Classification: BIOLOGICAL SCIENCES, Evolution

Title: Modelling the On-going Natural Selection of Educational Attainment in Contemporary Societies

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Abstract

I present a simulation model to examine the potential evolutionary trajectories of educational attainment (EA) under current selective pressure in western populations. With the assumption that EA is negatively correlated with fitness and has both a genetic component and a cultural component, I show that when prestige-biased transmission of the EA (i.e. people with more education are more likely to be copied) is present, the phenotype of EA is likely to keep increasing in the short term, yet the genetic component of EA may be undergoing a constant decline and eventually lead to a decrease in the phenotype.

Keywords: gene-culture coevolution, cultural evolution, prestige-biased transmission, education attainment, natural selection

Significance Statement

Contemporary humans live in very different environments from our ancestors and are subject to very different selective forces. The future evolutionary trajectory of current human populations is thus of both intellectual and pragmatic importance. The present simulation model sheds light on the short-term evolutionary dynamics of educational attainment, and demonstrates the significant contribution of systematically transmitted human culture in shaping the evolutionary paths and producing non-intuitive evolutionary outcomes.

Introduction

Today, the importance of taking evolutionary perspectives in understanding physical and physiological traits of humans is well-recognized (Albuquerque et al. 2015; Fan et al. 2016; Lieberman 2014), and there is little doubt that humans have experienced selection in relatively recent human evolutionary history (Pickrell et al. 2009). Indeed, many human physical and physiological trait variations have been celebrated as evidence of natural selection or gene-culture coevolution (Laland, Odling-Smee, and Myles 2010). Increasingly, researchers in the social sciences are paying more attention to evolutionary explanations for universality in human behavioral and cognitive traits (Buss 1999; Cosmides and Tooby 2013).

Nonetheless, to this day the majority of evolutionary studies on *homo sapiens* have focused on the past, using evolutionary theories to explain evolved physical, cognitive, and behavioral traits (Peyregne et al. 2017; Schier et al. 2014), with the notable exception of human behavioral ecology, which primarily focuses on human phenotypic plasticity in small scale societies within the fitness-maximization framework (Nettle et al. 2013; West and Burton-Chellew 2013). However, recent advances in genomic sciences and statistical techniques provide us with a better understanding of the genetic architecture of quantitative traits in humans, which, combined with observational studies on how selective pressure is operating in contemporary human populations, makes it possible to detect on-going selection and to potentially predict the future evolutionary trajectory of specific traits (Beauchamp 2016; Lee et al. 2018; Stulp et al. 2015). Of course, special attention needs to be paid when behavioral traits are under consideration, as the transmission of human culture could systematically and significantly affect the evolutionary dynamics (Boyd and Richerson 1985).

Educational attainment (EA) has long interested social scientists. It is strongly associated with fertility in a negative fashion (Ní Bhrolcháin and Beaujouan 2012; Soares 2005) and extensive research in behavioral genetics estimates significant heritability (Branigan, Mccallum, and Freese 2013; Tambs et al. 1989). Large scale Genome Wide Association Studies have identified specific loci that contribute to EA (Okbay et al. 2016; Rietveld et al. 2013), and two recent studies show that in certain western populations there is on-going negative selection against EA (Beauchamp 2016; Kong et al. 2017), where the genetic component of EA as measured by polygenic score has been declining for the past century in Iceland and the US respectively. On the other hand, the phenotype of EA, along with other cognitive measures such as the IQ score, has drastically increased in western populations (Fishlow 1966). To explore the usual evolutionary dynamics of EA, I construct an agent based simulation model to track the relative change in both genotypic value and phenotypic value of EA assuming that 1) EA is negatively associated with fitness; 2) the phenotype of EA is has both a genetic and a cultural component; 3) the genetic component is vertically transmitted; that is, from parents to offspring and the cultural component is obliquely transmitted (Feldman and Laland 1996); that is, from the parental to the offspring generation and 4) the cultural component is transmitted with a prestige-based bias, which refers to the psychological tendency of humans to copy the behaviors of the more prestigious individuals in a community (Henrich and Gil-White 2001). Alternatively, the tendency to copy from more educated people can be viewed as payoff biased transmission if education itself is considered as a desirable outcome.

