicted physical aggressiveness.

**Keywords:** formant frequency; fundamental frequency; sexual selection

## 1. INTRODUCTION

The acoustic properties of adult vocalizations are sexually dimorphic in many primates [1–3], including humans [4]. Males vocalize at a lower fundamental frequency ( $F_0$ ), the rate of vocal fold vibration and acoustic parameter closest to pitch, and with lower, more closely spaced formant frequencies [2,4], frequencies of high energy that affect vocal timbre. In humans,  $F_0$  may also vary less across an utterance in men than in women, resulting in a more monotone voice [5].

Except for  $F_0$  variation, the anatomical and physiological substrates for these vocal sex differences are well understood. At puberty, elevated testosterone levels [6] acting through androgen receptors in the vocal folds [7] cause the vocal folds to grow longer and thicker in boys than in girls [8]. Men's larger vocal folds consequently vibrate at approximately half the  $F_0$  of women's during phonation. Similarly, boys' larynges descend at puberty [9], producing a longer vocal tract and lower, more closely spaced formant frequencies [10].

## 2. SEXUAL SELECTION AND HUMAN VOICES

Recent research has focused on the evolutionary reasons for the pubertal development of vocal sex differences in humans [11] and non-human primates [12]. Of particular interest has been sexual selection, which operates through both mate choice, favouring ornaments and sexual displays

for attracting mates, and contest competition, favouring size, strength, aggression, weapons and threat displays [13].

Male mate choice of females may have contributed to higher fundamental and formant frequencies in women than in men [14–16]. However, sexual selection was probably stronger in shaping men's traits [17], and sexually selected traits tend to emerge at puberty [13], when precipitous changes in male, but not female, vocal anatomy occur. Moreover, low speaking  $F_0$  and a monotone voice predicted men's mating success in a US sample [18,19], and low speaking  $F_0$  predicted men's reproductive success among Hadza foragers of Tanzania [20], suggesting selection on men's voices.

Although some evidence implicates female mate choice in favouring deeper, more monotone voices in men [18,21–23], theoretical predictions and phylogenetic and functional data suggest that male contests played a larger role than female choice in shaping men's phenotypes [24]. Masculine fundamental and formant frequencies in particular increase perceptions of dominance and fighting ability among men more effectively than they increase attractiveness to women [11,25–27]. Likewise, a monotone voice appears more strongly related to perceptions of dominance than attractiveness [23]. Moreover, modulation of  $F_0$  affects and reflects dominance and submissiveness across animal species [28], including humans [11].

## 3. INFORMATION CONTENT OF MALE VOCAL SIGNALS

Although masculine voices evoke perceptions of dominance and fighting ability in men, it is less clear why they should do so. Does vocal masculinity advertise

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physical prowess? Attempts to answer this question have obtained mixed results.

Bruckert *et al.* [29] found that testosterone levels negatively predicted men's formant frequencies, and some have found that testosterone negatively predicted men's  $F_0$  [30,31], but others have not [29]. Because androgen is related to physical aggressiveness and competitiveness [32–34], a masculine voice may signal threat potential.

However, Sell *et al.* [35] found no relationships between men's fundamental or formant frequencies and upper body strength. Although some studies have found relationships between  $F_0$  and men's height [36] and weight [37], most have not [2,22,29,35,38–40]. Similarly, some studies have found relationships between formant structure and men's height [2,35,37,41], but others have not [22,29,42], and some have found relationships between formant structure and weight [37,42], but most have not [2,22,29,35]. To our knowledge, relationships between voice monotonicity and measures of physical threat potential are unexplored.

Rendall *et al.* [43] suggested that inferences of body size from  $F_0$  are misattributions resulting from (i) overgeneralizing age or sex differences, (ii) applying broader environmental sound-size associations to speech, or (iii) associations between behavioural traits (e.g. dominance) and both size and  $F_0$ . Similar reasoning could be applied to the inconsistent relationships between formant structure and men's physical characteristics. Alternatively, sexually dimorphic acoustic parameters may predict physical aggressive potential, including size, but varied results across studies could reflect differences in sample characteristics, small samples and varied methodologies.

