REPORT

HUMAN BEHAVIOR

Sex equality can explain the unique social structure of hunter-gatherer bands

M. Dyble,* G. D. Salali, N. Chaudhary, A. Page, D. Smith, J. Thompson, L. Vinicius, R. Mace, A. B. Migliano

The social organization of mobile hunter-gatherers has several derived features, including low within-camp relatedness and fluid meta-groups. Although these features have been proposed to have provided the selective context for the evolution of human hypercooperation and cumulative culture, how such a distinctive social system may have emerged remains unclear. We present an agent-based model suggesting that, even if all individuals in a community seek to live with as many kin as possible, within-camp relatedness is reduced if men and women have equal influence in selecting camp members. Our model closely approximates observed patterns of co-residence among Agta and Mbendjele BaYaka hunter-gatherers. Our results suggest that pair-bonding and increased sex egalitarianism in human evolutionary history may have had a transformative effect on human social organization.

ontemporary mobile hunter-gatherers cooperate extensively with unrelated individuals across multiple social and economic domains. Many communities of mobile hunter-gatherers (hereafter hunter-gatherers) share food extensively within camp and hunt, gather, and fish cooperatively (1). Alloparenting is also commonplace (2, 3). The importance of cooperative activities is reflected in many huntergatherer societies as a pervasive ethic of egalitarianism (4, 5). Like a number of non-human primate species, humans live in multimale, multifemale

University College London (UCL) Anthropology, 14 Taviton Street, London WC1H OBW, UK.

*Corresponding author. E-mail: mark.dyble.12@ucl.ac.uk

groups (6). However, we maintain enduring pair bonds, resulting in what have been described as "multifamily" groups (7). In addition, and in contrast to the bounded and territorial groups of chimpanzees (8, 9), bonobos (10) and gorillas (11), contemporary hunter-gatherers have fluid social networks where family units are relatively autonomous, with couples and their children moving often between bands (12), living with kin of either the husband or the wife. This residence pattern has been described as either "bilocal" or "multilocal" (13).

As well as being highly mobile, contemporary hunter-gatherer camps include a significant proportion of unrelated individuals (*14*) and are less closely related than groups of non-foraging smallscale societies (15). Given the inclusive fitness benefits of living with kin, why hunter-gatherers live with unrelated individuals is a puzzle, even more so if one considers that hunter-gatherers show a preference for living with siblings (13) and preferentially include kin in their campmate choices and social networks (16). Therefore, the mechanisms by which contemporary hunter-gatherers attempt to maximize co-residence and cooperation with kin, but nonetheless end up residing mostly with unrelated individuals, remain unclear.

Here, we offer a solution for this apparent paradox by demonstrating that, even where all individuals are actively assorting with kin, within-group relatedness is reduced if both sexes have influence over camp composition, as is the case among egalitarian, multilocal hunter-gatherers. We present a simulation of camp assortment where individuals attempt to reside with as many kin as possible under two conditions. In the egalitarian condition, men and women have equal influence on camp composition, whereas in the nonegalitarian condition only one sex has influence. We compared the results with previously unpublished data from two hunter-gatherer groups, the Palanan Agta (N = 4055 dyads) and Mbendjele BaYaka (5) (N = 1863 dyads), as well as one farming population, the Paranan (N = 1049 dyads). We demonstrate that low within-camp relatedness emerges naturally from men and women seeking to maximize the presence of related kin. In contrast, in societies where decision-making on coresidence rests on one sex only, as in the case of patrilocal farmers, low relatedness does not emerge. Our model offers a mechanism that reconciles individual-level preferences for kin with reduced camp-level relatedness. Assuming that extant hunter-gatherers live in social structures resembling the ones existing in past hominins, our model explains how the shift from an ancestral hierarchical, female-dispersal system, to a multilocal, egalitarian one would provide the selective context for expanded social networks, cumulative culture, and cooperation among unrelated individuals.



Fig. 1. Co-residence patterns across modeled and observed egalitarian populations. Chart area represents the proportion of all dyads across nine categories of relatedness for the egalitarian model (left), Agta (middle left), Mbendjele (middle right), Ache (bottom right), and Ju/'hoansi (top right). Ache and Ju/'hoansi data redrawn from Hill *et al.* (2011).

