

AperTO - Archivio Istituzionale Open Access dell'Università di Torino

**Artificial Neural Network approach for revealing individuality, group membership and age information in goat kid contact calls**

**This is the author's manuscript**

*Original Citation:*

*Availability:*

This version is available <http://hdl.handle.net/2318/141921> since 2022-07-18T12:34:50Z

*Published version:*

DOI:10.3813/AAA.918758

*Terms of use:*

Open Access

Anyone can freely access the full text of works made available as "Open Access". Works made available under a Creative Commons license can be used according to the terms and conditions of said license. Use of all other works requires consent of the right holder (author or publisher) if not exempted from copyright protection by the applicable law.

(Article begins on next page)



# UNIVERSITÀ DEGLI STUDI DI TORINO

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12

***This is an author version of the contribution published on:***

*Questa è la versione dell'autore dell'opera:*

*ACTA ACUSTICA UNITED WITH ACUSTICA Vol. 100 (2014) 782 – 789*

*DOI:10.3813/AAA.918758*

***The definitive version is available at:***

*La versione definitiva è disponibile alla URL:*

*[http://www.ingentaconnect.com/content/dav/aaua/2014/00000100/00000004/  
art00023](http://www.ingentaconnect.com/content/dav/aaua/2014/00000100/00000004/art00023)*

13        **Artificial Neural Network approach for revealing individuality, group**  
14                    **membership and age information in goat kid contact calls**

15

16    **Authors and affiliations**

17    Livio Favaro<sup>1</sup>, Elodie F. Briefer<sup>2</sup>, Alan G. McElligott<sup>3</sup>

18

19    <sup>1</sup> Department of Life Sciences and Systems Biology, University of Torino, Via Accademia Albertina  
20    13, 10123 Turin, Italy. livio.favaro@unito.it

21    <sup>2</sup> Institute of Agricultural Sciences, ETH Zürich, Universitätstrasse 2, 8092 Zürich, Switzerland.  
22    elodie.briefer@usys.ethz.ch

23    <sup>3</sup> Biological and Experimental Psychology, School of Biological and Chemical Sciences, Queen  
24    Mary University of London, Mile End Road, London E1 4NS, UK. a.g.mcelligott@qmul.ac.uk

25

26    **Corresponding authors:** LF and AGM

27 **Abstract**

28 Machine learning techniques are becoming an important tool for studying animal vocal  
29 communication. The goat (*Capra hircus*) is a very social species, in which vocal communication  
30 and recognition are important. We tested the reliability of a Multi-Layer Perceptron (feed-forward  
31 Artificial Neural Network, ANN) to automate the process of classification of calls according to  
32 individual identity, group membership and maturation in this species. Vocalisations were obtained  
33 from 10 half-sibling (same father but different mothers) goat kids, belonging to 3 distinct social  
34 groups. We recorded 157 contact calls emitted during first week, and 164 additional calls recorded  
35 from the same individuals at 5 weeks. For each call, we measured 27 spectral and temporal  
36 acoustic parameters using a custom built program in Praat software. For each classification task  
37 we built *stratified* 10-fold cross-validated neural networks. The input nodes corresponded to the  
38 acoustic parameters measured on each signal. ANNs were trained with the error-back-propagation  
39 algorithm. The number of hidden units was set to the number of attributes + classes. Each model  
40 was trained for 300 epochs (learning rate 0.2; momentum 0.2). To estimate a reliable error for the  
41 models, we repeated 10-fold cross-validation iterations 10 times and calculated the average  
42 predictive performance. The accuracy was  $71.13 \pm 1.16\%$  for vocal individuality,  $79.59 \pm 0.75\%$  for  
43 social group and  $91.37 \pm 0.76\%$  for age of the vocalising animal. Our results demonstrate that  
44 ANNs are a powerful tool for studying vocal cues to individuality, group membership and  
45 maturation in contact calls. The performances we achieved were higher than those obtained for the  
46 same classification tasks using classical statistical methods such as Discriminant Function  
47 Analysis. Further studies, investigating the reliability of these algorithms for the real-time  
48 classification of contact calls and comparing ANNs with other machine learning techniques are  
49 important to develop technology to remotely monitor the vocalisations of domestic livestock.

50

51 **Keywords**

52 bioacoustics, goats, livestock, machine learning, ungulates, vocal communication

53

54 **PACS no.** 43.80.Ka

## 55 **1. Introduction**

56 A crucial step in understanding animal vocal behaviour is the description and quantification of  
57 similarities and differences among acoustic signals [1, 2]. This step is essential in order to identify  
58 biologically meaningful categories of sound [1]. Indeed, several vocalisations encode a variety of  
59 information about animal sex, body size, age and even social status [3, 4, 5]. Vocal signal  
60 categorisation may also allow the detection of social context-dependent variability, ecological  
61 diversity, species recognition, and vocal individuality [6, 7]. Characterising animal sounds can also  
62 provide information on genetic and evolutionary relationships among different taxonomic units [8].