The present simulation model differs from traditional evolutionary models in two ways. First, this model is not aiming at identifying evolutionarily stable strategies (Smith 1984); it is rather an attempt to combine standard evolutionary logic and cultural evolutionary theories (Henrich and McElreath 2003) to explain and potentially predict genetic and phenotypic changes of a specific trait (EA) over time. Second, this model explicitly specifies the phenotype (EA) function as having a gene-culture interaction term in addition to the additive components of gene and culture. Orthodox gene-culture coevolution theory explicitly recognizes the importance of both factors, but typically models the phenotypic contribution of gene and culture in an additive fashion (Boyd and Richerson 1985), presumably for analytical tractability. In a recent proposal, Day and Bonduriansky (2011) provide a theoretical framework to account for both genetic and non-genetic inheritance under a generalized from of the Price equation (Price 1970). However, to quantitatively track evolutionary dynamics they also assume that the phenotype is additively determined by a genetic component and a non-genetic component. As will be seen, the inclusion of a gene-culture interaction term significantly alters the evolutionary trajectories of EA.

Methods

I employ the common assumptions of asexual reproduction and non-overlapping generations in theoretical evolution (Boyd and Richerson 1985; Day and Bonduriansky 2011). Each individual is represented by a vector [g, z] where g stands for the genotypic value, and p stands for the phenotypic value (educational attainment). In all simulation runs, the population is initially seeded with individuals drawn from the uniform distribution (0,1). "Cultural fitness", or the relative likelihood that a particular individual is picked as a model, is defined as z_i^{φ}, where φ is a population level parameter which roughly represents the idea of a "cultural reproductive skew". When $\varphi > 1$, individuals with high cultural fitness have disproportionally larger probabilities to produce cultural offspring by serving as models of culturally naïve individuals, and when $\varphi = 0$, all individuals in the population are equally likely to be picked as a model.

Starting from generation 2, for any given individual i, its phenotype of interest (educational attainment) is defined to be

$$z_i = g_i\beta + z_{model} (1-\beta) + g_i z_{model} \beta_{inter} \quad (1)$$

where $g_i\beta$ represents the genetic contribution from the focal individual, $z_{model} (1-\beta)$ represents the cultural contribution from the cultural model that the focal individual picked from the previous generation based on the model's cultural fitness, and $g_i z_{model} \beta_{inter}$ represents the interaction between the genetic component and the cultural component. I constrain the β coefficients for the first two terms to sum to 1 because it reduces the total number of parameters and I am more interested in the relative contribution from gene and culture and their magnitude compared to $\beta_{inter.}$

For any given individual i, its genetic fitness w_i is defined to be

$$w_i = w_b + \gamma z_i + \varepsilon \quad (2)$$

where w_b represents the baseline fitness, γz_i represents the fitness contribution from the phenotype of interest (EA), and ε is a uniformly distributed error term. Importantly, genetic

fitness is forced to be non-negative. In cases where the numeric value of an individual's genetic fitness becomes negative during simulation runs, it is automatically set to be 0.

In the beginning of the life cycle, an individual is born with a genotypic value inherited from his parent, and will sample the whole parent generation and pick an individual as his model based on cultural fitness. His own phenotypic value is then determined by the phenotype function (1). Once all individuals in the focal generation have acquired a phenotypic value, they become the new parental generation and start produce offspring based on their genetic fitness as determined by (2). At the end of each life cycle the both the genotypic value and phenotypic value are allowed to change (mutate) with some probability in frequency and magnitude.

To further explore the evolutionary dynamics in relevant contemporary contexts, I added a "demographic shock" condition by having EA uncorrelated with fitness up to a fixed generation number and then change the correlation into negative. This is to mimic the process of demographic transition which happened in most post-industrialized societies (Coale 1984; Van De Kaa 1987) in relatively short amount of time.