Some failures to replicate associations with formant frequencies may result from the method of measuring formant structure. The most commonly used measure is 'formant dispersion' ( $D_f$ ), the average distance between successive formants [44]. Formant dispersion is given by

$$\frac{\sum_{i=1}^n (F_{i+1} - F_i)}{n - 1},$$

where  $F_i$  is the  $i$ th formant frequency, and  $n$  is the number of formant frequencies measured. Although  $D_f$  is highly sexually dimorphic in humans and has been 'the most reliable known acoustic index of body size in humans, as well as other primates' [35], it is not without problems. Perhaps most importantly,  $D_f$  surprisingly contains no information about the middle formants used in its calculation. Consider the case where the first four formants are measured. Formant dispersion, then, is

$$\frac{(F_2 - F_1) + (F_3 - F_2) + (F_4 - F_3)}{3}.$$

This reduces to  $(F_4 - F_1)/3$ , so  $D_f$  actually measures only the distance between the highest and lowest formants measured. In addition, because the variance in a higher formant (e.g.  $F_4$ ) greatly exceeds the variance in  $F_1$  [2], and because the formant values are not standardized,  $D_f$  is mainly driven by the value of the highest formant measured. Finally, given that longer vocal tracts produce lower formants, the sum of the first  $n$  formants should be inversely related to vocal tract length. Adding one formant and subtracting another, as in  $D_f$  calculation,

captures information about formant spacing but partly cancels information about formant positions.

The present study therefore seeks to clarify relationships between sexually dimorphic acoustic properties ( $F_0$ , monotonicity, formant structure) and threat potential in men. Such evidence would elucidate the processes of sexual selection in humans and non-human primates and contribute to a key objective of acoustic communication research: clarify the information content of vocal signals [45]. We first introduce a new measure of formant structure, 'formant position' ( $P_f$ ), which we show is more sexually dimorphic and more strongly related to body size than is  $D_f$  in both a US sample and in a sample of Hadza foragers from Tanzania. We then explore relationships between vocal masculinity and body size, strength, testosterone and physical aggressiveness in these samples.

## 4. STUDY 1: US SAMPLE

### (a) Methods

#### (i) Participants

One hundred and seventy-six male (mean age =  $20.1 \pm 1.7$ , range = 18–26) and 268 female (139 taking hormonal contraception: mean age =  $20.4 \pm 1.6$ , range = 18–26; 129 normally cycling: mean age =  $19.6 \pm 1.5$ , range = 18–24) students from a large, northeastern US university participated in this IRB-approved study. Participants identified as 91.4 per cent White, 3.4 per cent Asian, 1.8 per cent Hispanic or Latino, 1.4 per cent Black or African American, 0.5 per cent American Indian or Alaska Native, 0.2 per cent Native Hawaiian or Other Pacific Islander and 1.4 per cent 'other'.

#### (ii) Procedures

Male and hormonally contracepting female participants were scheduled for 1 h morning sessions (beginning between 08.20 and 10.00 h) and evening sessions (beginning between 17.20 and 19.00 h) one week apart. Normally cycling women attended sessions between 13.00 and 16.00 h and were scheduled based on self-reported menstrual cycle data to attend one session during the late follicular phase and one session during the mid-luteal phase. Session order (morning or evening first, or follicular or mid-luteal first) was randomized. Voice, anthropometric, hormonal (in this study, men's testosterone) and psychometric data were collected at both sessions.

#### (iii) Voice recording and measurement

Participants were recorded reading an excerpt from a standard voice passage, the Rainbow Passage [46], in an anechoic, soundproof booth using a Shure SM58 vocal cardioid microphone. A curved wire projection from the microphone stand kept the participant's mouth approximately 9.5 cm from the microphone. Voices were recorded into a computer using GOLDWAVE software in mono at a sampling rate of 44 100 Hz and 16 bit quantization, and saved as uncompressed '.wav' files.

Each recording (mean duration =  $30.6 \pm 3.8$  s) was analysed using PRAAT software (v. 4.4.11). For mean  $F_0$  and  $F_0$  variation across the utterance (standard deviation in  $F_0$ ,  $F_0 - \text{s.d.}$ ), pitch floors were 75 and 100 Hz, and pitch ceilings were 300 and 500 Hz, for men and women, respectively, in accordance with the programmers' recommendations [47]. Otherwise, default settings were used.

We measured  $F_1$  through  $F_4$  at each glottal pulse (automated detection by Praat, mean glottal pulses per recording =  $2972 \pm 889$ ) and averaged across measurements. This facilitated formant measurement along the entire utterance, thus sampling a greater range of vocal tract configurations than measuring individual vowels. In addition, this method sampled only voiced speech and avoided fricatives, which artificially decrease apparent vocal tract length because the sound source of fricatives is turbulence in the mouth rather than vocal fold vibration [48]. Because Praat occasionally shifts formants (e.g. misattributing  $F_2$  as  $F_1$ ), we omitted all formant measurements from glottal pulses for which any value exceeded a predetermined threshold (less than 2% of pulses). Thresholds were based on published data [2] for male and female formants and were selected to eliminate only clearly erroneous measurements. For men, thresholds were 1000, 2850, 3750 and 4500 Hz for  $F_1$  through  $F_4$ , and for women, thresholds were 1250, 3350, 4150 and 5100 Hz. Formant measurements obtained by this method correlated highly with measurements obtained by measuring and averaging across 11 individual vowels each from nine randomly selected men and nine randomly selected women:  $0.93 \leq r \leq 0.98$  (both sexes),  $0.86 \leq r \leq 0.94$  (men only).