63  
64 Traditionally, the classification of animal vocal signals has been performed using subjective  
65 methods [9], such as signal classification by multiple listeners using their pattern recognition  
66 abilities (e.g. the abilities of human observers to recognise vocal categories using their auditory  
67 system). More recently, technological improvements have allowed detailed acoustic  
68 measurements on recorded vocal parameters, followed by automated classification using statistical  
69 methods. These later, more advanced techniques, include Multivariate Analysis of Variance [10],  
70 Discriminant Function Analysis [11], Hierarchical Cluster Analysis [12], and Principal Components  
71 Analysis [13]. However, statistical methods frequently fail to detect biologically meaningful  
72 information in vocalisations [1]. A modern and alternative approach is to use mathematical  
73 computational techniques. Among these, machine learning (ML) algorithms have been suggested  
74 as an attractive, non-linear alternative to traditional statistical analyses [14, 15]. The biggest  
75 advantage of ML techniques is their ability to model complex and non-linear relationships among  
76 acoustic parameters without having to satisfy the restrictive assumptions required by conventional  
77 parametric approaches. Moreover, they allow modelling of non-linear associations with a variety of  
78 data types, and accommodate interactions among predictor variables with limited *a priori*  
79 specifications [16].

80  
81 More recently, the reliability of ML techniques for solving complex pattern recognition problems has  
82 been demonstrated in many ecological [i.e. 17], biomedical [i.e. 18] and behavioural studies [i.e.

83 19]. Although the application of these approaches to bioacoustics has increased in the last decade,  
84 the growth has been slower than in other disciplines, and there is still a good degree of scepticism  
85 with respect to the role of these techniques in quantitative analyses [20].

86

87 Artificial Neural Networks (ANNs) are the most common ML methods used for classification and  
88 recognition of mammal vocalisations. These algorithms were firstly introduced in marine  
89 bioacoustics to study the sonar system of bottlenose dolphins, *Tursiops truncatus* [21], and were  
90 further used for recognition of vocal units, caller and species in many different marine mammals [1,  
91 22, 23, 24]. ANNs have been successfully used to identify echolocating bat species [25], to classify  
92 several non-human primates vocalisations [26, 27], Gunnison's prairie dog, *Cynomys gunnisoni*  
93 [28], fallow deer, *Dama dama* [14], and even stress-linked calls of domestic pigs, *Sus scrofa* [29].

94

95 Domestic goats are very social animals, and vocal communication and recognition are important  
96 for social bonding and group cohesion [5, 30]. Goat kids produce one basic call type, the “contact  
97 call”, when isolated at short distance from other group members [5]. According to the source-filter  
98 theory of voice production [31, 32], calls are generated by vibrations of the vocal folds (source,  
99 determining the fundamental frequency, “F0”) and are subsequently filtered by the supralaryngeal  
100 vocal tract (filter, resulting in amplified frequencies called “formants”). In goat kid vocalisations, F0  
101 and formants parameters are known to be good indicators of caller identity [33], group membership  
102 [30], body size, sex and even age [5].

103

104 In this study, we tested the reliability of a Multi-Layer Perceptron (MLP), feed forward ANN, to  
105 automate classification of calls according to individual identity, group membership and maturation  
106 in a livestock species: the goat (*Capra hircus*). Our aim was to determine whether the MLP  
107 performances were better than those obtained for the same classification tasks using classical  
108 statistical methods such as Discriminant Function Analysis and, therefore, should be adopted in  
109 future vocal communication studies.

## 110 **2. Methods**

### 111 *2.1 Animals and recordings*

112 Contact calls were collected from 10 goat kids (9 males and 1 female), belonging to 3 distinct  
113 social groups at White Post Farm, Nottinghamshire, UK (53°06'N, 1°03'W). Goats use both open  
114 and closed mouth contact calls but, for the purpose of this study, we used only open mouthed calls,  
115 since closed mouth calls suppress or modify some formants [5]. All kids were half-sibling (same  
116 father but different mothers) born in July and December 2009, and March 2010, respectively. Each  
117 group (6.00 +/- 0.97 kids per group, mean +/- SE) was housed in an indoor communal pen of 4.4 m  
118 x 4.5 m. Vocalisations were recorded from the same individuals both early after parturition (1 week  
119 ± 5 days), and when young kids (5 weeks ± 5 days). To promote contact call production, we  
120 isolated kids from their mothers for 5 min periods, 2-3 times per day between 10 and 5 pm. The  
121 distance to the mother was set at 1 m (on average), during the first day of recordings and  
122 increased afterwards if necessary, until we obtained contact calls [i.e. low-affect vocalizations, 5]  
123 instead of distress calls [i.e. high-pitched vocalizations associated with high stress levels, 5]. Kids  
124 were isolated alone, except if they showed signs of stress during isolation even at 1 m. In these  
125 cases, they were isolated with their sibling(s).

126

127 Recordings were collected with a Sennheiser MKH70 directional microphone (frequency response  
128 50 Hz to 20 kHz ± 2.5 dB) connected to a Marantz PMD660 digital recorder (sampling rate set to  
129 44.1 kHz). During recording sessions, the microphone was placed at distances of 1 - 5 m from the  
130 vocalising animal. Segments containing acoustic recordings were saved in WAV format (16-bit  
131 amplitude resolution) and stored into an SD memory card. All the files were then transferred to a  
132 computer for later acoustic analyses.

