

HOW SELF-DOMESTICATION AND PROSOCIALITY MAY SHAPE CROSS-MODAL LANGUAGE

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An emerging literature in language evolution has highlighted the key role of self-domestication, with at least two crucial features for language evolving in other species through domestication: vocal learning in birds and the recognition of communicative intent in dogs (Thomas & Kirby, 2018). Selection for less aggressive individuals is also associated with the appearance of higher prosociality, and closely linked to increased levels of serotonin and other 'bonding' neurochemicals (Hare, 2017). Interestingly, substances that interact with such neurochemicals can heighten integration of the senses (synesthesia) in humans (Brang & Ramachandran, 2008; Luke & Terhune, 2013). Both synesthesia and the related phenomenon of shared cross-modal mappings may have played a key role in the early evolution of language (Bankieris & Simner, 2015; Cuskley & Kirby, 2013; Imai & Kita, 2014) as a way to bootstrap shared linguistic form-meaning mappings. The current paper links self-domestication and cross-modality, using a task intended to enhance participants' prosociality and measuring their sensitivity to linguistic cross-modal associations.

A total of 62 participants were recruited at the University of Edinburgh and paid £5. Half of the participants performed a social clapping task with the experimenter using a composed rhythm, aimed at stimulating prosociality: synchronising through a rhythm has been shown to increase prosociality in individuals (Kirschner & Tomasello, 2010; von Zimmermann et al., 2018). The remaining participants completed an asocial clapping task, using a random rhythm with the computer. Participants in both groups then performed a task designed to test their cross-modal sensitivity to linguistic stimuli, where pre-recorded

pseudowords had to be matched with a shape that could be manipulated in size and weight through a circular slider (see figure 1). The pseudowords were designed based on previously found strong associative reactions between consonant voicing and visual weight, and vowel openness and visual size (Schmidtke, Conrad, & Jacobs, 2014; Cuskley, 2013), and were therefore distinctive for both these features, as well as place of articulation (bilabial vs alveolar) to create more variation in the data. This created a total of 8 pseudowords: /ipi/, /ibi/, /iti/, /idi/, /apa/, /aba/, /ata/, and /ada/. Lastly, all participants were asked to answer an open question, where word count was used as a prosociality measure (Baumsteiger & Siegel, 2019).

Since size and weight were simultaneously adjusted with one slider, we refer to size for the shape produced. Pseudowords were grouped into 4 types (ordered): 1: *closed-voiceless*, 2: *closed-voiced*, 3: *open-voiceless*, 4: *open-voiced*, since participants were expected to map closed vowels and voiceless consonants with smaller sizes, where openness was expected to have a stronger effect, and no strong relationship between size and place of articulation was expected. Size in relation to condition, word type and prosociality was analyzed using linear mixed-effects models with maximum likelihood estimation (P-values calculated using the Satterthwaite's method). Sizes produced for word types followed the expected cross-modal pattern ($1 < 2 < 3 < 4$). Shape sizes were significantly larger for *open-voiceless* than for *closed-voiceless* ($b = 0.97 \pm 0.28$ SEM, $P < 0.001$) and for *open-voiced* items relative to *closed-voiceless* items ($b = 1.02 \pm 0.28$ SEM, $P < 0.001$). This also interacted with task final measures of prosociality: participants with a higher prosociality score had even larger shapes for both *open-voiceless* ($b = 0.39 \pm 0.18$ SEM, $P = 0.03$) and *open-voiced* words ($b = 0.40 \pm 0.18$ SEM, $P = 0.03$) relative to *closed-voiceless* items, suggesting a link between prosociality and cross-modal associations. However, the effect of the clapping task is less clear: social clapping only affected *open-voiceless* items, actually dampening associations relative to asocial clapping ($b = -0.36 \pm 0.15$ SEM, $P = 0.02$). Differences between the social and asocial clapping tasks may not have been sufficient, since both involved rhythmic coordination. Since the clapping task did not measurably influence cross-modal sensitivity but prosociality did, perhaps the effect relates more strongly to more stable neurochemical interactions. For example, while mu-opioids are more sensitive to prosocial manipulations (Manninen et al., 2017), serotonin is much more stable (Mitchell, 2006). Future work in this direction could take this into account.

Overall, this experiment presents an initial exploration of a role for prosociality in mechanisms facilitating intuitive meaning sharing. This could potentially open up new comparative and experimental research directions in the context of self-domestication.



Figure 1. Cross-modal interface. Left to right: Slider at initial position, at largest position and at smallest position.

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