

Dogs recognise dog and human emotions

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20 Abstract: The perception of emotional expressions allows mammals to evaluate the social intentions and motivations of each other; this usually takes place within species; however in the 21 case of domestic dogs, it might be advantageous to recognise the emotions of humans as well as 22 other dogs. In this sense, the combination of visual and auditory cues to categorise others' 23 emotions facilitates the information processing and indicates high-level cognitive 24 representations. Using a cross-modal preferential looking paradigm, we presented dogs with 25 either human or dog faces with different emotional valences (happy/playful vs angry/aggressive) 26 paired with a single vocalization from the same individual with either a positive or negative 27 valence or Brownian noise. Dogs looked significantly longer at the face whose expression was 28 congruent to the valence of vocalization, for both conspecifics and heterospecifics, an ability 29 previously known only in humans. These results demonstrate that dogs can extract and integrate 30 bimodal sensory emotional information, and discriminate between positive and negative 31 emotions from both humans and dogs. 32

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41 **1.** Introduction

The recognition of emotional expressions allows animals to evaluate the social intentions and motivations of others (1). This provides crucial information about how to behave in different situations involving the establishment and maintenance of long-term relationships (2). Therefore reading the emotions of others has enormous adaptive value. The ability to recognise and respond appropriately to these cues has biological fitness benefits for both signaller and the receiver (1).

During social interactions, individuals use a range of sensory modalities, such as visual 48 and auditory cues to express emotion, with characteristic changes in both face and vocalization, 49 50 which together produce a more robust percept (3). Although facial expressions are recognised as a primary channel for the transmission of affective information in a range of species (2), the 51 perception of emotion through cross-modal sensory integration enables faster, more accurate and 52 more reliable recognition (4). Cross-modal integration of emotional cues has been observed in 53 some primate species with conspecific stimuli, such as matching a specific facial expression with 54 the corresponding vocalisation or call (5-7). However, there is currently no evidence of 55 emotional recognition of heterospecifics in non-human animals. Understanding heterospecific 56 emotions is of particular importance for animals such as domestic dogs, who live most of their 57 lives in mixed species groups and have developed mechanisms to interact with humans (8). 58 Some work has shown cross-modal capacity in dogs relating to the perception of specific 59 activities (e.g. food-guarding) (9) or individual features (e.g. body size) (10), yet it remains 60 61 unclear whether this ability extends to the processing of emotional cues, which inform individuals about the internal state of others. 62

Dogs can discriminate human facial expressions and emotional sounds (e.g. 11-18), however, there is still no evidence of multimodal emotional integration and these results relating to discrimination could be explained through simple associative processes. They do not demonstrate emotional recognition, which requires the demonstration of categorisation rather than differentiation. The integration of congruent signals across sensory inputs requires internal categorical representation (19-22) and so provides a means to demonstrate the representation of emotion.

In this study, we used a cross-modal preferential looking paradigm without 70 familiarization phase to test the hypothesis that dogs can extract and integrate emotional 71 information from visual (facial) and auditory (vocal) inputs. If dogs can cross-modally recognise 72 emotions, they should look longer at facial expressions matching the emotional valence of 73 simultaneously presented vocalizations, as demonstrated by other mammals (e.g. 5-7,21-22). 74 Due to previous findings of valence (5), side (22), sex (11,22) and species (12,23) biases in 75 perception studies, we also investigated whether these four main factors would influence the 76 dogs' response. 77

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79 2. Materials and Methods

Seventeen healthy socialised family adult dogs of various breeds were presented simultaneously with two sources of emotional information. Pairs of grey-scale gamma-corrected human or dog face images from the same individual but depicting different expressions (happy/playful vs angry/aggressive) were projected onto two screens at the same time as a sound was played (Fig. 1A). The sound was a single vocalization (dog barks or human voice in an unfamiliar language) of either positive or negative valence from the same individual or a neutral

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sound (Brownian noise). Stimuli (Fig. 1B) were taken from one female and one male of both
species. Unfamiliar individuals and languages (Brazilian Portuguese) were used to rule out the
potential influence of previous experience with model identity and human language.

