Evolutionary genomic patterns of recent natural selection on loci associated with sexually differentiated human body size and shape phenotypes

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Abstract

Levels of sex differences for human body size and shape phenotypes are hypothesized to have been reduced following the agricultural transition, as an evolutionary response to changing socioeconomic behaviors including relatively more equal divisions of labor and the adoption of new technologies. Here we apply a multi-step approach to study the recent evolutionary history of genetic variants associated with five sexually differentiated human phenotypes: height, body mass, hip circumference, body fat percentage, and waist circumference. We first analyzed genome-wide association study (GWAS) results for UK Biobank individuals (~197,000 females and ~167,000 males) in order to identify a total of 119,023 single nucleotide polymorphisms (SNPs) significantly associated with at least one of the studied phenotypes in either females or males or in both sexes ($P<5x10^{-8}$). From the set of phenotype-associated loci we then identified 3,016 SNPs (2.5%) with significant differences in the strength of association between the female and male-specific GWAS results at a low false-discovery rate (FDR<0.001; n=148 SNPs after pruning for linkage disequilibrium). Genes with known roles in sexual differentiation are significantly enriched for co-localization with one or more of these SNPs versus SNPs associated with the phenotypes generally but not with sex differences (2.83-fold enrichment; permutation test; P=0.0045). We then used the singleton density score statistic, which quantifies recent (within the last ~3,000 years) changes in the frequencies of alleles underlying polygenic traits, to identify signatures of recent positive selection on genetic variants associated with sex differences. Specifically, we observed significant increases in the frequencies of alleles associated with greater body fat percentage in females relative to SNPs associated with body fat percentage generally (permutation test; P=0.0038; FDR=0.038) and otherwise no evidence of positive selection for sex difference-associated alleles for any other trait. Given that females have higher average body fat percentages than males in historic and contemporary populations, our results contradict and otherwise fail to support expectations based on longstanding hypotheses concerning the direction of recent evolutionary change in human body size and shape sex differences.

Significance statement

There is uncertainty regarding the recent evolutionary history of human sex differences for quantitative body size and shape phenotypes. In this study we identified thousands of genetic loci that differentially impact body size and shape trait variation between females and males using a large sample of UK Biobank individuals. After confirming the biological plausibility of these loci, we used a population genomic approach to study the recent (over the past 3,000 years) evolutionary histories of these loci in this population. We observed significant increases in the frequencies of alleles associated with greater body fat percentage in females. This result is contradictory to longstanding hypotheses that sex differences have adaptively decreased following subsistence transitions from hunting and gathering to agriculture.

Main Text

Introduction

In many vertebrate species, it is not uncommon for morphological phenotypes to have average size and shape differences between females and males (1). Such traits are often generally referred to as 'sexually dimorphic.' However, that term is technically specific to non-overlapping traits; for example, exaggerated ornamentation in male guppies, peacocks, and mandrills (2–5). Traits with average phenotype values that differ by sex but with overlapping trait distributions – such as human height and body fat percentage – are better described as sexually differentiated traits.

Some sexually differentiated traits are believed by many researchers to be the result of sexual selection. In species with high inter-male competition for mates, larger males may have a competitive advantage that results in increased fitness (6). Perhaps as a result, the magnitude of sexually differentiated phenotypes are often greater in polygynous species with high competition (e.g., gorillas) and lower in monogamous species (e.g., gibbons) (6–8). Large ranges in the degree of body size and shape sexually differentiated traits are repeatedly observed among closely related groups of species, suggesting the potential for relatively rapid evolution (6).

In humans, females and males exhibit significant but relatively subtle differences in many anthropometric phenotypes (9). For example, European and African males are an average of approximately 9% taller in height and 15% heavier in body mass than females from the same populations (10, 11). Humans also exhibit sexually differentiated biometric and disease phenotypes, especially those related to immune function (12, 13).

There is little consensus regarding the evolutionary history of sexually differentiated traits in our species (11, 14, 15). Current levels of human sex differences may partially reflect an evolutionary history of complex interactions between our biology and culture, but the timing and direction of any adaptive changes in these patterns are debated. For example, it has been hypothesized that the degree of sexual differentiation for body size phenotypes in Europe likely decreased following recent (within the past ~10,000 years) shifts from a foraging-based subsistence strategy to one relying primarily on food production, in response to less pronounced divisions of labor and mobility (16). However, other scholars suggest that sexually differentiated phenotypes would respond to selection at too slow an evolutionary rate to respond to recent environmental and cultural changes; thus, any such recent changes are more likely to reflect genetic drift and/or non-genetic responses to environmental changes rather than natural selection (17).

In this study, we combined genomic and evolutionary analyses to quantify how recent changes in lifestyle and culture might be affecting the underlying genetic basis of human sexually differentiated traits. First, we applied a genome wide association study (GWAS)-based approach to identify genetic variants associated with sexually differentiated phenotypes based on data from the UK Biobank study (18). With the recent availability of a greatly increased participant sample size, this analysis represents a powerful extension of the several previous GWAS-based

approaches that have studied the genetic architecture of human sexually differentiated anthropometric traits (19). We then used the Singleton Density Score (SDS), a statistic that identifies signatures of polygenic adaptation that acted within the last ~3,000 years (20), a period following the transition to agriculture in present-day United Kingdom (21).

