



ELSEVIER

The genetics of human fertility

Yuri Kim and James J Lee

Heritable variation in fitness — survival and reproduction — is the fuel of evolution by natural selection. Many human societies have dramatically reduced mortality before and during the prime reproductive years, making fertility a reasonably good proxy for the whole of fitness in much of our species. For this reason, empirical knowledge regarding the genetics of fertility must be an essential part of any framework for understanding past and ongoing trends in human adaptive evolution. Here we use R.A. Fisher's analysis of human fertility as a starting point and find strong support from more recent research for his main contentions: fertility is a moderately heritable trait, where much of the genetic influences are shared with psychological characteristics.

Address

Department of Psychology, University of Minnesota Twin Cities, 75 East River Parkway, Minneapolis, MN 55455, USA

Corresponding authors: Kim, Yuri (kimx5361@umn.edu), Lee, James J (leex2293@umn.edu)

Current Opinion in Psychology 2019, **27**:41–45

This review comes from a themed issue on **Genetics**

Edited by **Brian B Boutwell** and **Michael A White**

For a complete overview see the [Issue](#) and the [Editorial](#)

Available online 13th August 2018

<https://doi.org/10.1016/j.copsyc.2018.07.011>

2352-250X/© 2018 Elsevier Ltd. All rights reserved.

All else being equal, heritable variation in fitness leads to future generations being more like the most reproductively successful members of previous generations. Ronald A. Fisher formalized this notion in the Fundamental Theorem of Natural Selection (FTNS): 'the rate of increase in fitness of any organism at any time is equal to its genetic variance in fitness at that time' [1]. A common misinterpretation of the FTNS, however, has occasionally hindered research on human fertility and quantitative traits more generally. It is worthwhile at the outset to set aside this confusion, hopefully for good.

The mistaken interpretation of the FTNS is that fitness-increasing alleles will necessarily reach fixation in the population, thus eventually resulting in a heritability of zero (i.e. no genetic variation). In particular, to the extent that fertility is a good proxy for overall fitness, it should have a near-zero heritability. However, what the FTNS

actually says is that the rate at which fitness increases, due solely to natural selection, is equal to the population's genetic variance in fitness [2,3]. It would take us too far afield to explicate why this statement of the FTNS does not in fact imply a low heritability of fitness [4,5,6**]. For our purposes it suffices to point out that Fisher's own remarks on human fertility in the very same book presenting the FTNS were not at all consistent with its interpretation as a prediction of near-zero heritability. He did not argue for a negligible heritability of fertility; in fact, he estimated the heritability of human fertility to be about 0.40 on the basis of ancestor-descendant correlations. This estimate is reasonably close to those obtained in subsequent twin studies — from 0.24 to 0.43 for women, from 0.24 to 0.28 for men [7]. The DNA-based GREML method gives an estimate of 0.10 [8], showing that not all of the heritability of fertility is due to rare variants of evolutionarily recent origin.

Fisher went beyond the estimation of heritability to provide an intriguing account of the mechanisms by which genetic variation affects fertility in humans. He emphasized the exceptionality of humans in post-forager societies, among whom differences in fertility might be more ascribable to differences in psychology ('mental and moral qualities') than to reproductive anatomy and physiology. This emphasis has proven to be quite prescient [9]. A recent genome-wide association study (GWAS) of fertility has found a near-unity genetic correlation between male and female fertility [6**]; this result is unintuitive if individual differences act chiefly through anatomy and physiology, in light of the physical differences between men and women, but becomes reasonable if the mediating traits are behavioral in nature. Indeed, this review will cover many behavioral correlates of fertility documented in recent studies. GWAS can now pinpoint the specific regions of the human genome harboring variation affecting fertility, and the first 'hits' strongly support the genetic overlap of fertility with behavior in our species; many of the polymorphic sites associated with age at first birth (an important component of overall fertility) are also strongly associated with years of education [10].

An increasing role for the psychological component of fertility since Fisher's own day may be driving a remarkable trend in modern industrialized societies: people are having children later in life and fewer children overall [11]. In many countries this trend has reached the point of subreplacement fertility — that is, a number of births insufficient to maintain the native-born population at its current size. This is certainly a very curious

phenomenon from the standpoint of evolutionary biology. A possible explanation is that a relaxation of traditional expectations with respect to family structure have enabled the greater expression of psychological differences affecting fertility preference [12–14]. It will be useful to keep this overarching hypothesis in mind as we review the particular traits appearing to mediate the heritability of human fertility.

