

1 **Anecdote:** When courtship blurs: a case of male-directed sexual signaling in house
2 crickets (*Acheta domesticus*)

3 M.A. Sekhar^{1,2}

4 ¹ Environmental and Conservation Sciences Program, North Dakota State University, ND,
5 USA

6 ² Department of Biological Sciences, North Dakota State University, ND, USA

7 *Corresponding Author: M.A. Sekhar

8 Email: sekhar.ma@ndsu.edu

9 ORCID ID: 0009-0009-7807-2753

10

11 **Abstract:**

12 Same-sex sexual behaviour has been reported across diverse animal taxa and is
13 increasingly understood as a flexible outcome of sexual signaling rather than an
14 evolutionary anomaly. In insects, such behaviours are often temporary, context dependent,
15 and expressed alongside opposite-sex courtship. This pattern has led to increasing
16 emphasis on condition-dependent mating filters, defined as how selectively individuals
17 direct sexual signals toward potential mates. Here, I report a rare and, to my knowledge,
18 first observation of male-directed sexual signaling in the house cricket (*Acheta*
19 *domesticus*). In a video recorded during routine behavioural assays, an adult male
20 produced sexual acoustic signals, courtship calls, toward another male. Simultaneously,
21 this male directed aggressive signals toward other individuals within the same arena. This
22 behaviour occurred under standardized laboratory conditions with a high-quality diet,
23 ruling out nutritional stress as an explanation. Notably, the individuals involved originated
24 from highly inbred laboratory lines, suggesting that extreme homozygosity may represent
25 an overlooked source of variation influencing sexual signaling. This observation highlights
26 the value of documenting rare behaviours and suggests new avenues for exploring the role
27 of genetic context in mating filters and sexual communication.

28

29 Key words: same-sex sexual behaviour, sexual signaling, mating filters, inbreeding

30

31 Introduction

32 Same-sex sexual behaviour has been documented across diverse animal taxa, including
33 mammals, birds, reptiles, fishes and insects (Adkins-Regan, 2002; Fruth et al., 2006; Bailey
34 & Zuk, 2009; Levan, Fedina, & Lewis, 2009a; Grinham, 2026). These behaviours include
35 courtship displays, mounting, copulation attempts and long-term pair bonds, which could
36 occur in both wild and captive contexts (Bailey & Zuk, 2009). Early interpretations often
37 framed same-sex sexual behavior as an evolutionary paradox, largely because such
38 interactions do not directly result in reproduction (Young, Zaun, & VanderWerf, 2008; Bailey
39 & Zuk, 2009; Levan, Fedina, & Lewis, 2009b). However, a growing body of literatures shifted
40 this view, emphasizing that same-sex sexual behaviour can arise from flexible mating
41 systems, lenient recognition thresholds, or context dependent decision rules rather than
42 from maladaptive error alone (Bailey & Zuk, 2009; Richardson, Hoversten, & Zuk, 2024).

43 Two well-studies examples illustrate this broader perspective. In bottlenose dolphins
44 (*Tursiops spp.*), male-male sexual interactions appears to function in alliance formation
45 and social bonding, with potential downstream benefits for reproductive success (Mann,
46 2006). In birds such as Laysan albatrosses (*Phoebastria immutabili*), female-female pair
47 bonds can persist for multiple breeding seasons and may represent a conditional strategy
48 in populations with skewed sex ratios (Young, Zaun, & VanderWerf, 2008). These cases
49 shows that same-sex sexual behavior can be seen within social and ecological context that
50 shape its expression and consequences, rather than existing as an isolated studies.

51 In insects, same sex sexual behavior has been reported in multiple orders including
52 Diptera, Coleoptera, and Orthoptera (Scharf & Martin, 2013). In many cases these
53 behaviors are temporary, context dependent, and occur alongside opposite-sex courtship
54 suggesting broad acceptance thresholds for sexual signaling (Scharf & Martin, 2013). This
55 pattern has motivated recent theoretical and empirical work framing same-sex sexual
56 behavior as an outcome of flexible 'mating filters', where individuals vary in how selectively
57 their direct sexual signals towards potential mates (Richardson & Zuk, 2023).

58 Crickets provide an excellent system for studying sexual signaling, as males produce
59 distinct acoustic signals associated with mate attraction, courtship, and aggression
60 (Alexander, 1961, 1962). In some cricket species, males occasionally direct courtship
61 behavior toward other males, including the production of courtship song or mounting
62 attempts (Boutin et al., 2016). Recent experimental work has demonstrated that such
63 behaviors can be condition dependent. In Pacific field crickets (*Teleogryllus oceanicus*),
64 males reared in low quality diets showed reduced expression of same-sex sexual behavior
65 compared to males in good condition, while maintaining normal levels of courtship toward
66 females (Richardson, Hoversten, & Zuk, 2024). These findings support the idea that mating

67 filters can narrow when the energetic or opportunity costs of misdirected courtship
68 increase.