Results

We analyzed genome-wide association study (GWAS) summary statistics for the following five sexually differentiated anthropometric phenotypes that were produced by the Neale Lab (22) using data from the UK Biobank (18): height, body mass, hip circumference, body fat percentage, and waist circumference. We chose these traits given their relevance to the motivating evolutionary hypothesis and because they have been extensively studied from anthropological and/or genomics perspectives (9, 23–25). We did not include body mass index following concerns about this metric (26).

Identification of phenotype-associated SNPs

We analyzed summary statistics from GWAS that were performed separately in ~194,000 females and ~167,000 males of white British genetic ancestry on ~13.8 million autosomal SNPs (22). SNPs with minor allele frequencies < 0.05 and low imputation quality were filtered out. We further restricted our analysis to SNPs that 1) passed these filters in both females and males and 2) had SDS values available (20), given our motivation to conduct downstream evolutionary analyses. These filtering steps resulted in ~4.4 million genome-wide SNPs for each sex-specific GWAS. For each phenotype, we identified significantly phenotype-associated SNPs present in females, males, or both using the genome-wide significance threshold of $P = 5x10^{-8}$ commonly applied in UK Biobank studies (27–29). We identified the following total (not yet pruned for linkage disequilibrium) numbers of phenotype-associated SNPs significant in either females, males, or both: 67,738 for height, 15,669 for body mass, 12,580 for hip circumference, 10,538 for body fat percent, and 7,765 for waist circumference (**Figure 1A-B**; Supplementary Table 1).

SNPs disproportionately associated with female or male trait variation

From the sets of phenotype-associated SNPs that were significant in females, males, or both for each phenotype (SexDiff-associated SNPs), we identified those SNPs with significant differences in the statistical strengths of association for the female vs. male-specific GWAS results (t-SexDiff) (19). To account for multiple testing, we performed this analysis with four different false-discovery rate (FDR) cutoffs for each phenotype: 0.05, 0.01, 0.005, and 0.001.

At the most stringent FDR cutoff of 0.001, we identified the following number of SexDiff-associated SNPs: 677 for height, 541 for body mass, 808 for hip circumference, 439 for body fat percentage, and 551 for waist circumference (**Figure 1C-D**; Supplementary Table 1). In addition to identifying SexDiff-associated SNPs, for each phenotype we calculated the ratio of SexDiff-associated SNPs at the FDR threshold of 0.001 to the number of phenotype-associated SNPs, which ranged from 0.0010 for height to 0.0718 for waist circumference.