We then computed formant dispersion ( $D_f$ ) and formant position ( $P_f$ ). We define  $P_f$  as the average standardized formant value for the first  $n$  formants, where formants are standardized using between-sex means and standard deviations. Thus,

$$P_f = \frac{\sum_{i=1}^n F'_i}{n},$$

where  $F'_i$  is the standardized  $i$ th formant, and  $n$  is the number of formants measured. This method effectively assigns each standardized formant a unit weight rather than, for example, a beta weight obtained via regressing formants on sex or height. We chose this approach following Cohen [49], who suggested that unit weights have better predictive power than beta weights derived from moderate-sized samples. So as not to bias the mean or standard deviation towards either sex during standardization, we obtained means and standard deviations using bootstrapping methods, randomly selecting (with replacement) 10 000 samples of 129 men and all 129 normally cycling women who provided voice recordings for both sessions. With both sexes included, mean  $F_1 = 472.3 \pm 49.7$  Hz, mean  $F_2 = 1594.4 \pm 146.7$  Hz, mean  $F_3 = 2604.5 \pm 184.4$  Hz and mean  $F_4 = 3612.8 \pm 267.9$  Hz. These values were then used to standardize formants for all participants.

#### (iv) Anthropometry

Flexed biceps circumference was measured at its widest point for left and right arms using a tape measure. Left- and right-hand strengths were obtained using a JAMAR hydraulic hand dynamometer. Biceps size and hand strength are good predictors of overall upper body strength [35]. Height was measured from a metre stick affixed to a wall, and weight was obtained using an electronic scale.

#### (v) Testosterone assays

Saliva was collected for testosterone (T) assays during morning and evening sessions. Contamination of saliva

samples was minimized by having participants not eat, drink (except plain water), smoke, chew gum or brush their teeth for 1 h before their session. Participants rinsed their mouths with water before chewing a piece of sugar-free Trident gum (inert in salivary hormone assays) to stimulate saliva flow. Approximately 9 ml of saliva was collected in a sodium azide-coated polystyrene tube. The tube was capped and left upright at room temperature for 18–24 h to allow mucins to settle. Tubes were then frozen at  $-20^\circ\text{C}$  until hormone analysis.

We obtained salivary unbound ('free') T concentrations, which correlate strongly with serum T concentrations [50]. The Salivary Radioimmunoassay Laboratory at the University of Western Ontario performed T radioimmunoassay on 330 male saliva samples, 175 from session 1 and 155 from session 2. All samples went through double ether extraction, followed by radioimmunoassay in duplicate using a Coat-A-Count kit for total T (Diagnostic Products, Los Angeles, CA, USA), modified for use with saliva (for details, see [51]). The average intra-assay coefficient of variation was 6.3 per cent, and sensitivity was  $5\text{--}10\text{ pg ml}^{-1}$ .

#### (vi) Psychometric data collection

Following anthropometry and saliva collection, each participant completed the Buss & Perry [52] Aggression Questionnaire, and instruments not used here, at a private computer station. The Aggression Questionnaire comprises 29 items, including nine targeting physical aggression (e.g. 'Once in a while I can't control the urge to strike another person'). Items are assessed on a 5-point scale, anchored at 'extremely uncharacteristic of me' and 'extremely characteristic of me'. Scores were summed for a composite measure of physical aggression.

## 5. RESULTS

### (a) Correlations across sessions and sides of the body

Mean  $F_0$  and  $F_0 - \text{s.d.}$  correlated across sessions  $r_{391} = 0.98$  and  $0.92$ , respectively (controlling for sex,  $r_{388} = 0.83$  and  $0.78$ , respectively). Formants correlated across sessions  $0.92 \leq r_{391} \leq 0.98$  (controlling for sex,  $0.82 \leq r_{388} \leq 0.89$ ). Men's anthropometric measurements (women's not reported here) correlated  $0.68 \leq r_{155} \leq 0.99$  across sessions and  $0.74 \leq r_{155} \leq 0.96$  across sides of the body. Men's physical aggression scores (Cronbach's  $\alpha = 0.85$ ) correlated across sessions  $r_{154} = 0.90$ .