69 Importantly, this experimental work links same-sex sexual behavior to nutritional condition
70 rather than to developmental abnormality or sensory failure (Richardson, Hoversten, & Zuk,
71 2024). Under this framework, same-sex courtship emerges as an expected consequence of
72 permissive signaling rules under certain conditions, rather than as a mistake requiring a
73 special explanation (Richardson & Zuk, 2023; Richardson, Hoversten, & Zuk, 2024).
74 However, most empirical studies in crickets have focused on diet quality or short-term
75 condition as drivers of variation in same-sex sexual behavior, leaving other potential
76 sources of behavioral variation largely unexplored.

77 **A rare observation in the house cricket (*Acheta domesticus*)**

78 Here, I report the rare and, to my knowledge, previously undocumented occurrence of
79 male-directed sexual signaling in the house cricket (*Acheta domesticus*). The observation
80 of this behavior comes from a video recording collected during routine behavioral assays
81 conducted under controlled laboratory conditions.

82 In the video, one adult male clearly produces sexual signals, courtship calls, directed
83 toward another male (Figure 1a). These signals include acoustic elements typically
84 associated with mating contexts, rather than purely aggressive interactions. Notably, within
85 the same observation period, the focal male also produces aggressive signals toward
86 multiple individuals in the same arena (Figure 1b), suggesting that sexual and agonistic
87 signaling are expressed toward different social partners within a shared space. The

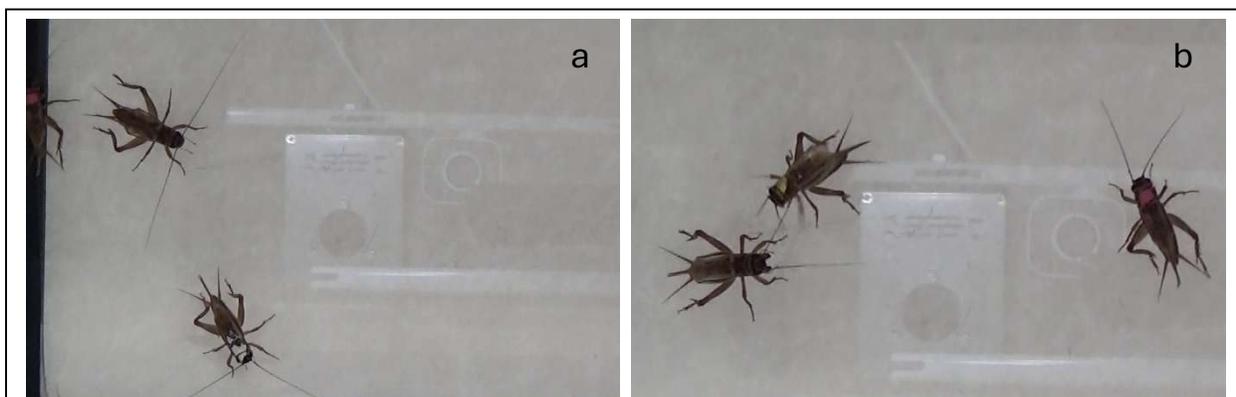


Figure 1: Context dependent signaling in *A. domesticus*. (a) Focal male (unmarked pronotum) directing mating calls toward a pink-marked male. (b) The same male displaying aggressive signals toward yellow-marked males while the pink-marked individual remains present.

88 signaling male also performed stereotypical behaviors and motor patterns that are part of
89 the Gryllidae copulatory sequence (Supplementary File 1, (Sakai & Kumashiro, 2004)).

90 The video (Supplementary File 1) and still frames (Figure 1) illustrate the focal male's
91 signaling behavior, the spatial arrangement of individuals, and the absence of females in
92 the immediate signaling context.

93 This observation is striking for several reasons. First, same-sex sexual signaling has not
94 been reported in *A. domesticus*, despite the species being one of the most intensively
95 studied crickets in laboratory research. Male house crickets are well known for producing
96 distinct signals during aggression, including mandible flaring and escalated acoustic
97 displays, while sexual signaling is typically directed toward females and occurs in specific
98 courtship contexts (Gray, 1997; Buena & Walker, 2008; Sekhar et al., 2025). Females do not
99 exhibit these signaling behaviours, and male signals are generally interpreted as sex
100 specific and context dependent (Alexander, 1962; Gray, 1997).