Results

General result

The main result from my simulation model is that phenotypic value of EA can increase as the genotypic value of EA decreases (Figure 1, top left plot), at least for a short period of time (~23 generations for benchmark parameter settings). There are two opposing forces at work here: the negative correlation between phenotype (EA) and genetic fitness which constantly drives down genotypic value, and the prestige-biased transmission which increases population mean phenotypic value. After running a large number of simulations I identified two key parameters that contribute to this pattern of initial divergence of genotypic value and phenotypic value. First, increase in phenotype depends on the presence of variation in phenotype. In the model, the main source of phenotypic variation is supplied by random "cultural mutation" at the end of each life cycle. This result is largely intuitive: if every individual in the population has roughly the same phenotypic value, then there is less potential room for biased transmission to increase population mean. Second, the interaction term contributes to mean phenotypic value increase by magnifying the effect of biased transmission, as it places extra weight on the model's phenotypic value in constructing the focal individual's phenotype. When β_{inter} is set to be 0, the initial phenotypic increase was much milder (Figure 1, top right plot).

As genetic fitness is negatively correlated with phenotype (EA), genotypic value always monotonically decreases from the beginning until variation is exhausted in all simulation runs. The initial increase under certain parameter settings does not last indefinitely and we quickly see a subsequent drop because over time the variance in both genotypic and phenotypic values decreases due to stabilizing selection (Lande and Shannon 1996), and eventually the increase in phenotypic value due to picking a model with higher EA becomes insufficient to compensate for the overall population level decrease in phenotypic value due to natural selection acting against the phenotype.

In the vast majority of simulation runs genotypic value stabilizes after ~ 120 generations. This phenomenon is entirely due to the exhaustion of genetic variance. In the initial population, genotypic values and phenotypic values are randomly seeded, and I assume a very small mutation rate for the genetic component of EA because genetic mutation rate is vastly lower than environmental variations (Kondrashov 2003) and the simulation only looks at relatively short time period (~120 generation) in evolutionary time scales. As a result, once natural selection has used up all standing genetic variation no further genetic evolution can occur. In cases where variation in environmental contribution (cultural mutation) is large, all observed phenotypic variation would be due to environmental differences (heritability = 0); in cases where there is no variation in environmental contribution, there would be no phenotypic variation nor genotypic variation (i.e. every individual has the same genotypic value and phenotypic value).

Figure 1. The short term evolutionary trajectory of the genotypic value and phenotypic value of EA. Red line blue line represent the evolutionary trajectory of genotypic value and phenotypic value respectively. The significant diverging pattern (increase in phenotypic value but decrease in genotypic value) occurs only when both cultural mutation and gene-culture interaction term are both present.

Adding demographic shock

In the "fitness shock" condition, I model educational attainment as first contributing nothing to fitness ($\gamma=0$); then at some generations (generation 20 in the figure) EA suddenly becomes negatively associated with fitness ($\gamma=-0.5$). As shown in Figure 2, the fitness shock does not immediately drive down average phenotypic value; rather, there is a lag in the

phenotype response. Genotypic value, on the other hand, remained constant for the "pre-shock" period and started to decrease immediately after the shock.

Unlike the "non-shock" condition where the contribution of EA to fitness, γ, is always negative, the first 20 generations for the "shock" condition (γ=0) does not incur genetic evolution as genetic fitness is uncorrelated with phenotype. As expected, we observe no change in genotypic value up to generation 20. However, in the "pre-shock" period cultural evolution in the form of prestige-biased copying happens which significantly boosts the phenotype of EA. This cultural evolutionary process reduces phenotypic variation in the absence of genetic change, and consequently the diverging pattern only lasted a relatively short period of time (from generation 20 to 30), compared to the non-shock condition (from generation 0 to 20).

Figure 2. Evolutionary trajectories of EA with a shock at generation 20 (dashed line). The coefficient of effect of phenotype on fitness, γ, changes from 0 to -0.5.

Effect of prestige-biased transmission and other parameters

To systematically examine the effect of other parameters, I thus used the end point (generation 150) phenotypic and genotypic values to represent the quasi long term evolutionary outcomes within this short time window (Figure 3). As expected, most genotypic values are significantly lower at the end of the simulation run than at the beginning due to natural selection. In accordance with the logic of evolution, larger β (more genetic contribution to phenotype) and larger γ (more phenotypic contribution of EA to fitness) increases the strength of natural selection on genes, and when both β and γ are small we barely observe any decline in genotypic value (top left plot, figure 3). Most end-point variation in phenotypic values, on the other hand, is primarily due to the varying degree of prestige-biased transmission. For example, when the ego's