SCIENCE sciencemag.org

F1

F2

F3

Among the Agta, we collected data from 191 adults across 11 camps, coding a total of 4055 dvadic relationships. Among the Mbendjele, we collected data from 103 adults across nine camps, totaling 1863 dyadic relationships. Mean experienced camp size was 18.09 adults (SD = 8.62) for the Mbendjele and 21.23 adults (SD = 8.61) for the Agta. Both populations were multilocal, with husbands and wives living with similar numbers of consanguineal (genetic) kin (table S1 and fig. S1). In both groups, around 25% of dyads represented consanguineal kin, 25% were close affinal kin, and around 50% of dyads were distant affinal kin or unrelated individuals (Fig. 1 and table S2). These results are similar to those reported for the Ache and Ju/'hoansi by Hill and colleagues (14); see Fig. 1.

In contrast to the unbiased residence patterns of the Agta and Mbendjele, Paranan farmers (n = 49 adults, 1049 dyads) demonstrate a significant male bias in residence, with men living with a larger number of primary kin (n = 23, mean = 2.65, SD = 2.29) than women (n = 26, mean = 1.27, SD = 2.05; P = 0.031). Despite having a comparable group size of 21.4 adults (SD = 9.30), the Paranan live with fewer unrelated individuals than the hunter-gatherers (4.2% versus 16.7%) ($\chi^2 = 108.93$, P < 0.001) (Fig. 2).

Although it is possible that low within-camp relatedness could result from random dispersal, with households moving randomly between camps and living with related individuals only by chance, our results do not suggest that this is the case. Rather, the observed frequency of primary kin co-residence was significantly higher than would be expected if individuals assorted randomly across camps (Mbendjele, $\chi^2 = 451.62$, P < 0.001; Agta, $\chi^2 = 982.00$, P < 0.001). Thus, huntergatherer co-residence patterns are notable not only in their low-relatedness but because this low relatedness occurs despite the positive assortment of kin.

We developed a model to understand how hunter-gatherers come to co-reside with a large number of unrelated individuals at the group level, despite a preference toward living with kin at the individual level. We ran two versions of the model: one egalitarian, where both husband and wife have equal influence over where their household resides, and a non-egalitarian one, where only one sex has influence. Even at relatively small population sizes, these two conditions result in large differences in group composition. Across 100 simulations at a population size of 20, for example, there was a significantly larger proportion of unrelated dyads in the modeled egalitarian camps (12.0% \pm 8.4) compared with the non-egalitarian, single-sex dispersal camps $(0.6\% \pm 1.5)$ ($\chi^2 = 4372.36$, P < 0.001; Figs. 1 and 2). Although it is known that group relatedness decreases with increased group size (15), modeled egalitarian camps show higher proportions of unrelated individuals irrespective of camp size (Fig. 3).

The modeled co-residence patterns also mirror our observed data. The proportion of unrelated dyads in the model at a comparable group size (*n* = 20 agents) (12.0% ± 8.4) was not significantly different from the observed proportion of unrelated co-residency among the Agta (11.2%, χ^2 = 1.98, *P* = 0.016). Although the Mbendjele had significantly larger numbers of unrelated individuals in the camps (28.6%) than predicted by the model (χ^2 = 440.76, *P* < 0.001), this was in the direction consistent with our hypothesis. The observed proportion of unrelated dyads among the Paranan (4.2%) was larger than the modeled proportion (0.6% ± 1.5, χ^2 = 183.41, *P* < 0.001), but it was lower than either of the observed hunter-gatherer populations (see above) and the egalitarian model (χ^2 = 58.65, *P* < 0.001).

Our results suggest that pair-bonding alone is not sufficient to explain the low levels of relatedness seen in hunter-gatherer groups. Rather, both pair-bonding and sex equality in residential decisionmaking act together to constrain the overall relatedness of groups, leading to the co-residence of individuals unrelated through either genetic or affinal ties.

It has been proposed elsewhere that the combination of pair-bonding, cooperation among unrelated males, and increased mobility derived from male alliances could account for the low relatedness of hunter-gatherer camps (7). We argue instead that low within-camp relatedness is a consequence of sex equality in hunter-gatherer couples, with husbands and wives having an equal influence over camp composition. Given sex equality, we have shown that unrelated individuals come to



Fig. 2. Co-residence patterns across modeled and observed non-egalitarian populations. Chart area represents the proportion of all dyads across nine categories of relatedness for the non-egalitarian model (left) and Paranan (right).