101 Second, the behavior described here cannot be attributed to poor nutritional condition. All
102 individuals involved were reared on a standardized, high-quality diet under consistent
103 environmental conditions. This distinguishes the present observation from previous work
104 linking same-sex sexual behavior in crickets to dietary manipulation or reduced condition.
105 This observation raises the question of whether unusual genetic or social contexts might
106 contribute to the expression of such behavior. One relevant aspect of the individuals
107 involved is their genetic background.

108 **Inbreeding as a potential contributor**

109 The individuals involved in this observation originated from laboratory lines of *A.*
110 *domesticus* generated through repeated full-sibling matings following standard
111 quantitative genetic protocols (Falconer & Mackay, 1983). These lines were derived from a
112 stock population obtained in 2021 and maintained under controlled laboratory conditions
113 with standardized, high-quality diet and husbandry. Full-sibling matings were conducted
114 for 14 consecutive generations, resulting in an expected inbreeding coefficient exceeding
115 0.94 for the individuals observed. Thus, the behavior described here occurred in a context
116 of extreme homozygosity, rather than nutritional stress or atypical rearing conditions.

117 Inbreeding is well known to influence a range of fitness-related traits, including mating
118 behavior, aggression, and signal production (Pusey & Wolf, 1996; Drayton et al., 2007;
119 Duffield et al., 2019; Townsend, Williams, & Nannas, 2024). While inbreeding is most often
120 discussed in terms of reduced performance or viability, (i.e. inbreeding depression), it can
121 also lead to altered behavioral expression by modifying neural or sensory pathways
122 underlying social interactions (Bansal et al., 2021; Townsend, Williams, & Nannas, 2024).

123 In the present case, extreme inbreeding may have altered the integration of sensory cues or
124 the balance between aggressive and sexual signaling pathways, resulting in the expression
125 of sexual signals toward an atypical social target.

126 Importantly, this interpretation remains speculative. The nature of the observation
127 excludes causal inference, and no claims are made here about adaptive function. The
128 occurrence of same-sex sexual behavior is also, apparently, rare in *A. domesticus*. The
129 behavior was observed from only one male out of 448 individuals observed over a total of
130 28 hours of recordings for this population of *A. domesticus*. However, the combination of
131 normal rearing conditions and extreme inbreeding distinguishes this case from existing
132 studies and suggests a novel axis of variation worth further investigation.

133 **Implications and future directions**

134 Although based on a single observation, this anecdote raises several broader questions
135 relevant to the study of sexual signaling and social behavior. First, it suggests that same-
136 sex sexual signaling may occur in *A. domesticus* under rare or previously overlooked
137 conditions. Second, it highlights inbreeding as a potential contributor to variation in mating
138 filters, independent of nutritional condition. Third, it underscores the importance of
139 reporting well-documented but unexpected behaviors, even when they fall outside the
140 scope of formal hypothesis-driven experiments.

141 Future work could explicitly test whether inbreeding influences mating filter breadth, signal
142 discrimination, or the coupling between aggression and courtship in crickets. Comparative
143 studies across inbred and outbred lines, combined with manipulations of social context,
144 could help determine whether the behavior reported here reflects altered sensory
145 processing, relaxed acceptance thresholds, or context-dependent signaling strategies.
146 More broadly, integrating genetic background into studies of same-sex sexual behavior may
147 reveal additional sources of variation that have been underappreciated in largely outbred
148 experimental systems.

149 By making this observation available to the scientific community, I hope to encourage
150 further documentation, replication, or contrast by other researchers working with crickets
151 or related taxa. In this sense, the anecdote serves not as a conclusion, but as an invitation
152 for reflection, comparison, and future study, aligning with the goal of protecting rare but
153 potentially informative behavioral observations from being lost to anonymity.

154 **Acknowledgement**

155 SMA thanks Prof. Ned Dochtermann for guidance and support throughout this work. Thank
156 you also to Zachary Sweep, Katy Takumi, Hieu Le, Piper Zander, and Elizabeth Frias for their

157 assistance with cricket rearing and husbandry. I also thank the Environmental and
158 Conservation Sciences Program at North Dakota State University for financial support.