Our review is encapsulated in [Table 1](#). In order for a correlate of fertility to have an evolutionary impact on the species, the phenotypic correlation must have some genetic basis, and in [Table 1](#) we indicate whether we have been able to find evidence in the literature for the presence of a genetic correlation. Some readers may take the term ‘fertility’ to mean the biological capacity or potential to have children (i.e. fecundity), but here we follow the demographic definition of fertility as the actual number of biological children produced over the entire lifespan.

Socioeconomic traits

The most well-established correlate of fertility may be years of education, particularly in women. The prolonging of education probably has at least some causal effect on later age at first reproduction [15–17], which inevitably depresses total fertility in turn [6**,7]. Since the mid-twentieth century, many countries have experienced a massive rise in the proportion of the population awarded a college degree, particularly among women. Women postpone having their first child in order to pursue higher education, along with other career-enhancing opportunities [18]. This is a convincing example of a social or cultural change leading to individuals, perhaps in particular those with certain genetic dispositions, altering their

reproductive behavior. Years of education and fertility do share a genetic basis. Polygenic scores for education successfully predict fertility [19], and significant genetic correlations in the expected directions have been found between years of education and various measures of fertility. The genetic correlation between years of education and age at first birth is particularly high (~ 0.7) [6**].

It has recently been shown that this negative correlation between fertility and education is changing the genetic composition of Western populations [20,21*]. For example, the average polygenic score for education is falling over time in Iceland, as a result of more highly educated individuals having fewer children.

Most studies find intelligence to be negatively correlated with fertility [22–24], and some of these studies suggest that this is not entirely because more intelligent individuals seek more education [25]. Raw IQ scores, however, have not declined over most of this time period but rather have increased [26]. As we have already said, educational credentials have proliferated as well. This apparent masking of genetic decline by environmental improvements illustrates the complexity of evolutionary change in our peculiar species. Nevertheless we think the authors of one relevant study justified in pointing out that ‘[i]t is remarkable to report changes in [the education polygenic score] across the several decades covered by this study. In evolutionary time, this is a blink of an eye. However, if this trend persists over many centuries, the impact could be profound’ [21*, p. E730].

Some studies have found higher income to be associated with decreased fertility [27]. Others have questioned the general applicability of this finding, particularly for high-

Table 1

Traits that may mediate genetic influences on fertility in humans

Correlate of fertility	Sign of correlation	Genetic correlation	References
Socioeconomic traits			
Years of education	–	Yes	[6**,15,19,20,21*]
Intelligence	–	Yes	[22–25]
Income	–	No	[15,27]
Antisocial behavior	+	Yes	[29]
Psychopathology			
Autism spectrum disorder	–	Yes	[6**,33]
Schizophrenia	–	No	[6**,33]
Bipolar disorder	+	Yes	[6**,33]
Depression	+	Yes	[19,33,34]
Reproductive behavior			
Intended fertility	+	No	[35,36]
Physical attractiveness	+	No	[37–39]
Age at first sex	–	Yes	[6**,41**]
Religion and politics			
Religiosity	+	No	[43,44,46]
Conservatism	+	No	[45,46]

‘Genetic correlation,’ whether we have been able to find a study documenting a significant genetic correlation between the two traits. Some of the relationships with fertility may be nonlinear; see the text for details.

earning men [15]. The relationship may be U-shaped, with the highest-income and lowest-income individuals having higher fertility. Further research is necessary. If it is true that the overall correlation between income and fertility is negative, the sign of this correlation may have undergone a reversal since the medieval and early modern period [28]. Antisocial behavior now shows a positive genetic correlation with fertility [29], and the sign of this correlation may also have undergone a reversal [30].

Personality and psychopathology

We will not review the literature on personality and fertility, as this has been recently done in this journal [31]. It is worth repeating the finding of this review that studies of the personality-fertility associations are sometimes inconsistent, almost certainly as a result of low statistical power. There have been few studies of whether the observed correlations have a genetic basis [6^{**},9,32], but with the growing availability of personality GWAS data there will certainly be more.