Men's testosterone concentrations correlated more modestly across morning and evening sessions ( $r_{154} = 0.50$ ), as expected given temporal variability in T secretion. Testosterone concentrations also decreased significantly from morning to evening sessions (paired  $t_{153} = 10.4$ ,  $p < 0.0001$ ), indicating that we were able to capture significant diurnal variation.

Because all measurements highly correlated across sessions and sides of the body, we averaged across sessions and sides of the body (tables 1 and 2). Only values from session 1 were used for participants who did not return for session 2 (9% of cases).

### (b) Sexual dimorphism in the voice

All acoustic parameters were highly sexually dimorphic ( $2.7 \leq d \leq 5.7$ , table 1). Importantly,  $P_f$  ( $d = 5.4$ ) was

Table 1. Sex differences in voice measurements.

		women ( <i>n</i> = 268)	men ( <i>n</i> = 176)	<i>t</i>	<i>d</i>
study 1	$F_0$ (Hz)	201.7 ± 18.0	111.4 ± 14.2	58.8***	5.7
	$F_0$ - s.d. (Hz)	35.5 ± 8.5	15.0 ± 3.6	35.1***	3.6
	$F_1$ (Hz)	511.8 ± 34.3	434.3 ± 26.7	26.7***	2.6
	$F_2$ (Hz)	1724.9 ± 64.3	1456.0 ± 55.7	45.4***	4.3
	$F_3$ (Hz)	2781.3 ± 91.9	2442.9 ± 78.5	41.5***	4.1
	$F_4$ (Hz)	3874.0 ± 125.6	3390.9 ± 108.9	43.0***	4.3
	$D_f$ (Hz)	1120.7 ± 41.6	985.5 ± 39.0	34.3***	3.3
	$P_f$	0.90 ± 0.38	-0.85 ± 0.27	56.9***	5.4
		women ( <i>n</i> = 43)	men ( <i>n</i> = 32)	<i>t</i>	<i>d</i>
study 2	$F_0$ (Hz)	208.5 ± 29.4	114.9 ± 15.3	17.9***	4.4
	$F_0$ - s.d. (Hz)	28.8 ± 11.3	14.7 ± 5.7	7.3***	1.8
	$F_1$ (Hz)	454.0 ± 62.4	418.1 ± 62.5	2.5*	0.6
	$F_2$ (Hz)	1431.9 ± 123.2	1286.1 ± 120.8	5.1***	1.2
	$F_3$ (Hz)	2970.9 ± 153.2	2611.4 ± 149.4	10.2***	2.4
	$F_4$ (Hz)	4000.1 ± 121.8	3591.1 ± 149.9	13.3***	3.1
	$D_f$ (Hz)	1174.6 ± 47.5	1058.9 ± 51.8	10.0***	2.3
	$P_f$	0.68 ± 0.43	-0.80 ± 0.40	15.3***	3.6

\* $p < 0.05$ .\*\*\* $p < 0.0001$ .

Table 2. Descriptive statistics (mean (s.d., range)) for male participants in US and Hadza samples.

variable	US sample	Hadza sample
height (cm)	178.8 (6.9, 158.5–197.3)	162.5 (7.2, 145.8–179.3)
weight (kg)	78.7 (13.4, 53.1–131.9)	51.4 (6.3, 38.8–67.2)
biceps (cm)	33.4 (3.2, 26.1–42.0)	25.1 (1.6, 22.0–28.5)
hand strength (kg)	49.5 (8.1, 31.3–69.3)	35.0 (5.6, 21.5–45.0)
triceps skinfold (cm)	—	5.3 (1.8, 3.0–9.7)
physical aggression	21.8 (6.6, 9–39)	—
T levels (pg ml <sup>-1</sup> )	92.9 (34.2, 36.3–254.5)	—

more sexually dimorphic than  $D_f$  ( $d = 3.3$ ) (test for difference between dependent correlation coefficients (correlations between sex and  $P_f$  and  $D_f$ ):  $t_{441} = 8.97$ ,  $p < 0.0001$ ). When sex and height were entered simultaneously to predict  $P_f$  and  $D_f$  in separate general linear models, sex more strongly predicted  $P_f$  controlling for height (partial  $\eta^2 = 0.73$  compared with 0.54 for  $D_f$ , both  $p < 0.0001$ ), and height more strongly predicted  $P_f$  controlling for sex (partial  $\eta^2 = 0.11$ ,  $p < 0.0001$  compared with 0.01 for  $D_f$ ,  $p = 0.021$ , electronic supplementary material, table S1). All other acoustic parameters remained highly sexually dimorphic after controlling for height ( $0.36 \leq \text{partial } \eta^2 \leq 0.77$ ).