Experiments took place in a quiet, dimly-lit test room and each dog received two 10-trial 89 90 sessions, separated by two weeks. Dogs stood in front of two screens and a video camera recorded their spontaneous looking behaviour. A trial consisted of the presentation of a 91 combination of the acoustic and visual stimuli and lasted five seconds (see Supplementary 92 Information for details). Each trial was considered valid for analyses when the dog looked at the 93 images for at least 2.5 seconds. The 20 trials presented different stimulus combinations: 4 face-94 pairs (2 human and 2 dog models) \times 2 vocalizations (positive and negative valence) \times 2 face 95 positions (left and right), in addition to 4 control trials (4 face-pairs with neutral auditory 96 stimulus). Therefore, each subject saw each possible combination once. 97

We calculated a congruence index=(C-I)/T where C and I represent the amount of time 98 99 the dog looked at the congruent (facial expression matching emotional vocalization, C) and incongruent faces (I), and T represents total looking time (looking left + looking right + looking 100 101 at the centre) for the given trial, to measure the dog's sensitivity to audiovisual emotional cues 102 delivered simultaneously. We analysed the congruence index across all trials using a General 103 Linear Mixed Model (GLMM) with individual dog included in the model as a random effect. 104 Only emotion valence, stimulus sex, stimulus species and presentation position (left vs right) 105 were included as the fixed effects in the final analysis because first and second order interactions were not significant. The means were compared to zero and confidence intervals were presented 106 for all the main factors in this model. A backward selection procedure was applied to identify the 107 108 significant factors. The normality assumption was verified by visually inspecting plots of

residuals with no important deviation from normality identified. To verify a possible interaction between the sex of subjects and stimuli, we used a separate GLMM taking into account these factors. We also tested whether dogs preferentially looked at a particular valence throughout trials and at a particular face in the control trials (see Supplementary Material for details of index calculation).

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115 **3. Results**

Dogs showed a clear preference for the congruent face in 67% of the trials (n=188). The 116 mean congruence index was 0.19 ± 0.03 across all test trials, and was significantly greater than 117 118 zero (t_{16} =5.53; p<0.0001), indicating dogs looked significantly longer at the face whose expression matched the valence of vocalization. Moreover, we found a consistent congruent 119 looking preference regardless of the stimulus species (dog: t_{167} =5.39, p<0.0001; human: 120 $t_{167}=2.48$, p=0.01; Fig. 2A), emotional valence (negative: $t_{167}=5.01$, p<0.0001; positive: 121 $t_{167}=2.88$, p=0.005; Fig. 2B), stimulus gender (female: $t_{167}=4.42$, p<0.0001; male: $t_{167}=3.45$, 122 p < 0.001; Fig. 2C) and stimulus position (left side: $t_{167} = 2.74$, p < 0.01; right side: $t_{167} = 5.14$, 123 p < 0.0001; Fig. 2D). When a backwards selection procedure was applied to the model with the 124 four main factors, the final model included only stimulus species. The congruence index for this 125 126 model was significantly higher for viewing dog than human faces (dog: 0.26±0.05, human: 0.12 ± 0.05 , $F_{1.170}=4.42$; p=0.04, Fig 2A), indicating that dogs demonstrated greater sensitivity 127 towards conspecific cues. In a separate model, we observed no significant interaction between 128 subject sex and stimulus sex ($F_{1,169}=1.33$, p=0.25) or main effects (subject sex: $F_{1,169}=0.17$, 129 p=0.68; subject stimulus: $F_{1,169}=0.56$, p=0.45). 130

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Dogs did not preferentially look at either of the facial expressions in control conditions when the vocalization was the neutral sound (mean: 0.04 ± 0.07 ; $t_{16}=0.56$; p=0.58). The mean preferential looking index was -0.05 ± 0.03 that was not significantly different from zero ($t_{16}=-1.6$, p=0.13), indicating that there was no difference in the proportion of viewing time between positive and negative facial expressions across trials.