The evidence for a relationship between fertility and certain mental illnesses is more secure. Individuals diagnosed with schizophrenia and autism spectrum disorder, particularly males, show dramatically reduced fertility [33]. One oddity is that whereas individuals diagnosed with bipolar disorder or unipolar depression show baseline or slightly reduced fertility, the genetic correlations of these two disorders with fertility are both positive [6^{**},34]. A possibility worthy of further investigation is that increasing liability to bipolar disorder increases fertility for some reason, until the point where further liability produces the behavioral problems leading to a diagnosis of the disorder. Consistent with this hypothesis, siblings of individuals affected by these disorders do show somewhat elevated fertility [33].

Reproductive behavior

Intended fertility unsurprisingly affects actual fertility; those who want to have more children, tend to do so [35,36]. The proportion of children born to unmarried parents has dramatically increased in many countries [11]. Interestingly, in times and places where marriage does predict fertility, one of the mediating pathways through which physical attractiveness leads to higher fertility appears to be a greater probability of marriage [37,38]. Married women who are more attractive may have higher fertility even after conditioning on duration of marriage, which is consistent with female attractiveness providing cues to additional indicators of fecundity beyond age [39]. Physical attractiveness is heritable [40], but a genetic correlation with fertility has not been demonstrated to our knowledge.

Age at first sexual intercourse shows a very strong positive genetic correlation with age at first reproduction and hence a negative genetic correlation with overall fertility

(i.e. those who start having sex earlier tend to end up with more children) [6^{**},41^{**}]. In part, an early age at first sexual intercourse may reflect a fast life history [42], consistent with small and positive genetic correlations with age at menarche and age at voice breaking [41^{**}].

Religion and politics

People with more conservative or traditional beliefs beget more children [43–46]. These findings combine in an interesting way with the relationships between fertility and reproductive behavior described above; social conservatism is associated with earlier and more stable marriage (a positive correlate of fertility), but also with later age at first sexual intercourse and fewer sexual partners over the lifespan (negative correlates of fertility) [47]. Overall, there is a suggestion of two different reproductive strategies proving to be successful in modern Western societies: a strategy associated with socially conservative values, including a high commitment to the bearing of children within marriage; and a strategy associated with antisocial behavior, early sexual experimentation, a variety of sexual partners, low educational attainment, low commitment to marriage, haphazard pregnancies, and indifference to politics. This notion of distinct lifestyles characterized in common by relatively high fertility deserves further empirical and theoretical study.

Conclusion

R.A. Fisher was a pioneer in the study of human fertility, as in so many other fields. Using the quantitative-genetic principles that he had himself developed, he found that fertility is a moderately heritable trait. He went on to posit that much of this heritability overlaps with genetic influences on behavior. Recent research has amply confirmed these hypotheses and early findings.

Potential future directions are numerous. Table 1 suggests the importance of confirming a genetic contribution to the correlation between fertility and more traditional religious and political views. One would have liked an opportunity to ask Fisher for his thoughts about the psychological basis of this relationship. Fisher himself was a conservative, an English patriot, a professed Christian, and a father of nine, and it is natural to think that a mind so penetrating would have gleaned some insight if trained introspectively. GWAS of fertility should continue to increase the sample size, and one promising application of the resulting data might be testing the ‘Fisher-Muller hypothesis’ and its variants for the advantage of sexual reproduction itself [1,48], in a manner analogous to recent studies of mostly model organisms [49,50^{*}].

Conflict of interest statement

Nothing declared.

References and recommended reading

Papers of particular interest, published within the period of review, have been highlighted as:

- of special interest
- of outstanding interest

1. Fisher RA: *The Genetical Theory of Natural Selection: A Complete Variorum Edition*. Oxford University Press; 1999.
 2. Ewens WJ: **What is the gene trying to do?** *Brit J Philos Sci* 2011, **62**:155-176 <http://dx.doi.org/10.1093/bjps/axq005>.
 3. Lee JJ, Chow CC: **The causal meaning of Fisher's average effect.** *Genet Res* 2013, **95**:89-109 <http://dx.doi.org/10.1017/S0016672313000074>.
 4. Bürger R: *The Mathematical Theory of Selection, Recombination, and Mutation*. Wiley; 2000.
 5. Goddard ME: **Genetic architecture and the evolution of quantitative traits.** In *Proceedings of the Twentieth Conference of the Association for the Advancement of Animal Breeding and Genetics*. Edited by Villalobos NL. *Proceedings of the Twentieth Conference of the Association for the Advancement of Animal Breeding and Genetics AAABG*; 2013:122-125.
 6. Barban N, Jansen R, de Vlaming R, Vaez A, Mandemakers J, •• Trof FC, Shen X, Wilson JF, Chasman DI, Nolte IM *et al.*: **Genome-wide analysis identifies 12 loci influencing human reproductive behavior.** *Nat Genet* 2016, **48**:1462-1472 <http://dx.doi.org/10.1038/ng.3698>.
- A genome-wide association study of age at first reproduction and total number of children ever born. The sample sizes for these two respective traits are 251 151 and 343 072.
7. Mills MC, Trof FC: **The biodemography of fertility: a review and future research frontiers.** *Kölnher Z Soziol Soz* 2015, **67**:397-424 <http://dx.doi.org/10.1007/s11577-015-0319-4>.
 8. Trof FC, Stulp G, Barban N, Visscher PM, Yang J, Snieder H, Mills MC: **Human fertility, molecular genetics, and natural selection in modern societies.** *PLOS ONE* 2015, **10**:e0126821 <http://dx.doi.org/10.1371/journal.pone.0126821>.
 9. Briley DA, Trof FC, Mills MC: **What explains the heritability of completed fertility? Evidence from two large twin studies.** *Behav Genet* 2017, **47**:36-51 <http://dx.doi.org/10.1007/s10519-016-9805-3>.
 10. Lee JJ, Wedow R, Okbay A, Kong E, Maghzian O, Zacher M, Nguyen-Viet TA, Bowers P, Sidorenko J, Linne'r RK *et al.*: **Gene discovery and polygenic prediction from a 1.1-million-person GWAS of educational attainment.** *Nat Genet* 2018, **50**:1112-1121 <http://dx.doi.org/10.1038/s41588-018-0147-3>.
 11. Lesthaeghe R: **The second demographic transition: a concise overview of its development.** *Proc Natl Acad Sci U S A* 2014, **111**:18112-18115 <http://dx.doi.org/10.1073/pnas.1420441111>.
 12. Udry JR: **Biosocial models of low-fertility societies.** *Popul Dev Rev* 1996, **22**:325-336 <http://dx.doi.org/10.2307/2808017>.
 13. Barber JS: **Ideational influences on the transition to parenthood: attitudes toward childbearing and competing alternatives.** *Soc Psychol Q* 2001, **64**:101-127 <http://dx.doi.org/10.2307/3090128>.
 14. Briley DA, Harden KP, Tucker-Drob EM: **Genotype × cohort interaction on completed fertility and age at first birth.** *Behav Genet* 2015, **45**:71-83 <http://dx.doi.org/10.1007/s10519-014-9693-3>.
 15. Weeden J, Abrams MJ, Green MC, Sabini J: **Do high-status people really have fewer children? Education, income, and fertility in the contemporary U.S.** *Hum Nat* 2006, **17**:377-392 <http://dx.doi.org/10.1007/s12110-006-1001-3>.
 16. Ní Bhrolcháin M, Beaujouan É: **Fertility postponement is largely due to rising educational enrolment.** *Popul Stud* 2012, **66**:311-327 <http://dx.doi.org/10.1080/00324728.2012.697569>.
 17. Trof FC, Mandemakers JJ: **Is the association between education and fertility postponement causal? The role of family background factors.** *Demography* 2017, **54**:71-91 <http://dx.doi.org/10.1007/s13524-016-0531-5>.
 18. Balbo N, Billari FC, Mills MC: **Fertility in advanced societies: a review of research.** *Eur J Popul* 2013, **29**:1-38 <http://dx.doi.org/10.1007/s10680-012-9277-y>.