Fig. 3. Modeling relatedness and equality. Results of the egalitarian model (top) and non-egalitarian model (bottom) at camp sizes between 10 and 80. From bottom to top, areas represent consanguineal (genetic) kin, affinal kin, distant affinal kin, and unrelated individuals. Exact proportions are given in tables S3 and S4.



sciencemag.org SCIENCE

co-reside even when they display a strong individual preference to live with kin, exemplified in hunter-gatherers by the frequent co-residence of brothers and sisters (14) and the higher frequencv of related individuals in campmate and gift networks (16). Therefore, our simulations provide a mechanism for the emergence of low withincamp relatedness in hunter-gatherers by solving the apparent contradiction between individuallevel preferences for living with kin and grouplevel co-residence with non-kin. Gender inequality reappeared in humans with the transition to agriculture and pastoralism (17). Once heritable resources, such as land and livestock, became important determinants of reproductive success, sex-biased inheritance and lineal systems started to arise, leading to wealth and sex inequalities (18). This predicted effect was demonstrated in our non-egalitarian model and data from Paranan agriculturalists. Our results also provide further evidence that multilocality, rather than patrilocality, is the norm among mobile hunter-gatherers.

Understanding hunter-gatherer sex egalitarianism and the shift from hierarchical male philopatry typical of chimpanzees and bonobos to a multilocal residence pattern is key to theories of human social evolution. A possible clue for the evolution of sex equality in the hominin lineage was the increase in the cost of human reproduction associated with larger brain sizes in early Homo (19). Higher offspring costs would require investment from both mothers and fathers (20), as seen among extant hunter-gatherers (3, 21). The need for biparental investment predicts increased sex equality (22), reflected in the high frequency of monogamy and the reproductive schedules of male hunter-gatherers, who typically stop reproducing early and exhibit long life spans after their last reproduction, in contrast to male farmers and pastoralists, whose reproductive spans extend well into late life (23). The recognition of affinal ties throughout our long life span has been argued to be an important step in human social evolution, and household residence may also be influenced by a tug of war between a husband and his affinal kin, who may want to live with their daughter or sister (7). The possibility of recruiting help from both maternal and paternal kin by moving camps might have been an important adaptation to meet reproductive costs in unpredictable environments, for example, by increasing the frequency of coresidence with grandmothers, who have an important provisioning role in many hunter-gatherer societies (24). Increased reproductive costs, cooperative breeding, and sex equality in residential decision-making can explain why hunter-gatherer parents live in groups containing multiple mated pairs, why hunter-gatherers recruit help both from related and unrelated individuals, and why huntergatherer camps exhibit low levels of relatedness.

Sex equality and the resulting low within-camp relatedness had many important consequences. Co-residence with unrelated individuals set the selective environment for the evolution of hypercooperation and prosociality (25). Sex equality suggests a scenario where cooperation among unrelated individuals can evolve in the absence of wealth accumulation, reproductive inequalities, and intergroup warfare (26). Couples freely moving between camps and sharing interests with kin and affines would be able to maintain cooperation without the need for more complex systems, such as cultural group selection and altruistic punishment (27).

Last, this social system may have allowed hunter-gatherers to extend their social networks, buffering environmental risk and promoting levels of information exchange required for cumulative culture (*28–31*).

REFERENCES AND NOTES

- 1. M. Gurven, Behav. Brain Sci. 27, 543-583 (2004).
- 2. P. K. Ivey, Curr. Anthropol. 41, 856-866 (2000).
- C. L. Meehan, R. Quinlan, C. D. Malcom, Am. J. Hum. Biol. 25, 42–57 (2013).
- 4. J. Woodburn, Man 17, 431-451 (1982).
- J. Lewis, in *Hunter-Gatherers of the Congo Basin: Cultures, Histories, and Biology of African Pygmies, B. S. Hewlett, Ed.* (Transaction, City, NJ, 2014), pp. 219–244.
- L. Rodseth, R. W. Wrangham, A. M. Harrigan, B. B. Smuts, *Curr. Anthropol.* 32, 221–254 (1991).
- B. Chapais, Primeval Kinship: How Pair-Bonding Gave Birth to Human Society (Harvard Univ. Press, Cambridge, MA, 2009).
 M. L. Wilson, R. W. Wrangham, Annu. Rev. Anthropol. 32,
- 363–392 (2003).
- 9. J. Mitani, D. Watts, Behaviour 138, 299–327 (2001).
- R. W. Wrangham, D. Peterson, Demonic Males: Apes and the Origins of Human Violence (Houghton Mifflin Harcourt, City, 1996).