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137 **4. Discussion**

The findings are the first evidence of the integration of heterospecific emotional 138 expressions in a species other than humans, and extend beyond primates the demonstration of 139 140 cross-modal integration of conspecific emotional expressions. These results show that domestic dogs can obtain dog and human emotional information from both auditory and visual inputs, and 141 integrate them into a coherent perception of emotion (21). Therefore, it is likely that dogs 142 143 possess at least the mental prototypes for emotional categorisation (positive vs negative affect) and can recognise the emotional content of these expressions. Moreover, dogs performed in this 144 145 way without any training or familiarisation with the subjects, suggesting that these emotional signals are intrinsically important. This is consistent with this ability conferring important 146 adaptive advantages (24). 147

Our study shows that dogs possess a similar ability to some non-human primates in being able to match auditory and visual emotional information (5), but also demonstrates an important advance. In our study, there was not a strict temporal correlation between the recording of visual and auditory cues (e.g. relaxed dog face with open mouth paired with playful bark), unlike the earlier research on primates (e.g. 5). Thus the relationship between the modalities was not

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temporally contiguous, reducing the likelihood of learned associations accounting for the results.

154 This suggests the existence of a robust categorical emotion representation.

Although dogs showed the ability to recognise both conspecific and heterospecific 155 emotional cues, we found that they responded significantly more strongly towards dog stimuli. 156 This could be explained by a more refined mechanism for the categorization of emotional 157 information from conspecifics, which is corroborated by the recent findings of dogs showing a 158 greater sensitivity to conspecifics' facial expressions (12) and a preference for dog over human 159 images (23). The ability to recognise emotions through visual and auditory cues may be a 160 particularly advantageous social tool in a highly social species such as dogs and might have been 161 162 exapted for the establishment and maintenance of long-term relationships with humans. It is possible that during domestication, such features could have been retained and potentially 163 selected for, albeit unconsciously. Nonetheless, the communicative value of emotion is one of 164 the core components of the process and even less-social domestic species, such as cats, express 165 affective states such as pain in their faces (25). 166

It has been a long-standing debate as to whether dogs can recognise human emotions. 167 Studies using either visual or auditory stimuli have observed that dogs can show differential 168 169 behavioural responses to single modality sensory inputs with different emotional valences 170 (e.g.14,16). For example, Müller and colleagues (13) found that dogs could selectively respond 171 to happy or angry human facial expressions; when trained with only the top (or bottom) half of 172 unfamiliar faces they generalized the learned discrimination to the other half of the face. However, these human-expression-modulated behavioural responses could be attributed solely to 173 learning of contiguous visual features. In this sense, dogs could be discriminating human facial 174 expressions without recognizing the information being transmitted. 175

176 Our subjects needed to be able to extract the emotional information from one modality and activate the corresponding emotion category template for the other modality. This indicates 177 that domestic dogs interpret faces and vocalizations using more than simple discriminative 178 processes; they obtain emotionally significant semantic content from relevant audio and visual 179 stimuli that may aid communication and social interaction. Moreover, the use of unfamiliar 180 Portuguese words controlled for potential artefacts induced by a dog's previous experience with 181 specific words. The ability to form emotional representations that include more than one sensory 182 modality suggests cognitive capacities not previously demonstrated outside of primates. Further 183 the ability of dogs to extract and integrate such information from an unfamiliar human stimulus, 184 demonstrates cognitive abilities, not known to exist beyond humans. These abilities may be 185 fundamental to a functional relationship within the mixed species social groups in which dogs 186 187 often live. Moreover, our results may indicate a more widespread distribution of the ability to spontaneously integrate multimodal cues amongst non-human mammals, which may be key to 188 understanding the evolution of social cognition. 189