 19. Conley D, Laidley T, Belsky DW, Fletcher JM, Boardman JD, Domingue BW: **Assortative mating and differential fertility by phenotype and genotype across the 20th century.** *Proc Natl Acad Sci U S A* 2016, **113**:6647-6652 <http://dx.doi.org/10.1073/pnas.1523592113>.
 20. Beauchamp JP: **Genetic evidence for natural selection in humans in the contemporary United States.** *Proc Natl Acad Sci U S A* 2016, **113**:7774-7779 <http://dx.doi.org/10.1073/pnas.1600398113>.
 21. Kong A, Frigge ML, Thorleifsson G, Stefansson H, Young AI, Zink F, Jonsdottir GA, Okbay A, Sulem P, Masson G *et al.*: **Selection against variants in the genome associated with educational attainment.** *Proc Natl Acad Sci U S A* 2017, **114**:E727-E732 <http://dx.doi.org/10.1073/pnas.1612113114>.
- This study found that as a consequence of a negative correlation between fertility and years of education, the genetic potential for more education has declined in Iceland over the last hundred years. A strong example of how polygenic scores derived from genome-wide association studies can be used in follow-up research.
22. Van Court M, Bean FD: **Intelligence and fertility in the United States: 1912–1982.** *Intelligence* 1985, **9**:23-32 [http://dx.doi.org/10.1016/0160-2896\(85\)90004-2](http://dx.doi.org/10.1016/0160-2896(85)90004-2).
 23. Lynn R, Van Court M: **New evidence of dysgenic fertility for intelligence in the United States.** *Intelligence* 2004, **32**:193-201 <http://dx.doi.org/10.1016/j.intell.2003.09.002>.
 24. Hill WD, Marioni RE, Maghzian O, Ritchie SJ, Hagenaars SP, McIntosh AM, Gale CR, Davies G, Deary IJ: **A combined analysis of genetically correlated traits identifies 187 loci and a role for neurogenesis and myelination in intelligence.** *Mol Psychiatry* 2018 <http://dx.doi.org/10.1038/s41380-017-0001-5>. (in press).
 25. Meisenberg G: **The reproduction of intelligence.** *Intelligence* 2010, **38**:220-230 <http://dx.doi.org/10.1016/j.intell.2010.01.003>.
 26. Flynn JR: **Secular changes in intelligence.** In *The Cambridge Handbook of Intelligence*. Edited by Sternberg RJ, Kaufman SB. Cambridge University Press; 2011:647-665.
 27. Jones LE, Tertilt M: **An economic history of fertility in the United States: 1826–1960.** In *Frontiers of Family Economics*, vol 1. Edited by Rupert P, Emerald . 2008:165-230.
 28. Clark G: *A Farewell to Alms: A Brief Economic History of the World*. Princeton University Press; 2007.
 29. Tielbeek JJ, Johansson A, Polderman TJC, Rautiainen MR, Jansen P, Taylor M, Tong X, Lu Q, Burt AS, Tiemeier H *et al.*: **Genome-wide association studies of a broad spectrum of antisocial behavior.** *JAMA Psychiatry* 2017, **74**:1242-1250 <http://dx.doi.org/10.1001/jamapsychiatry.2017.3069>.
 30. Frost P, Harpending H: **Western Europe, state formation, and genetic pacification.** *Evol Psychol* 2015, **13**:230-243.
 31. Penke L, Jokela M: **The evolutionary genetics of personality revisited.** *Curr Opin Psychol* 2016, **7**:104-109 <http://dx.doi.org/10.1016/j.copsyc.2015.08.021>.
 32. Berg V, Lummaa V, Rickard IJ, Silventoinen K, Kaprio J, Jokela M: **Genetic associations between personality traits and lifetime reproductive success in humans.** *Behav Genet* 2016, **46**:742-753 <http://dx.doi.org/10.1007/s10519-016-9803-5>.
 33. Power RA, Kyaga S, Uher R, MacCabe JH, Langstrom N, Landen M, McGuffin P, Lewis CM, Lichtenstein P, Svensson AC: **Fecundity of patients with schizophrenia, autism, bipolar disorder, depression, anorexia nervosa, or substance abuse vs their unaffected siblings.** *JAMA Psychiatry* 2013, **70**:22-30 <http://dx.doi.org/10.1001/jamapsychiatry.2013.268>.
 34. Wray NR, Ripke S, Mattheisen M, Trzaskowski M, Byrne EM, Abdellaoui A, Adams MJ, Agero E, Air TM, Andlauer TMF *et al.*: **Genome-wide association analyses identify 44 risk variants and refine the genetic architecture of major depression.** *Nat*