### (c) Correlates of acoustic parameters

In subsequent analyses,  $F_0$  - s.d., weight, physical aggression and T levels were log-transformed to correct for positive skew. Hand strength and biceps circumference were standardized and then averaged to produce the composite measure 'arm strength'.

Measures of threat potential (height, weight, arm strength, physical aggression and testosterone), along with acoustic parameters as control variables, were entered simultaneously into separate multiple regressions (table 3) to explore their independent relationships to each sexually dimorphic acoustic parameter ( $F_0$ ,

$F_0$  - s.d. and  $P_f$ ). Age was not significantly correlated with any acoustic parameter (all  $p > 0.3$ ), so it was not included in these models. All variance inflation factors (VIFs) were less than 1.9. In addition, we explored zero-order correlations between measures of threat potential and acoustic parameters (table 4).

In the regression model for mean  $F_0$  ( $F_{7,167} = 15.99$ ,  $p < 0.0001$ ), testosterone ( $t = -2.87$ ,  $\beta = -0.17$ ,  $p = 0.005$ ), height ( $t = -2.46$ ,  $\beta = -0.16$ ,  $p = 0.015$ ) and  $F_0$  - s.d. ( $t = 9.51$ ,  $\beta = 0.60$ ,  $p < 0.0001$ ) were significant predictors. In zero-order correlations, mean  $F_0$  was significantly negatively related to height ( $r_{175} = -0.17$ ,  $p = 0.028$ ) and testosterone ( $r_{175} = -0.15$ ,  $p = 0.044$ ), although these correlations do not survive Bonferroni correction for multiple tests (five predictors).

Physical aggression ( $t = -3.40$ ,  $\beta = -0.20$ ,  $p < 0.001$ ) and mean  $F_0$  ( $t = 9.67$ ,  $\beta = 0.58$ ,  $p < 0.0001$ ) significantly predicted  $F_0$  - s.d. in the regression model for this parameter ( $F_{2,173} = 53.90$ ,  $p < 0.0001$ ). In zero-order correlations,  $F_0$  - s.d. was significantly negatively correlated with physical aggression ( $r_{175} = -0.23$ ,  $p = 0.003$ ) and marginally significantly correlated with arm strength ( $r_{175} = -0.15$ ,  $p = 0.053$ ).

Height ( $t = -2.11$ ,  $\beta = -0.17$ ,  $p = 0.036$ ) significantly predicted  $P_f$  in the final model for  $P_f$  ( $F_{7,167} = 4.22$ ,  $p < 0.001$ ). In zero-order correlations,  $P_f$  was significantly negatively correlated with height ( $r_{175} = -0.24$ ,



Table 3. Predictors of men's vocal acoustic parameters—multiple regression results.

		study 1		study 2	
		<i>t</i>	beta	<i>t</i>	beta
$F_0$	height	-2.46*	-0.16	-2.39*	-0.66
	weight	0.05	<0.01	2.53*	0.94
	arm strength	0.71	0.06	-2.16*	-0.47
	physical aggression	1.19	0.08	—	—
	testosterone	-2.87**	-0.17	—	—
	$F_0$ - s.d.	9.51***	0.60	5.09***	0.64
	$P_f$	0.15	0.01	1.15	0.17
$F_0$ - s.d.	height	0.82	0.05	1.13	0.37
	weight	1.64	0.13	-1.15	-0.53
	arm strength	-1.78 <sup>+</sup>	-0.14	1.01	0.26
	physical aggression	-2.53*	-0.16	—	—
	testosterone	1.56 <sup>+</sup>	0.09	—	—
	$F_0$	9.51***	0.58	5.09***	0.80
	$P_f$	1.52	0.10	-0.33	-0.05
$P_f$	height	-2.11*	-0.17	0.49	0.21
	weight	-0.79	-0.08	-1.45	-0.84
	arm strength	-1.43	-0.13	1.05	0.35
	physical aggression	-1.31	-0.10	—	—
	testosterone	1.06	0.08	—	—
	$F_0$	0.15	0.01	1.15	0.32
	$F_0$ - s.d.	1.52	0.14	-0.33	-0.08

<sup>+</sup> $p < 0.1$ .\* $p < 0.05$ .\*\* $p < 0.01$ .\*\*\* $p < 0.001$ .Table 4. Predictors of men's vocal acoustic parameters—zero-order correlations (and  $n$ ).