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191 **References**

- Schmidt KL, Cohn JF. 2001 Human expressions as adaptations: evolutionary questions in
 facial expression research. *Am. J. Phys. Anthropol.* 33, 3-24.
- Parr LA, Winslow JT, Hopkins WD, de Waal FBM. 2000 Recognizing facial cues:
 individual discrimination by Chimpanzees (*Pan troglodytes*) and Rhesus Monkeys
 (*Macaca mulatta*). J. Comp. Psychol. 114, 47–60.
- 197 3 Campanella S, Belin P. 2007 Integrating face and voice in person perception. *Trends*.
 198 *Cogn. Sci.* 11, 535–543.

199	4	Yuval-Greenberg S, Deouell LY. 2009 The dog's meow: asymmetrical interaction in
200		cross-modal object recognition. Exp. Brain. Res. 193, 603-614.
201	5	Ghazanfar AA, Logothetis NK. 2003 Facial expressions linked to monkey calls. Nature
202		423 , 937–938.
203	6	Izumi A, Kojima S. 2004 Matching vocalizations to vocalizing faces in a chimpanzee
204		(Pan troglodytes). Anim. Cogn. 7, 179–184.
205	7	Payne C, Bachevalier J. 2013 Crossmodal integration of conspecific vocalizations in
206		Rhesus macaques. PLoS One 8(11), e81825.
207	8	Nagasawa M Mitsui S, En S, Ohtani N, Ohta M, Sakuma Y, Onaka T, Mogi K, Kikusui T
208		2015 Oxytocin-gaze positive loop and the coevolution of human-dog bonds. Science 348,
209		333–336.
210	9	Faragó T, Pongrácz P, Range F, Virányi Z, Miklósi A. 2010 'The bone is mine': affective
211		and referential aspects of dog growls. Anim. Behav. 79, 917-925.
212	10	Taylor AM, Reby D, McComb K. 2011 Cross modal perception of body size in domestic
213		dogs (Canis familiaris). PLos One 6(2), e0017069.
214	11	Nagasawa M, Murai K, Mogi K, Kikusui T. 2011 Dogs can discriminate human smiling
215		faces from blank expressions. Anim. Cogn. 14, 525–533.
216	12	Racca A, Guo K, Meints K, Mills DS. 2012 Reading faces: differential lateral gaze bias
217		in processing canine and human facial expressions in dogs and 4-year-old children. PLoS
218		<i>One</i> 7(4) , e36076.
219	13	Müller CA, Schmitt K, Barber ALA, Huber L. 2015 Dogs can discriminate emotional
220		expressions of human faces. Curr. Biol. 25, 601-605.

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221	14	Buttelmann D, Tomasello M. 2013 Can domestic dogs (Canis familiaris) use referential
222		emotional expressions to locate hidden food? Anim. Cogn. 16, 137-145.
223	15	Flom R, Gartman P. 2015 Does affective information influence domestic dogs' (Canis
224		lupus familiaris) point-following behavior? Anim. Cogn. doi:10.1007/s10071-015-0934-
225		5.
226	16	Fukuzawa M, Mills DS, Cooper JJ. 2005 The effect of human command phonetic
227		characteristics on auditory cognition in dogs (Canis familiaris). J. Comp. Psychol. 119,
228		117–120.
229	17	Custance D, Mayer J. 2012 Empathic-like responding by domestic dogs (Canis
230		familiaris) to distress in humans: an exploratory study. Anim. Cogn. 15, 851-859.
231	18	Andics A, Gácsi M, Faragó T, Kis A, Miklósi A. 2014 Voice-sensitive regions in the dog
232		and human brain are revealed by comparative fMRI. Curr. Biol. 24, 574-578.
233	19	Kondo N, Izawa E-I, Watanabe S. 2012 Crows cross-modally recognize group member
234		but not non-group members. Proc. R. Soc. B. 279, 1937-1942.
235	20	Silwa J, Duhamel J, Pascalis O, Wirth S. 2011 Spontaneous voice-face identity matching
236		by rhesus monkeys for familiar conspecifics and humans. PNAS 108, 1735–1740.
237	21	Proops L, McComb K, Reby D. 2009 Cross-modal individual recognition in domestic
238		horses (Equus caballus). PNAS 106, 947–951.
239	22	Proops L, McComb K. 2012 Cross-modal individual recognition in domestic horses
240		(Equus caballus) extends to familiar humans. Proc. R. Soc. B. 282, 3131–3138.
241	23	Somppi S, Törnqvist H, Hänninen L, Krause C, Vainio O. 2014 How dogs scan familiar
242		and inverted faces: an eye movement study. Anim. Cogn. 17, 793-803.