- 11. P. Sicotte, Am. J. Primatol. 30, 21-36 (1993).
- 12. R. Kelly, J. Anthropol. Res. 39, 277-306 (1983).
- 13. F. Marlowe, Curr. Anthropol. 45, 277-284 (2004).
- 14. K. R. Hill et al., Science 331, 1286-1289 (2011).
- 15. R. S. Walker, Evol. Hum. Behav. 35, 384–388 (2014).
- C. L. Apicella, F. W. Marlowe, J. H. Fowler, N. A. Christakis, *Nature* **481**, 497–501 (2012).
- M. Martin, B. Voorhies, *Female of the Species* (Columbia Univ. Press, New York, 1975).
- 18. R. Mace, Evol. Anthropol. 22, 251-258 (2013).
- 19. L. Aiello, P. Wheeler, Curr. Anthropol. 36, 199-221 (1995).
- H. Kaplan, K. Hill, J. Lancaster, A. M. Hurtado, *Evol. Anthropol.* 9, 156–185 (2000).
- 21. K. Hill, A. M. Hurtado, *Proc. Biol. Sci.* **276**, 3863–3870 (2009).
- H. S. Kaplan, J. B. Lancaster, in Offspring: Human Fertility Behavior in Biodemographic Perspective, K. W. Wachter, R. A. Bulatao, Eds. (National Academies Press, Washington, DC, 2003), pp. 170–223.
- 23. L. Vinicius, R. Mace, A. Migliano, PLOS ONE 9, e112236 (2014).
- K. Hawkes, J. F. O'Connell, N. G. Jones, H. Alvarez, E. L. Charnov, *Proc. Natl. Acad. Sci. U.S.A.* 95, 1336–1339 (1998).
- 25. J. M. Burkart et al., Nat. Commun. 5, 4747 (2014).
- 26. D. P. Fry, P. Söderberg, Science 341, 270-273 (2013).
- 27. H. M. Lewis, L. Vinicius, J. Strods, R. Mace, A. B. Migliano, Nat. Commun. 5, 5789 (2014).
- M. Kempe, A. Mesoudi, *Evol. Hum. Behav.* 35, 285–290 (2014).
- K. R. Hill, B. M. Wood, J. Baggio, A. M. Hurtado, R. T. Boyd, PLOS ONE 9, e102806 (2014).
- R. Lee, *The! Kung San* (Cambridge Univ. Press, Cambridge, 1979).
- 31. R. Tonkinson, *The Mardu Aborigines* (Holt, Rinehard, and Winston, City, 1978).

ACKNOWLEDGMENTS

This project was funded by the Leverhulme Trust grant RP2011-R-045 to A.B.M. R.M. also received funding from European Research Council Advanced Grant AdG 249347. A.B.M. conceived the project; M.D. created the model; M.D., A.P., G.D.S., N.C., A.B.M., D.S., and J.T. collected the data; M.D. and G.D.S. wrote scripts and analyzed the data. M.D., A.B.M, L.V., and R.M. wrote the paper. We thank P. Gerbault, J. Stevenson, J. Lewis, and R. Schlaepfer for help in the field and the Human Evolutionary Ecology Group at UCL as well as three anonymous reviewers for valuable comments on the manuscript. Last, we thank our assistants in both Congo and the Philippines as well as the Agta, Paranan, and Mbendjele communities.

SUPPLEMENTARY MATERIALS

www.sciencemag.org/content/348/[issue no.]/[page]/suppl/DC1 Materials and Methods Figs. S1 and S2 Tables S1 to S9 References (*32–37*)

18 December 2014; accepted 16 April 2015 10.1126/science.aaa5139