phenotype depends more on the phenotypic value of a cultural model ($\beta = 0.2$), the magnitude of prestige-biased transmission φ is positively correlated with the end point phenotypic values. In other words, it "helps" drive up the population mean phenotypic value. Interestingly, an increase in end-point phenotypic value due to larger cultural reproductive skew is accompanied by a smaller end-point genotypic value under certain parameter settings ($\beta = 0.2$, $\beta_{inter} = 0.1$). This is because the same conditions that leads to large phenotypic variance also leads to large genotypic variance, thus as cultural evolution drives up mean phenotypic value, genetic evolution pushes down mean genotypic value. Expectedly, these conditions are also where we observe the initial diverging patterns, except when $\varphi = 0$ (no prestige-based transmission at all).

Figure 3. End point phenotypic and genotypic values under different parameter combinations. All data are multiplied by 100 and then log transformed to accommodate for a few extremely large values. Stars represent a definitive diverging pattern of genotypic values and phenotypic values (maximum original phenotypic value >0.9). The dashed lines represent the expected starting genotypic and phenotypic values. Each pair of bards is the average of 1000 independent simulation runs.

Discussion

Simulation as a way of studying gene-culture coevolution

That human cognitive and behavioral traits are significantly influenced by human culture is no news (P. J. Richerson, Boyd, and Henrich 2010), and much effort has been put into modeling the gene-culture coevolutionary dynamics analytically (Boyd and Richerson 1985; Feldman and Cavalli-Sforza 1976). However, most models are designed for analyzing long term allele frequencies and evolutionarily stable scenarios for discrete traits. These models are not ideal for examining short term evolutionary changes, especially when models are becoming more complex incorporating more assumptions and parameters.

The evolution of educational attainment serves a case in point. Although it is not difficult to envision a situation where phenotype and genotype undergo different selective pressures, the complexity of real-world biological systems often makes it difficult for humans to evaluate the not only the effect of individual assumptions and parameter settings but also the short-term/long term evolutionary outcomes. For example, the steady increase in educational attainment over the past century can be plausibly attributed to income increase (Fishlow 1966) or "top-down" government policies aiming at increasing literacy (Lutz, Cuaresma, and Sanderson 2008), but it can be difficult to know the genetic consequences of these environmental effects or to examine how certain cultural evolutionary mechanisms—such as prestige or payoff biased transmission influence the evolutionary dynamics of EA. In these situations simulation can be of great help. The technique of simulation itself has been widely used by the cultural evolution community to both derive novel insights (McElreath et al. 2008; Smaldino 2018) and provide validation to analytic predictions (Muthukrishna et al. 2018), yet there has been very little simulation attempt to track the evolutionary change of genetic and cultural component of quantitative traits simultaneously. In the present case, the non-intuitive hill-shape phenotypic value change provides important insight for both understanding the role of specific transmission biases in shaping genetic evolution and potentially predicting the short term evolutionary change in the trait of interest (EA).

Gene and culture evolving towards opposite directions: how should we interpret the result?

There can be many environmental factors that contribute to the phenotypic change of any human trait. In particular, the technological progress and vastly improved living conditions since the industrial revolution has drastically altered many human phenotype; for example, adult human height has substantially increased throughout the world over the past century due to improved nutrition and reduction in childhood disease (Bentham et al. 2016). What is especially interesting about EA is its consistent negative association with fertility. In general, more educated women delay the onset of childbearing and have fewer children overall compared to less educated women (Martín-García and Baizán 2006; Ní Bhrolcháin and Beaujouan 2012). This pattern is very robust in both developed (Brewster and Rindfuss 2002) and developing countries (Canning, Raja, and Yazbeck 2015), and various theories have been proposed to explicate the intrinsic, potentially causal relationship between education level and fertility (Blossfeld and Huinink 2002; Edin and Kefalas 2005).