133

### 134 *2.2 Acoustic analysis*

135 For each file, the waveform and FFT spectrogram (window length = 0.01 s, time steps = 1000,  
136 frequency steps = 250, Gaussian window shape, dynamic range = 50 dB) were generated in  
137 Seewave [34]. After visual examinations of sonograms, calls with high background noise levels

138 were discarded. Among the remaining vocalizations, we selected 157 good quality contact calls  
139 (13 to 23 calls per individual) emitted during early postnatal days, and 164 additional calls (13 to 24  
140 calls per individual) recorded from the same individuals at 5 weeks (Table 1, Figure 1).

141

142 For each call, we measured 27 spectral and temporal acoustic parameters (Table 2), which were  
143 potentially important for vocal distinctiveness. These included both temporal measures, such as  
144 call duration, related to lung capacity [35], source-related vocal features (F0) and filter-related  
145 acoustic vocal features (formants), [5, 32]. Acoustic measurements were carried out using a  
146 custom built program [36, 37] in Praat v.5.0.47 DSP Package [38].

147

#### 148 Source-related parameters

149 We extracted the F0 contour of each call using a cross-correlation method ([Sound: To Pitch (cc)  
150 command], 1 week old: time step = 0.005 s, pitch floor =300-400 Hz, pitch ceiling = 700-900 Hz; 5  
151 weeks old: time step = 0.005-0.015 s, pitch floor =200-300 Hz, pitch ceiling = 700-800 Hz). If the  
152 entire F0 contour could not be detected, calls were high-pass filtered before the analysis (cut-off  
153 frequency: 1 week old, 300 Hz; 5 weeks old, 200 Hz). For each extracted F0 contour, we  
154 measured the following vocal parameters: the frequency value of F0 at the start (F0Start) and at  
155 the end (F0End) of the call; the mean (F0Mean), minimum (F0Min) and maximum (F0Max) F0  
156 frequency values across the call; the percentage of the total call duration when F0 was maximum  
157 (TimeF0Max); and the F0 mean absolute slope (F0AbsSlope). Moreover, we calculated F0  
158 variation by measuring jitter (the mean absolute difference between frequencies of consecutive F0  
159 periods divided by the mean frequency of F0 [Jitter (local) command]) and shimmer (the mean  
160 absolute difference between the amplitudes of consecutive F0 periods divided by the mean  
161 amplitude of F0 [Shimmer (local) command]) parameters.

162

#### 163 Filter-related parameters

164 We extracted the contour of the first four formants of each call using Linear Predictive Coding  
165 analysis (LPC; [Sound: To Formant (burg) command], 1 week old: time step = 0.003 s, maximum



166 number of formants = 4–5, maximum formant = 9800–12000 Hz, window length = 0.01–0.04 s; 5  
167 weeks old: time step = 0.01–0.025 s, maximum number of formants = 4–5, maximum formant =  
168 8000–10000 Hz, window length = 0.01–0.05 s). To check if the Praat software accurately tracked  
169 the formants, the outputs of the LPC analysis were visually inspected together with the  
170 spectrograms. Spurious values and inter-segment values were deleted and we corrected for  
171 octave jumps when necessary. For each call we collected the mean (F1-4Mean) minimum (F1-  
172 4Min), and maximum (F1-4Max) values of the formants. Further, we estimated the minimum  
173 formant dispersion (DfMin) and the vocal tract length of vocalising kids (estVTL) using the methods  
174 described by Reby and McComb [36] and validated for goats by Briefer & McElligott [5]. Finally, we  
175 measured the frequency values at the upper limit of the first (Q25%), second (Q50%) and third  
176 (Q75%) quartiles of energy, using a linear amplitude spectrum applied to the whole call, and we  
177 included in the analyses the total duration of each call (Dur).

178

### 179 *2.3 Classification tasks*

180 We tested the reliability of a neural network to automate classification of goat kids contact calls  
181 according to:

- 182 1) Caller individual identity
- 183 2) Caller group membership
- 184 3) Caller age

185 For the classification tasks 1 and 2, we used 157 contact calls recorded when goat kids were 1  
186 week old (Table 1). For the classification task 3, we introduced in the analysis 164 calls recorded  
187 from the same individuals at 5 weeks of age (Table 1).

188

### 189 *2.4 Artificial Neural Network*

#### 190 Architecture

191 For this study, we used a supervised Multi-Layer Perceptron (MLP), feed-forward artificial neural  
192 network, computed in the WEKA v. 3.6.9 software package [39]. The MLP was trained with the  
193 error-back-propagation method developed by Rumelhart et al. [40]. In this MPL architecture, the

194 processing elements are arranged in the following layered structure: (a) the input nodes, (b) the  
195 hidden layers and (c) an output layer. Each neuron is connected to the other adjacent elements by  
196 axons, and the signals are transmitted forward only: from the input nodes to the output neurons  
197 through the hidden layers. The input nodes of our MLP corresponded to the acoustic parameters  
198 measured on each contact call. The output neurons corresponded, for the three classification tasks,  
199 to the identities of the callers, the group memberships of the callers, and the ages of the callers,  
200 respectively. A schematic representation of the Multi-Layer Perceptron used is presented in Figure  
201 2.