- Genet* 2018, **50**:668-681 <http://dx.doi.org/10.1038/s41588-018-0090-3>.
35. Schoen R, Astone NM, Kim YJ, Nathanson CA, Fields JM: **Do fertility intentions affect fertility behavior?** *J Marriage Fam* 1999, **61**:790-799 <http://dx.doi.org/10.2307/353578>.
 36. Liefbroer AC: **Changes in family size intentions across young adulthood: a life-course perspective.** *Eur J Popul* 2009, **25**:363-386 <http://dx.doi.org/10.1007/s10680-008-9173-7>.
 37. Jokela M: **Physical attractiveness and reproductive success in humans: evidence from the late 20th century United States.** *Evol Hum Behav* 2009, **30**:342-350 <http://dx.doi.org/10.1016/j.evolhumbehav.2009.03.006>.
 38. Prokop P, Fedor P: **Physical attractiveness in fluences reproductive success of modern men.** *J Ethol* 2011, **29**:453-458 <http://dx.doi.org/10.1007/s10164-011-0274-0>.
 39. Püger LS, Oberzaucher E, Katina S, Holzleitner IJ, Grammer K: **Cues to fertility: perceived attractiveness and facial shape predict reproductive success.** *Evol Hum Behav* 2012, **33**:708-714 <http://dx.doi.org/10.1016/j.evolhumbehav.2012.05.00>.
 40. Mitchem DG, Purkey AM, Grebe NM, Carey G, Garver-Apgar CE, Bates TC, Arden R, Hewitt JK, Medland SE, Martin NG *et al.*: **Estimating the sex-specific effects of genes on facial attractiveness and sexual dimorphism.** *Behav Genet* 2014, **44**:270-281 <http://dx.doi.org/10.1007/s10519-013-9627-5>.
 41. Day FR, Helgason H, Chasman DI, Rose LM, Loh PR, Scott RA, Helgason A, Kong A, Masson G, Magnusson OT *et al.*: **Physical and neurobehavioral determinants of reproductive onset and success.** *Nat Genet* 2016, **48**:617-623 <http://dx.doi.org/10.1038/ng.3551>.
- A genome-wide association study of age at first sexual intercourse, using a sample size of 125 667. Earlier age at first sexual intercourse was found to be genetically correlated with more sexual partners and higher fertility
42. Stearns SC: *The Evolution of Life Histories*. Oxford University Press; 1992.
 43. Frejka T, Westoff CF: **Religion, religiousness and fertility in the US and in Europe.** *Eur J Popul* 2008, **24**:5-31 <http://dx.doi.org/10.1007/s10680-007-9121-y>.
 44. Blume M: **The reproductive benefits of religious affiliation.** In *The Biological Evolution of Religious Mind and Behavior*. Edited by Voland E, Schiefelhovel W. Springer; 2009:117-126.
 45. Kaufmann E, Goujon A, Skirbekk V: **American political affiliation, 2003–43: a cohort component projection.** *Popul Stud* 2012, **66**:53-67 <http://dx.doi.org/10.1080/00324728.2011.628047>.
 46. Friesen A: **Religion, politics, and the social capital of children.** *J Women Polit Policy* 2013, **34**:197-218 <http://dx.doi.org/10.1080/1554477X.2013.805100>.
 47. Weeden J, Kurzban R: *The Hidden Agenda of the Political Mind: How Self-Interest Shapes Our Opinions and Why We Won't Admit It*. Princeton University Press; 2014.
 48. Barton NH: **Why sex and recombination?** *Cold Spring Harb Symp Quant Biol* 2009, **74**:187-195 <http://dx.doi.org/10.1101/sqb.2009.74.030>.
 49. McDonald MJ, Rice DP, Desai MM: **Sex speeds adaptation by altering the dynamics of molecular evolution.** *Nature* 2016, **531**:233-236 <http://dx.doi.org/10.1038/nature17143>.
 50. Sohail M, Vakhrusheva OA, Sul JH, Pulit SL, Francioli LC, Genome of the Netherlands Consortium, Alzheimer's Disease Neuroimaging Initiative, van den Berg LH, Veldink JH, de Bakker PIW *et al.*: **Negative selection in humans and fruit flies involves synergistic epistasis.** *Science* 2017, **356**:539-542 <http://dx.doi.org/10.1126/science.aah5238>.
- This study found evidence from rare loss-of-function mutations in humans and fruit flies consistent with a certain class of theories explaining the advantage of sexual over clonal reproduction. Data from genome-wide association studies of human fertility might be put to a similar use.