This negative association between EA and fertility, which is a reasonable proxy for fitness, has important evolutionary consequences, as both theoretical predictions and empirical evidence suggest that the genetic component of EA is undergoing a temporal decline (Kong et al. 2017). The phenotype of EA, on the other hand, may experience a very different type of selective

pressure. A large literature in sociology shows that educational attainment and socio-economic status are associated (Ohlendorf, Sewell, and Haller 1970; Thomson 2018), and cultural evolutionary theory predicts that humans readily copy the behaviors of the those who are perceived as more prestigious or successful (Henrich and Gil-White 2001; Henrich and McElreath 2003). Although educational attainment cannot be "copied" in a literal sense, individuals who pick models with high EA are likely to be more motivated in learning and committed to their academic studies, and as a result become more likely to obtain higher EA themselves. Thus, genetic fitness and cultural fitness of the same trait (EA) invites selection in opposite directions, and what is shown in the simulation model is that biased transmission alone can lead to a short term increase in phenotype.

In reality, the massive increase in EA and over the past century is almost certainly a combination of various social, economic, and cultural factors, and the substantial increase in the phenotype of EA indicates that whatever the mechanism is, the decline of genetic component of EA has been vastly overcompensated by enhanced environmental inputs. However, the simulation results reveal a crucial insight: short term change in phenotypic value does not guarantee long-term phenotypic change in the same direction. A natural question to ask is whether there is some ceiling effect of environmental input: that is, whether the environmental factors can drive up the phenotype of EA indefinitely. It seems reasonable to assume some upper limit of environmental influences, which means at some point genetics will become the limiting factor in determining the phenotype of EA, which may eventually decline as a result of natural selection.

Even in the short term, the decline of the genetic component of EA may have non-trivial consequences. Although people can reasonably question whether educational attainment is the trait under selection (Woodley 2016), it is not crucially important what the precise trait is. What matters in this evolutionary process is that genetic variants associated with EA are being selective against. We know that these genetic variants are primarily involved in brain development processes and neuron-to-neuron communication (Lee et al. 2018), which accords with previous research showing that the same set of SNPs is associated with both educational attainment and cognitive function (Rietveld et al. 2013). Thus, it seems that at least part of the genetic component of EA comes in the form cognitive ability, which is important for not only EA but also other aspects of social outcomes such as income (Ceci and Williams 1997) and job performance (Hunter 1986).

The puzzle of contemporary humans: are they fitness maximizer?

A long-standing question in evolutionary biology is whether organisms behave as if they are maximizing fitness (Lehmann, Alger, and Weibull 2015). Humans, as an evolved primate, must be subject to the logic of evolution via natural selection (Cosmides and Tooby 2013); yet on the other hand, due to the presence of a second, cultural inheritance system (Boyd and Richerson 1985), seem to require a different type of analysis. Richerson and Boyd (2005) offer a plausible account of how cultural transmission could result in maladaptions –when genetic fitness is not perfectly aligned with cultural fitness for any particular trait. More broadly, our capacity for culture evolved so that we could acquire information socially, which conferred significant fitness advantage in the evolutionary past; yet the very same psychological

mechanism allows for maladaptations at the same time. In the modern environment, copying the most prestige individuals may not be the most adaptive strategy (for an extreme case, see De Leo and Heller 2008 on the social transmission of suicide).

However, this view does not go unchallenged by evolutionary biologists. El Mouden et al. (2014) presents a narrative of why genetic fitness should always align with cultural fitness by arguing that "genetic selection will shape psychological mechanisms to avoid cultural traits that bear a genetic fitness cost". This line of thinking assumes an infinite strategy space, in particular the possible psychological mechanisms that are capable of detecting genetic fitness consequences in novel environments. Even if we ignore all constraints on the potential evolution of human cognition, it remains the case that novel mutations reaching evolutionary equilibrium state typically requires some significant amount of time (Lynch 2016).

What is notable about contemporary humans is that the rate of environmental change can be very rapid. In post-industrialized societies, both the ecological environment and the cultural environment (norms, values) are vastly different from not very long ago. Substantially reduced infant mortality (Corsini and Viazzo 1997), dissociation of sexual activity from reproduction (Tone 2001), and the reversed association between status and fertility (Haines 1989) happened only in the past few hundred years, and it is unrealistic for genetic evolution to generate new psychological mechanisms to keep up with the cultural change. Thus, to understand the short-term evolutionary dynamics it would be better to focus on what is known about the selective pressure and the heritability of the target of selection.

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