159 **Conflict of interest**

160 The author declares no conflict of interest.

161

162 **Supplementary files**

163 **Video files:** <https://doi.org/10.6084/m9.figshare.31366156>

164

165 **References**

- 166 Adkins-Regan, E. (2002). Development of sexual partner preference in the zebra finch: a socially
167 monogamous, pair-bonding animal– Arch. Sex. Behav. 31: 27–33.
- 168 Alexander, R.D. (1961). Aggressiveness, territoriality, and sexual behavior in field crickets
169 (Orthoptera: Gryllidae)– Behaviour 130–223.
- 170 Alexander, R.D. (1962). Evolutionary change in cricket acoustical communication– Evolution (N
171 Y). 16: 443–467.
- 172 Bailey, N.W. & Zuk, M. (2009). Same-sex sexual behavior and evolution– Trends Ecol. Evol. 24:
173 439–446.
- 174 Bansal, R., Nagel, M., Stopkova, R., Sofer, Y., Kimchi, T., Stopka, P., Spehr, M., & Ben-Shaul, Y.
175 (2021). Do all mice smell the same? Chemosensory cues from inbred and wild mouse
176 strains elicit stereotypic sensory representations in the accessory olfactory bulb– BMC Biol.
177 19: 133.
- 178 Boutin, S.R.T., Harrison, S.J., Fitzsimmons, L.P., McAuley, E.M., & Bertram, S.M. (2016). Same-sex
179 sexual behaviour in crickets: understanding the paradox– Anim. Behav. 114: 101–110.
- 180 Buena, L.J. & Walker, S.E. (2008). Information asymmetry and aggressive behaviour in male
181 house crickets, *Acheta domesticus*– Anim. Behav. 75: 199–204.
- 182 Drayton, J.M., Hunt, J., Brooks, R., & Jennions, M.D. (2007). Sounds different: inbreeding
183 depression in sexually selected traits in the cricket *Teleogryllus commodus*– J. Evol. Biol. 20:
184 1138–1147.
- 185 Duffield, K.R., Hampton, K.J., Houslay, T.M., Hunt, J., Sadd, B.M., & Sakaluk, S.K. (2019).
186 Inbreeding alters context-dependent reproductive effort and immunity in male crickets– J.
187 Evol. Biol. 32: 731–741.
- 188 Falconer, D.S. & Mackay, T.F.C. (1983). Quantitative genetics. Longman London.
- 189 Fruth, B., Hohmann, G., Vasey, P., & Sommer, V. (2006). Social grease for females? Same-sex
190 genital contacts in wild bonobos– Homosexual behaviour in animals: An evolutionary
191 perspective 389.
- 192 Gray, D.A. (1997). Female house crickets, *Acheta domesticus*, prefer the chirps of large males–
193 Anim. Behav. 54: 1553–1562.
- 194 Grinham, L.R. (2026). Same-sex sexual behaviour in nonhuman primates– Nature Reviews
195 Biodiversity 1.

196 Levan, K.E., Fedina, T.Y., & Lewis, S.M. (2009a). Testing multiple hypotheses for the maintenance
197 of male homosexual copulatory behaviour in flour beetles– *J. Evol. Biol.* 22: 60–70.

198 Levan, K.E., Fedina, T.Y., & Lewis, S.M. (2009b). Testing multiple hypotheses for the maintenance
199 of male homosexual copulatory behaviour in flour beetles– *J. Evol. Biol.* 22: 60–70.

200 Mann, J. (2006). Establishing trust: socio-sexual behaviour and the development of–
201 Homosexual behaviour in animals: An evolutionary perspective 107.

202 Pusey, A. & Wolf, M. (1996). Inbreeding avoidance in animals– *Trends Ecol. Evol.* 11: 201–206.

203 Richardson, J., Hoversten, I.P., & Zuk, M. (2024). Male crickets in poor condition engage in less
204 same-sex sexual behavior– *Proceedings of the National Academy of Sciences* 121:
205 e2408811121.

206 Richardson, J. & Zuk, M. (2023). Rethinking same-sex sexual behaviour: male field crickets have
207 broad mating filters– *Proceedings of the Royal Society B: Biological Sciences* 290.

208 Sakai, M. & Kumashiro, M. (2004). Copulation in the cricket is performed by chain reaction–
209 *Zoolog. Sci.* 21: 705–718.

210 Scharf, I. & Martin, O.Y. (2013). Same-sex sexual behavior in insects and arachnids: prevalence,
211 causes, and consequences– *Behav. Ecol. Sociobiol.* 67: 1719–1730.

212 Sekhar, M.A., Rothamer, B.A., Gillam, E., & Dochtermann, N.A. (2025). A saga of agonistic
213 interactions in house crickets (*Acheta domesticus*): A direct and indirect effects
214 perspective– *bioRxiv* 2021–2025.

215 Townsend, A.K., Williams, K.E.G., & Nannas, N.J. (2024). Inbreeding and cognitive impairment in
216 animals– *Behavioral Ecology* 36.

217 Young, L.C., Zaun, B.J., & VanderWerf, E.A. (2008). Successful same-sex pairing in Laysan
218 albatross– *Biol. Lett.* 4: 323–325.

219