202

### 203 Training and testing

204 For each of the three classification tasks, the number of hidden units was set to the number of  
205 attributes + classes. Each model was trained for 350 epochs (learning rate 0.2; momentum 0.2).  
206 We determined these optimal values empirically, by studying the performances of different cross-  
207 validated MLP with a trial-and-error approach [26, 14]. We used 10-fold cross-validation to build  
208 robust models. For each classification task, the dataset was randomly reordered and then split into  
209 10 folds of equal size. In each iteration, one fold was used for the testing phase and the other 9  
210 folds for the training phase. In particular, we performed a *stratified* cross-validation. This means  
211 that folds were created to reflect the same class distributions in each fold as in the complete  
212 dataset. We chose this approach, because non cross-validated machine-learning algorithms are  
213 likely to overfit the training, and to lose their accuracy and ability to generalize during the test  
214 phase [41]. To estimate a reliable error of the models, we repeated 10-fold *stratified* cross-  
215 validation iterations 10 times and calculated the average predictive performance. Finally, before  
216 building the models, all the features were scaled by applying the feature normalization algorithm  
217 implemented in WEKA. This pre-processing procedure can improve ANN efficiency by keeping the  
218 connection weights from becoming too large and swamping the model during training phase [42].

219

### 220 ANN performance evaluation

221 The performances of the model were assessed (for each classification task) by calculating the  
222 following three retrieval metrics:

223 1) Accuracy (ACC). This value shows the percentage of test instances that were correctly  
224 classified by the neural network;

225 2) Kappa statistic (kappa). This value assesses whether the performance of the neural network  
226 differed from expectations based on chance alone [43, 44]. Kappa can vary between 1 (perfect  
227 classification) and 0 (classification expected by chance);

228 3) Area under the receiver operating characteristic curve (AUC). The AUC of a classifier is  
229 equivalent to the probability that it ranks a randomly chosen positive instance higher than a  
230 randomly chosen negative one. AUC values can vary between 0 and 1.

231 It is important to note that, since kappa and AUC metrics are computed in WEKA for the binary  
232 class case, we handled the multiclass classification (caller identity and group membership tasks)  
233 using the "one against all" strategy. In particular, we treated each class value in turn as the  
234 "positive" class and all others as the "negative" class.

235

### 236 Comparison of ANN with Discriminant Function Analysis (DFA)

237 In order to provide a direct comparison of the ANN accuracy with a more classical multivariate  
238 technique, we performed a discriminant function analysis in SPSS v. 19 (SPSS, Inc. 2010) for each  
239 of the three classification tasks, using the same dataset presented to the MLP. Firstly, to meet the  
240 assumption of independence between predictor variables, we performed a principal component  
241 analysis (PCA). Principal Components (PC) showing eigenvalues  $> 1$  were used to classify  
242 vocalisations with a cross-validated (leave-one-out) DFA.

243

## 244 **3. Results**

245 The Multi-Layer Perceptron succeeded in classifying most of the contact calls according to  
246 individuality, group membership and age of the goat kids. The average and standard deviations of  
247 the ACC showed limited variation within each classification task (Table 3). Average predictive  
248 performances for each classification task were, respectively,  $71.13 \pm 1.16$  % for the caller

249 individual identity ( $N = 10$  individuals and 157 calls),  $79.52 \pm 0.76$  % for the caller group  
250 membership ( $N = 10$  individuals and 157 calls) and  $91.37 \pm 0.76$  % for the caller age ( $N = 10$   
251 individuals and 321 calls). The average kappa and AUC values of the neural network models were,  
252 respectively: caller identity task =  $0.62 \pm 0.02$  and  $0.78 \pm 0.03$ , group membership =  $0.68 \pm 0.01$   
253 and  $0.92 \pm 0.01$ , caller age =  $0.85 \pm 0.02$  and  $0.98 \pm 0.01$  (Figure 3). The PCA explained 78.33 %  
254 of the total variability with 5 PCs showing eigenvalues exceeding 1. The cross-validated DFA  
255 performed using this PCA factor solution correctly classified 43.0 % of the vocal signals according  
256 to the caller individual identity, 73.50 % according to the caller group membership and 87.50 % to  
257 the caller age. To summarise, the Multi-Layer Perceptron used in this study achieved a higher  
258 accuracy than the DFA and yielded reliable predictions (none based on chance), in classifying the  
259 contact calls according to individuality, group membership and age of emitters.