		$F_0$	$F_0$ - s.d.	$P_f$	$D_f$
study 1	height ( $n = 175$ )	-0.17*	-0.06	-0.24**	-0.02
	weight ( $n = 175$ )	0.01	0.02	-0.23**	-0.01
	arm strength ( $n = 176$ )	-0.04	-0.15 <sup>+</sup>	-0.26***	-0.02
	physical aggression ( $n = 176$ )	-0.04	-0.23**	-0.19**	-0.02
	testosterone ( $n = 176$ )	-0.15*	0.01	0.11	0.05
study 2	height ( $n = 32$ )	-0.20	-0.09	-0.38*	-0.11
	weight ( $n = 32$ )	-0.08	-0.04	-0.39*	-0.04
	arm strength ( $n = 31$ )	-0.17	-0.07	-0.23	0.10

<sup>+</sup> $p < 0.1$ .\* $p < 0.05$ .\*\* $p < 0.01$ .\*\*\* $p < 0.001$ .

$p = 0.001$ ), weight ( $r_{175} = -0.23$ ,  $p = 0.002$ ), physical aggressiveness ( $r_{175} = -0.19$ ,  $p < 0.01$ ) and arm strength ( $r_{175} = -0.26$ ,  $p < 0.001$ ).

No variable significantly or marginally significantly predicted  $D_f$  (regression model:  $F_{7,167} = 0.16$ ,  $p = 0.992$ , zero-order correlations: all  $p > 0.4$ ).

## 6. STUDY 2: HADZA SAMPLE

### (a) Methods

#### (i) Participants

The Hadza occupy remote savannah-woodland areas in northern Tanzania around Lake Eyasi and subsist on game killed with bow and arrow and foraged foods such as berries, baobab fruit and tubers. The Hadza number approximately 1000 and live in camps of approximately

30 individuals. Data were gathered from participants in eight Hadza camps. Thirty-two Hadza men (mean age =  $29.6 \pm 6.2$ , range = 19–40) and 43 Hadza women (mean age =  $28.9 \pm 6.3$ , range = 18–39) participated in this IRB-approved study.

#### (ii) Voice recording and measurement

Participants were recorded speaking the word 'hujambo', which loosely translates to 'hello' in English, inside a Land Rover with a Sennheiser MKH-60 microphone. Recordings were encoded in mono directly onto computer hard disk, using Sonic Foundry's SOUND FORGE at 44 100 Hz sampling rate and 16 bit quantization, and saved as uncompressed '.wav' files.

Voice recordings were measured using methods identical to those in study 1 (table 1). Mean number of glottal

pulses per recording was  $128 \pm 73$ . Less than 3 per cent of glottal pulses were excluded owing to formant misattributions in Praat. Formant means and standard deviations used to compute  $P_f$  were obtained using bootstrapping methods from 10 000 random samples with replacement of 32 women and all 32 men. With both sexes included, mean  $F_1 = 425.3 \pm 49.8$  Hz, mean  $F_2 = 1372.8 \pm 138.1$  Hz, mean  $F_3 = 2794.4 \pm 221.1$  Hz and mean  $F_4 = 3779.2 \pm 240.3$  Hz.

### (iii) Anthropometry

Height was measured using a portable stadiometer, and weight was obtained using an electronic scale (Tanita Ultimate Scale 2000). Triceps skinfold measurements for the left arm were made in triplicate (Cronbach's  $\alpha = 0.96$ ) using Mitutoyo Absolute Digimatic CD 6 calipers and then averaged. Mid-upper arm circumference for the left arm was measured using a flexible tape measure. A dynamometer was used to measure left- and right-hand strengths, which correlated  $r_{31} = 0.83$  and were thus averaged. (See table 2 for descriptive statistics.) Upper arm muscle mass was computed from the formula

$$\frac{(a - \pi s)^2}{4\pi} - 10,$$

where  $a$  is the mid-upper arm circumference (cm), and the  $s$  is the average of three triceps skinfold measurements (cm). Hand strength and upper arm muscle mass were then standardized and averaged to produce the composite measure arm strength.