243	24 Guo K, Meints K, Hall C, Hall S, Mills D. 2009 Left gaze bias in humans, rhesus
244	monkeys and domestic dogs. Anim. Cogn. 12, 409-418.
245	25 Holden E, Calvo G, Collins M, Bell A, Reid J, Scot EM, Nolan AM. 2014 Evaluation of
246	facial expression in acute pain in cats. J. Small Anim. Pract. 55, 615-621.
247	
248	
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256	Figure captions
257	Fig. 1. (A) Schematic apparatus. R2: researcher, C: camera, S: screens, L: loudspeakers, P:
258	projectors, R1: researcher; (B) Example of stimuli used in the study: faces (human angry vs
259	happy, dog aggressive vs playful) and their correspondent vocalizations.
260	Fig. 2. Dogs' viewing behaviour (calculated as congruence index). (A) Species of stimulus; (B)
261	Valence of stimulus; (C) Sex of stimulus; (D) Side of stimulus presentation.

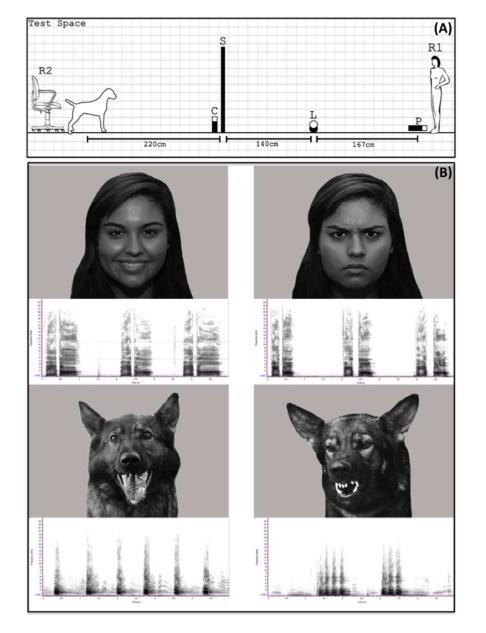


Fig. 1. (A) Schematic apparatus. R2: researcher, C: camera, S: screens, L: loudspeakers, P: projectors, R1: researcher; (B) Example of stimuli used in the study: faces (human angry vs happy, dog aggressive vs playful) and their correspondent vocalizations. 254x338mm (72 x 72 DPI)

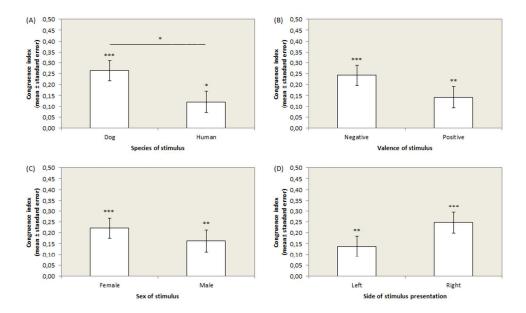


Fig. 2. Dogs' viewing behaviour (calculated as congruence index). (A) Species of stimulus; (B) Valence of stimulus; (C) Sex of stimulus; (D) Side of stimulus presentation. 343x205mm (72 x 72 DPI)

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