#### 260 **4. Discussion**

261 We investigated whether a Multi-Layer Perceptron (MLP), feed-forward artificial neural network  
262 (ANN), could reliably classify goat kids contact calls according to the caller identity, group  
263 membership and age. To this end, we used a database of vocalisations recorded from 10 kids  
264 during the immediate postnatal period (1 week) and additional calls recorded from the same  
265 individuals at 5 weeks. For each vocalisation, we measured 27 spectral and temporal acoustic  
266 parameters, which were then presented to the neural network as input variables. The MLP showed  
267 a higher level of accuracy (ACC) compared to the results obtained with the cross-validated DFA. In  
268 particular, the DFA correctly classified 43.0 % of 1 week kid calls according to the emitter, while  
269 the MLP achieved an average ACC of 71.13 %. The MLP obtained a higher ACC also in the group  
270 membership identification of the caller (79.52 % vs 73.50 %) and suggested the presence of a  
271 social effect on the ontogeny of vocalisations in this species. Accordingly, Briefer and McElligott  
272 [30] showed that the social group influenced the energy distribution in the spectrum (energy  
273 quartiles) and the second and third formants. This probably results from changing the shape and  
274 length of the vocal tract. Finally, the MLP proved more reliable than the DFA also in classifying  
275 calls according to caller age (91.37 % vs 87.50 %), revealing the age-related changes in the vocal  
276 parameters of contact calls [5]. Overall, the MLP accuracy performances suggest that these  
277 algorithms can be used as a modern and reliable alternative to traditional statistical methods in  
278 bioacoustics.

279

280 The MLP we used showed average kappa values of  $0.62 \pm 0.02$  (caller individuality task),  $0.68 \pm$   
281  $0.01$  (group membership task),  $0.85 \pm 0.02$  (caller age task). Fleiss [44] suggested that kappa  
282 values greater than 0.75 can be considered to represent excellent agreement beyond chance,  
283 values below 0.40 indicate poor agreement beyond chance, and values between 0.40 and 0.75  
284 may be taken to represent fair to good agreement beyond chance. According to Fleiss [44], we  
285 suggest that the MLP presented in this study show reliable predictions and matching not based on  
286 chance, in each of the three classification tasks. The average area under the receiver operating  
287 characteristic curve (AUC) values were  $0.78 \pm 0.03$  (caller individuality task),  $0.92 \pm 0.01$  (group

288 membership task) and  $0.98 \pm 0.01$  (caller age task) respectively. Fawcett [45] showed that random  
289 guessing classification produces an AUC of 0.5 and suggested that realistic classifiers should have  
290 AUC higher than 0.5. Accordingly, we consider the values observed in this study as a very good  
291 discrimination in each classification task.

292

293 Overall, our results confirmed that the MLP can process a variety of spectral and temporal acoustic  
294 parameters to classify vocal signals [26]. In particular, we used temporal measures related to lung  
295 capacity (i.e. duration), source-related vocal features (F0) and filter-related acoustic vocal features  
296 (formants) to show that the MLP can be used to study vocalisations from a source-filter perspective  
297 [35]. Moreover, although ANNs have been previously used in the study of wild mammal  
298 vocalisations [27], very few reports exist for the use and potential of these techniques in farm  
299 animal research [29]. In particular, our study is the first to show the reliability of these algorithms  
300 for the classification of domestic livestock vocalisations. Developing novel tools to understand  
301 which animals are calling and to extract biological meaningful information from vocalisations has  
302 great potential for remotely monitoring domestic livestock, especially on farms with large numbers  
303 of animals. In future, the technology could be used to investigate whether the calls uttered indicate  
304 that the animals are in positive or negative states, and even to investigate their emotions [46].

305

306 We used a MLP to analyse a particular animal call type: the contact call. Contact calls are very  
307 complex signals, mostly used by birds and mammals, encoding a great deal of information about  
308 the emitter [47]. The results achieved by the MLP in grouping these calls provide evidence that  
309 ANN algorithms have the capacity to extract and categorise the biological meaningful information  
310 encoded in mammal vocalisations.

311

312 In conclusion, our results show successful examples of signal recognition by a MLP for individuality,  
313 group membership and maturation in domestic goat kids, suggesting that ANNs could be  
314 considered a reliable tool to study vocalisations of domestic livestock from a source-filter  
315 perspective. ANNs also have the potential to exhibit substantially greater predictive power than

316 traditional statistical approaches and we argue that these algorithms can be adopted to classify  
317 contact calls of many different species. Further research, comparing ANNs with other machine  
318 learning techniques would be especially valuable. We also recommend additional investigations to  
319 evaluate whether ANNs could classify contact calls in real-time and therefore be suitable to  
320 develop effective passive acoustic monitoring systems.

321

## 322 **5. Acknowledgments**

323 The authors would like to thank E. Antill, C. Booth, E. Cant, C. Charpin, K. Cho Geun-A, C.  
324 Farrington, F. Galbraith, E. Landy, M. Padilla de la Torre and M. Wang for their help with data  
325 collection. We are grateful to D. Reby for providing the custom built program in Praat. Special  
326 thanks are due to D. Stowell and D. Pessani for their helpful comments. We thank the staff of  
327 White Post Farm (<http://whitepostfarmcentre.co.uk/>) for their help and free access to their animals.  
328 L. Favaro was supported by the University of Torino through a Fiat Group Automobiles S.p.A.  
329 research grant. E. Briefer was funded by a Swiss National Science Foundation fellowship during  
330 data collection. We acknowledge the financial support of the University of London Central  
331 Research Fund for recording equipment.