## 7. RESULTS

### (a) Sexual dimorphism in the voice

All acoustic parameters were highly sexually dimorphic ( $1.5 \leq d \leq 4.4$ , table 1). Importantly,  $P_f$  ( $d = 3.6$ ) was more sexually dimorphic than  $D_f$  ( $d = 2.4$ ) (test for difference between dependent correlation coefficients (correlations between sex and  $P_f$  and  $D_f$ ):  $t_{72} = 2.81$ ,  $p = 0.003$ ). When both sex and height were entered to predict  $P_f$  and  $D_f$  in separate general linear models, sex more strongly predicted  $P_f$  controlling for height (partial  $\eta^2 = 0.59$  compared with 0.41 for  $D_f$ , both  $p < 0.0001$ ), and height more strongly predicted  $P_f$  controlling for sex (partial  $\eta^2 = 0.11$ ,  $p = 0.004$  compared with partial  $\eta^2 < 0.01$ ,  $p > 0.5$  for  $D_f$ , electronic supplementary material, table S1). All other acoustic parameters remained highly sexually dimorphic after controlling for height ( $0.20 \leq$  partial  $\eta^2 \leq 0.66$ ). In subsequent analyses,  $F_0 - \text{s.d.}$  was log-transformed to correct positive skew.

### (b) Correlates of acoustic parameters

Measures of threat potential (height, weight and arm strength), along with acoustic parameters as control variables, were entered simultaneously into separate multiple regressions (table 3) to explore independent relationships to each sexually dimorphic acoustic parameter ( $F_0$ ,  $F_0 - \text{s.d.}$  and  $P_f$ ). Age was not significantly correlated with any acoustic parameter (all  $p > 0.1$ ), so it was not included in these models. The VIF for weight was between 9.4 and 11.1, indicating that some caution should be used in interpreting results for this variable [53]. In addition, we explored zero-order correlations between measures of threat potential and acoustic parameters (table 4).

In the regression model for mean  $F_0$  ( $F_{5,25} = 8.38$ ,  $p < 0.0001$ ), height ( $t = -2.39$ ,  $\beta = -0.66$ ,  $p = 0.025$ ), weight ( $t = 2.53$ ,  $\beta = 0.94$ ,  $p = 0.018$ ), arm strength ( $t = -2.16$ ,  $\beta = -0.47$ ,  $p = 0.041$ ) and  $F_0 - \text{s.d.}$  ( $t = 5.09$ ,  $\beta = 0.64$ ,  $p < 0.0001$ ) were significant predictors. In zero-order correlations, a lower, more masculine  $F_0$  was associated with greater threat potential for all measures, but no correlation attained statistical significance. Although no variable significantly predicted  $P_f$  in the model for  $P_f$  ( $F_{5,25} = 1.51$ ,  $p = 0.222$ ), a more masculine  $P_f$  was associated with a significantly greater height ( $r_{32} = -0.38$ ,  $p = 0.030$ ) and weight ( $r_{32} = -0.39$ ,  $p = 0.026$ ) in zero-order correlations. However, these relationships are only marginally statistically significant after Bonferroni correction for multiple tests (three predictors). Only mean  $F_0$  significantly predicted  $F_0 - \text{s.d.}$  ( $t = 5.09$ ,  $\beta = 0.80$ ,  $p < 0.0001$ ) in the model for this variable ( $F_{5,25} = 5.65$ ,  $p = 0.001$ ), and no zero-order correlation between  $F_0 - \text{s.d.}$  and measures of threat potential attained statistical significance. No variable significantly or marginally significantly predicted  $D_f$  in the regression model ( $F_{5,25} = 0.34$ ,  $p = 0.886$ ) or in zero-order correlations (all  $p > 0.5$ ).

## 8. DISCUSSION

We proposed a novel method of measuring formant structure, formant position ( $P_f$ ), and showed in two diverse human samples that  $P_f$  was more sexually dimorphic than the most widely used measure of formant structure, formant dispersion ( $D_f$ ). In both samples,  $P_f$  was more sexually dimorphic than  $D_f$  when height was held constant, and more strongly related to height when sex was held constant. Mean fundamental frequency ( $F_0$ ) and the variation in  $F_0$  across an utterance ( $F_0 - \text{s.d.}$ ) were also highly sexually dimorphic in both populations. Especially remarkable was the finding that  $F_0 - \text{s.d.}$  differed by sex when measured from a single three-syllable word (hujambo). This variable was highly sexually dimorphic whether measured in Hertz or in equivalent rectangular bandwidths (ERBs), a unit that scales more nearly linearly with perception [54]. Thus, it is unlikely that women's  $F_0$  varies more dramatically simply because, at the higher frequencies at which women speak, larger changes are required for the same perceptual effects.