332

## 333 **6. References**

- 334 [1] V. B. Deecke, V. M. Janik: Automated categorization of bioacoustic signals: Avoiding perceptual  
335 pitfalls. *J. Acoust. Soc. Am.* **119** (2006) 645-653,.
- 336 [2] M. V. Torriani, E. Vannoni, A. G. McElligott: Mother-young recognition in an ungulate hider  
337 species: a unidirectional process. *Am. Nat.* **168** (2006) 412-420.
- 338 [3] M. B. Manser, R. M. Seyfarth and D. L. Cheney: Suricate alarm calls signal predator class and  
339 urgency. *Trends Cogn. Sci.* **6** (2002) 55-57.
- 340 [4] E. Briefer, E. Vannoni, A. G. McElligott. Quality prevails over identity in the sexually selected  
341 vocalisations of an ageing mammal. *BMC Biology* **8** (2010) 35.
- 342 [5] E. Briefer, A. G. McElligott: Indicators of age, body size and sex in goat kid calls revealed using  
343 the source-filter theory. *Appl. Anim. Behav. Sci.* **133** (2011) 175-185.

- 344 [6] V. M. Janik, L. S. Sayigh, R. S. Wells: Signature whistle shape conveys identity information to  
345 bottlenose dolphins. *Proc. Natl. Acad. Sci. USA*, **103** (2006) 8293-8297.
- 346 [7] E. Vannoni, A. G. McElligott: Individual acoustic variation in fallow deer (*Dama dama*) common  
347 and harsh groans: a source-filter perspective. *Ethology* **113** (2007) 223–234.
- 348 [8] J. B. Stachowicz, E. Vannoni, B. J. Pitcher, E. F. Briefer, E. Geffen, A. G. McElligott: Acoustic  
349 divergence in the rut vocalisation of Persian and European fallow deer. *J. Zool.* **292**  
350 (2014) 1-9.
- 351 [9] L. S. Sayigh, P. L. Tyack, R. S. Wells, M. D. Scott: Signature whistles of free-ranging bottlenose  
352 dolphins *Tursiops truncatus*: Stability and mother-offspring comparisons. *Behav. Ecol.*  
353 *Sociobiol.* **26** (1990) 247-260.
- 354 [10] F. Range, J. Fischer: Vocal repertoire of Sooty Mangabeys (*Cercocebus torquatus atys*) in the  
355 Tai National Park. *Ethology* **110** (2004) 301-321.
- 356 [11] E. C. Garland, A. W. Goldizen, M. L. Rekdahl, R. Constantine, C. Garrigue, N. Daeschler  
357 Hauser, M.M. Poole, J. Robbins, M. J. Noad: Dynamic horizontal cultural transmission of  
358 humpback whale song at the ocean basin scale. *Curr. Biol.* **21** (2011) 687-691.
- 359 [12] V. M. Janik: Pitfalls in the categorization of behaviour: A comparison of dolphin whistle  
360 classification methods. *Anim. Behav.* **57** (1999) 133-143.
- 361 [13] J. J. Gros-Louis, S. E. Perry, C. Fichtel, E. Wikberg, H. Gilkenson, S. Wofsy, A. Fuentes:  
362 Vocal repertoire of *Cebus capucinus*: acoustic structure, context, and usage. *Int. J.*  
363 *Primatol.* **29** (2008) 641-670.
- 364 [14] D. Reby, S. Lek, I. Dimopoulos, J. Joachim, J. Lauga, S. Aulagnier: Artificial neural networks  
365 as a classification method in the behavioural sciences. *Behav. Process.* **40** (1997) 35-43.
- 366 [15] M. A. Acevedo, C. J. Corrada-Bravo, H. Corrada-Bravo, L. J. Villanueva-Rivera, T. M. Aide:  
367 Automated classification of bird and amphibian calls using machine learning: A  
368 comparison of methods. *Ecol. Inform.* **4** (2009) 206-214.
- 369 [16] J. D. Olden, J. J. Lawler, N. L. Poff: Machine learning methods without tears: a primer for  
370 ecologists. *Q. Rev. Biol.* **83** (2008) 171–193.



- 371 [17] T. Tirelli, L. Favaro, M. Gamba, D. Pessani: Performance comparison among multivariate and  
372 data mining approaches to model presence/absence of *Austropotamobius pallipes*  
373 complex in Piedmont (North Western Italy). *C. R. Biol.* **334** (2011) 695-704.
- 374 [18] M. Khanmohammadi, A. B. Garmarudi, K. Ghasemi: Back-propagation artificial neural network  
375 and attenuated total reflectance-Fourier transform infrared spectroscopy for diagnosis of  
376 basal cell carcinoma by blood sample analysis. *J. Chemom.* **23** (2009) 538-544.
- 377 [19] S. Ghirlanda, M. Enquist: Artificial neural networks as models of stimulus control. *Anim.*  
378 *Behav.* **56** (1998) 1383–1389.
- 379 [20] S. Huebner: Bioacoustic Classifier System Design as a Knowledge Engineering Problem. In:  
380 Computational Bioacoustics for Assessing Biodiversity. K.H. Frommolt, R. Bardeli, M.  
381 Clausen (Eds.). Federal Agency for Nature Conservation, Vilm, Germany, 2008.
- 382 [21] W. W. Au: Comparison of sonar discrimination: dolphin and an artificial neural network. *J.*  
383 *Acoust. Soc. Am.* **95** (1994) 2728-2735.
- 384 [22] E. Mercado III, A. Kuh: Classification of humpback whale vocalizations using a self-organizing  
385 neural network. *Proceedings of the International Joint Conference on Neural Networks* **2**  
386 (1998) 1584-1589.
- 387 [23] S. O. Murray, E. Mercado, H.L. Roitblat: The neural network classification of false killer whale  
388 (*Pseudorca crassidens*) vocalizations. *J. Acoust. Soc. Am.* **104** (1998) 3626–3633.
- 389 [24] M. Marcoux, M. Auger-Méthé, M. M. Humphries: Variability and context specificity of narwhal  
390 (*Monodon monoceros*) whistles and pulsed calls. *Mar. Mammal Sci.* **28** (2011) 649-665.
- 391 [25] S. Parsons, G. Jones: Acoustic identification of twelve species of echolocating bat by  
392 discriminant function analysis and artificial neural networks. *J. Exp. Biol.* **203** (2000) 2641-  
393 2656.
- 394 [26] L. Pozzi, M. Gamba, C. Giacoma: The use of Artificial Neural Networks to classify primate  
395 vocalizations: a pilot study on black lemurs. *Am. J. Primatol.* **72** (2010) 337-348.
- 396 [27] A. Mielke, K. Zuberbühler: A method for automated individual, species and call type  
397 recognition in free-ranging animals. *Anim. Behav.* **86** (2013) 475-482.

- 398 [28] J. Placer, C. N. Slobodchikoff: A fuzzy-neural system for identification of species-specific  
399 alarm calls of Gunnison's prairie dogs. *Behav. Process.* **52** (2000) 1-9.
- 400 [29] P. C. Schön, B. Puppe, G. Manteuffel: Linear prediction coding analysis and self-organizing  
401 feature map as tools to classify stress calls of domestic pigs (*Sus scrofa*). *J. Acoust. Soc.*  
402 *Am.* **110** (2001) 1425-1431.
- 403 [30] E. F. Briefer, A. G. McElligott: Social effects on vocal ontogeny in an ungulate, the goat, *Capra*  
404 *hircus*. *Anim. Behav.* **83** (2012) 991-1000.
- 405 [31] G. Fant: Acoustic theory of speech production. The Hague, Mouton, 1960.
- 406 [32] A. Taylor, D. Reby: The contribution of source-filter theory to mammal vocal communication  
407 research. *J. Zool.* **280** (2010) 221-236.
- 408 [33] E. Briefer, A. G. McElligott: Mutual mother-offspring vocal recognition in an ungulate hider  
409 species (*Capra hircus*). *Anim. Cogn.* **14** (2011) 585-598.
- 410 [34] J. Sueur, T. Aubin, C. Simonis: Equipment Review: Seewave, a free modular tool for sound  
411 analysis and synthesis. *Bioacoustics* **18** (2008) 213-226.
- 412 [35] W. T. Fitch: Production of Vocalizations in Mammals. *In: Encyclopedia of Language and*  
413 *Linguistics: 115-121*, Brown K. (Ed.), Oxford, Elsevier, 2006.
- 414 [36] D. Reby, K. McComb: Anatomical constraints generate honesty: acoustic cues to age and  
415 weight in the roars of red deer stags. *Anim. Behav.* **65** (2003) 519-530.
- 416 [37] B. D. Charlton, Z. Zhihe, R. J. Snyder: Vocal cues to identity and relatedness in giant pandas  
417 (*Ailuropoda melanoleuca*). *J. Acoust. Soc. Am.* **126** (2009) 2721-2732.
- 418 [38] P. Boersma: Praat, a system for doing phonetics by computer. *Glott International* **5** (2001) 341-  
419 345.
- 420 [39] M. Hall, E. Frank, G. Holmes, B. Pfahringer, P. Reutemann, I. H. Witten: The WEKA Data  
421 Mining Software: An Update. *SIGKDD Explorations* **11** (2009) 1-18.
- 422 [40] D. E. Rumelhart, G. E. Hinton, R. J. Williams: Learning representations by back-propagating  
423 errors. *Nature* **323** (1986) 533-536.
- 424 [41] C. Schittenkopf, G. Deco, W. Brauer: Two Strategies to Avoid Overfitting in Feedforward  
425 Networks. *Neural Networks* **10** (1997) 505-516.