Sex differences in mean  $F_0$  and formant frequencies depend largely on the anatomy of the vocal apparatus [2]. By contrast, it seems likely that the sex difference in  $F_0 - \text{s.d.}$  mainly reflects neuropsychological differences affecting control of the vocal apparatus. In this regard, a highly varied  $F_0$  may be analogous to smiling—both signal affiliation and deference [23], and both are expressed to a greater degree among women [55]. Perhaps reliably emerging neurodevelopmental processes, possibly influenced by sex hormones, underlie the apparent cross-cultural ubiquity of both sex differences, but these processes are also modifiable by social learning [55,56].

Each sexually dimorphic acoustic parameter in each sample was predicted by one or more measures of men's physical threat potential, with the exception of  $F_0 - \text{s.d.}$  in the Hadza sample. This last result is unsurprising, given the modest sample size and paucity of vocal information (one three-syllable word) available for assessing individual differences in  $F_0 - \text{s.d.}$  In the larger US

sample with more vocal information, a low (masculine)  $F_0$  – s.d. was related to increased physical aggressiveness and marginally significantly greater arm strength.

Taller men with higher testosterone levels spoke in a lower (more masculine)  $F_0$  in the US sample. Hadza men who spoke in a lower  $F_0$  were also taller and were stronger. Interestingly, they also tended to be lighter, with height and arm strength statistically controlled, suggesting less body fat. However, entering skinfold thickness into the model did not change these relationships.

Across men and women in both US and Hadza samples, controlling for sex, taller individuals spoke with a more masculine (lower)  $P_f$ , indicating longer vocal tracts. In the US sample, men who spoke with a lower  $P_f$  were taller, heavier, stronger and reported more physical aggression. Only height predicted a lower  $P_f$  when other predictors were held statistically constant via multiple regression, although the non-significance of other predictors probably reflects collinearity. For example, when height and arm strength were entered alone into a multiple regression model, both were significant predictors ( $p < 0.01$ ). In the Hadza sample, taller and heavier men also spoke with a more masculine  $P_f$ , although these associations were only marginally statistically significant after Bonferroni correction for multiple tests. No measure of physical formidability significantly predicted  $P_f$  in the multiple regression model, again probably reflecting collinearity among predictor variables ( $0.43 < r < 0.85$ , for correlations among height, weight and strength).

Height significantly predicted  $D_f$  only in the US dataset with both sexes included, perhaps owing to the increased sample size.  $D_f$  was unrelated to any other measure of physical formidability in either US or Hadza samples.

## 9. CONCLUSION

Lower fundamental and formant frequencies in men reflect the hypertrophic growth of the male larynx [8] and its descent in the trachea a full vertebra lower than in women [57]. These pubertal changes may have evolved in order to exaggerate body size [44], but same-sex competitors and potential mates should evolve to attend to signals in proportion to the signals' reliability at communicating useful information. Our results support the notion that certain acoustic features of men's voices are honest signals of threat potential. It is possible that adult physiology and/or developmental influences such as pubertal testosterone and growth hormone affect both vocal anatomy and physical prowess. Certain psychological states (e.g. self-confidence) may also modulate the position of the larynx and vocal fold tension, and these psychological states may be related to physical formidability. These results thus help clarify widespread attributions of physical prowess to men with masculine voices. The two most sexually dimorphic acoustic features,  $F_0$  and  $P_f$ , in particular showed the strongest relationships with size, strength and testosterone levels across our samples.

Each of our samples has advantages and disadvantages, for example, greater sample size (the largest of which we are aware to explore such relationships) and tighter methodological control in the US sample versus

greater ecological validity in the Hadza sample. Our results agree with some previous results and contrast with others, highlighting the need for future work to address these discrepancies. One result contrasting with previous research was the significant negative relationship that we observed in both samples between men's height and  $F_0$ . We note that the zero-order correlation was not significant in the Hadza sample (though it was of greater magnitude than in the US sample:  $r = -0.20$  versus  $-0.17$ ) and suggest that the discrepancy is due to sample size. Similarly, the weighted mean correlation between men's height and  $F_0$  measured from continuous speech was approximately  $-0.17$  across four samples reported by Sell *et al.* [35], yet individual sample sizes were modest (20–63), and no correlation attained statistical significance.

Future research should explore additional correlates of physical competitive ability (e.g. past success in fights and peer rankings of physical formidability), particularly in traditional societies. Future work should especially address  $F_0$  – s.d., as little evidence is available on this promising acoustic parameter, and should explore the utility of  $P_f$  as an inverse measure of apparent vocal tract length and vocal masculinity. In unisex samples, within-sex means and standard deviations may be used in calculating  $P_f$  values; doing so had a minimal effect on the present results. Finally, researchers should continue to investigate the information content of these acoustic parameters across non-human primates.

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