- 426 [42] R. C. Eberhart, R. W. Dobbins: Neural Network PC Tools. Academic Press, New York, NY,  
427 1990.
- 428 [43] K. Titus, J. A. Mosher, B.K. Williams: Chance corrected classification for use in discriminant  
429 analysis. *Am. Midl. Nat.* **111** (1984) 1-7.
- 430 [44] J. L. Fleiss: Statistical methods for rates and proportions. Second Edition. John Wiley, New  
431 York, 1981.
- 432 [45] T. Fawcett: An introduction to ROC analysis. *Pattern Recogn. Lett.* **27** (2006) 861–874.
- 433 [46] E. Briefer: Vocal expression of emotions in mammals: mechanisms of production and  
434 evidence. *J. Zool.* **288** (2012) 1–20.
- 435 [47] N. Kondo, S. Watanabe: Contact calls: information and social function. *Jpn. Psychol. Res.* **51**  
436 (2009) 197–208.

437 **FIGURE CAPTIONS**

438

439 **Figure 1.** Spectrogram (window length: 0.009 s, time steps = 1000, frequency steps = 500,  
440 Gaussian window shape, dynamic range = 70 dB) of two contact calls recorded from the same  
441 goat kid at 1 week (left) and 5 weeks (right) of age. F0 indicates the fundamental frequency while  
442 F1-F4 indicate formant frequencies.

443 **Figure 2.** Schematic representation of the feed-forward Multi-Layer Perceptron.

444 **Figure 3.** Kappa statistic ( $\kappa$ ) and area under the ROC curve (AUC) values obtained for each  
445 classification task (I = Caller individual identity, M = Caller group membership, A = Caller age). T-  
446 bars represent 95% confidence interval.

447 **TABLES**

448 **Table 1.** Group membership and number of calls recorded for each goat kid at 1 week ( $N = 157$ )  
449 and 5 weeks of age ( $N = 164$ ).

<b>Goat kid</b>	<b>Group membership</b>	<b>1 week</b>	<b>5 weeks</b>
1	A	15	16
2	A	15	15
3	A	15	16
4	B	13	13
5	B	15	16
6	B	14	15
7	C	23	24
8	C	15	15
9	C	17	16
10	C	15	18

450

451 **Table 2.** Abbreviations and brief descriptions of the vocal parameters measured on each call.

<b>Abbreviation</b>	<b>Parameter</b>
<b>F0Start (Hz)</b>	Frequency value of F0 at the start of the call
<b>F0End (Hz)</b>	Frequency value of F0 at the end of the call
<b>F0Mean (Hz)</b>	Mean F0 frequency value across the call
<b>F0Min (Hz)</b>	Minimum F0 frequency value across the call
<b>F0Max (Hz)</b>	Maximum F0 frequency value across the call
<b>%TimeF0Max (%)</b>	Percentage of the total call duration when F0 is maximum
<b>F0AbsSlope (Hz/s)</b>	F0 mean absolute slope
<b>Jitter (%)</b>	Mean absolute difference between frequencies of consecutive F0 periods divided by the mean frequency of F0
<b>Shimmer (%)</b>	Mean absolute difference between the amplitudes of consecutive F0 periods divided by the mean amplitude of F0
<b>F1Mean (Hz)</b>	Mean frequency value of the first formant
<b>F2Mean (Hz)</b>	Mean frequency value of the second formant
<b>F3Mean (Hz)</b>	Mean frequency value of the third formant
<b>F4Mean (Hz)</b>	Mean frequency value of the fourth formant
<b>F1Min (Hz)</b>	Minimum frequency value of the first formant
<b>F2Min (Hz)</b>	Minimum frequency value of the second formant
<b>F3Min (Hz)</b>	Minimum frequency value of the third formant
<b>F4Min (Hz)</b>	Minimum frequency value of the fourth formant
<b>F1Max (Hz)</b>	Maximum frequency value of the first formant
<b>F2Max (Hz)</b>	Maximum frequency value of the second formant
<b>F3Max (Hz)</b>	Maximum frequency value of the third formant
<b>F4Max (Hz)</b>	Maximum frequency value of the fourth formant
<b>DfMin (Hz)</b>	Minimum spacing of the formants
<b>EstVTL</b>	Estimation of the vocal tract length
<b>Q25% (Hz)</b>	Frequency value at the upper limit of the first quartiles of energy
<b>Q50% (Hz)</b>	Frequency value at the upper limit of the second quartiles of energy
<b>Q75% (Hz)</b>	Frequency value at the upper limit of the third quartiles of energy
<b>Dur (s)</b>	Duration of the call

452

453 **Table 3.** Accuracy (ACC) for each classification task. Each RUN is an average among 10 different  
 454 *stratified* cross-validations.

	<b>Caller individual identity</b>	<b>Caller group membership</b>	<b>Caller age</b>
RUN	ACC	ACC	ACC
1	73.46	79.22	91.29
2	69.70	80.98	90.97
3	71.00	80.10	91.59
4	70.12	78.74	90.35
5	70.37	78.95	91.58
6	72.16	78.77	90.64
7	71.01	79.19	90.66
8	72.13	80.10	91.57
9	71.23	80.08	92.83
10	70.12	79.02	92.21
Average	71.13	79.52	91.37
St. Dev.	1.16	0.75	0.76

455