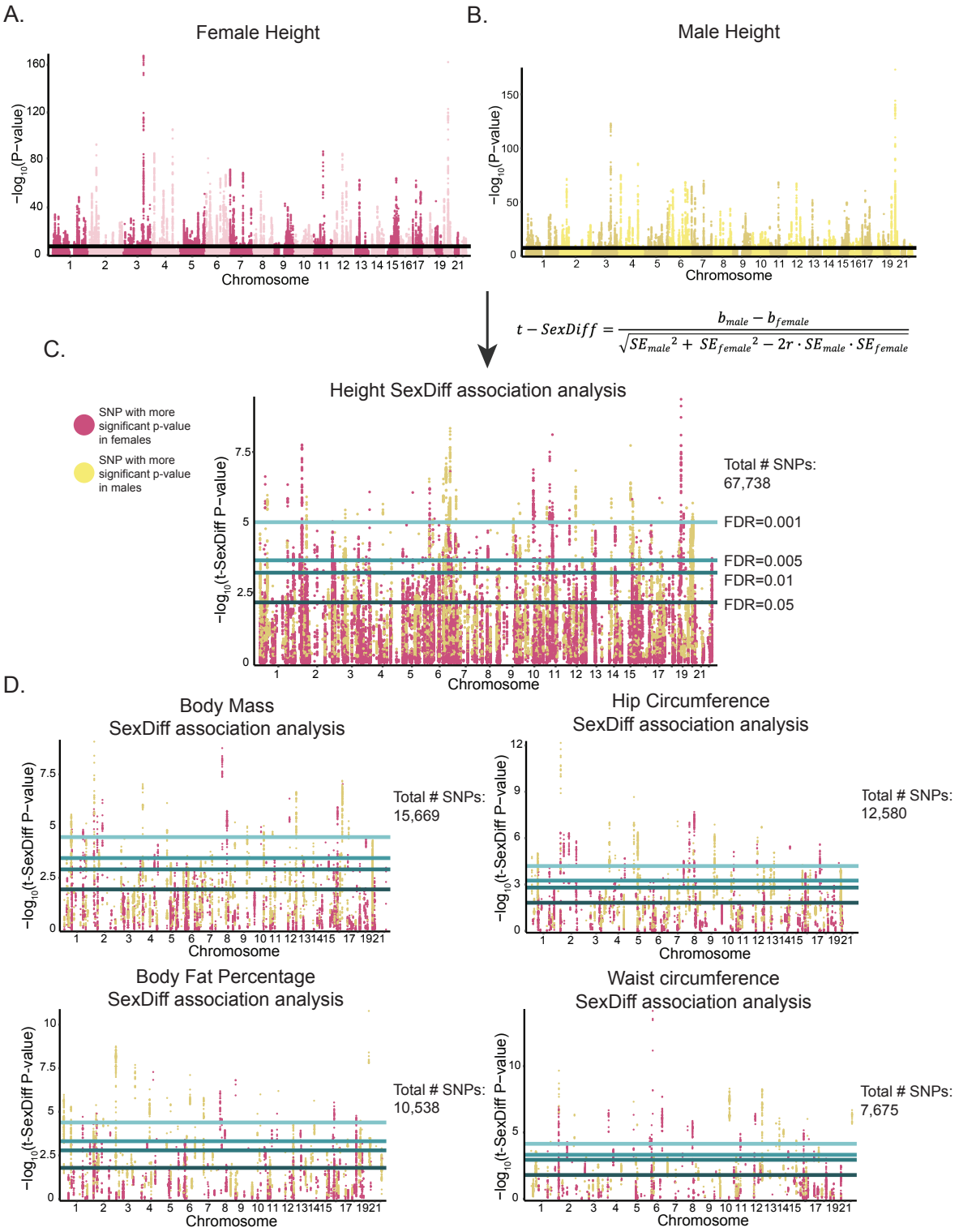

Figure 1. SexDiff-associated SNPs for five anthropometric phenotypes. (A) Manhattan plot depicting -$\log_{10}$ P-values for the association of each genome-wide SNP with female height. The black line corresponds to the genome-wide significance threshold of $P=5x10^{-8}$. (B) Manhattan plot for SNP associations with male height. (C) For each of the 67,738 SNPs significantly associated female and/or male height, we used the equation shown to test whether the SNP was disproportionately associated with height between the sexes (height SexDiff-associated SNPs). The plot depicts -$\log_{10}$ P-values for the t-SexDiff statistic. Green bars correspond to four different FDR cutoffs. (D) SexDiff association analyses for significant phenotype-associated SNPs (number of SNPs included in each analysis is shown to the right of each plot) for four additional anthropometric traits: body mass, hip circumference, body fat percentage, and waist circumference.

Genes with known roles in sexual differentiation are enriched for SexDiff-associated SNPs

Prior to conducting evolutionary analyses on the anthropometric trait SexDiff-associated SNPs, we assessed their biological plausibility by testing whether these SNPs are significantly more likely to be located within or nearby (+/- 10,000 base pairs) genes previously known to be involved in sexual differentiation (Gene Ontology term GO:0007548) compared to SNPs that are significantly associated with the phenotype but not significantly associated with sex differences (30, 31). Although not all SexDiff-associated SNPs are expected to be located nearby genes already known to be involved in sexual differentiation, we would expect at least some enrichment if we are identifying true SexDiff associations with our approach.

For each t-SexDiff FDR threshold, we determined the number of unique sexual differentiation (GO:0007548) genes with one or more co-localized SexDiff-associated SNPs. We similarly identified the number of all genes (of those in the GO database) that were co-localized with at least one SexDiff-associated SNP and calculated the proportion of the number of GO:0007548 genes to the total number of genes. We conducted the same analysis for the set of SNPs associated with the phenotypes in general but *not* associated with sexual differentiation. Finally, we estimated the GO:0007548 enrichment ratio for SexDiff-associated to non SexDiff-associated SNPs by dividing the two proportions.

For example, at our most stringent FDR threshold (0.001), the 3,016 total SexDiff-associated SNPs were located within or nearby 9 unique GO:0007548 genes and 135 total genes (to be included in Dryad Digital Repository deposition). Thus, the proportion of GO:0007548 genes = 0.0667 (9/135). In contrast, the 100,646 non-SexDiff-associated but still phenotype-associated SNPs for this same FDR threshold were located within or nearby 52 unique GO:0007548 genes and 2,207 total genes, for a proportion of 0.0236, resulting in an enrichment ratio = 2.83 (0.0667/0.0236). In other words, GO:0007548 genes with known roles in sexual differentiation are >2.8 times more likely to be co-localized with one or more SexDiff-associated SNPs than with one or more non-SexDiff-associated but still phenotype-associated SNPs at the FDR<0.001 analysis level.

We repeated this analysis for each FDR threshold (**Figure 2A**; Supplementary Table 2). The enrichment ratio steadily increased with increasingly stringent FDR significance thresholds. This pattern is consistent with expectations if our analyses are identifying true SexDiff-associated loci.

To test whether the proportion of GO:0007548 genes co-localized with at least one SexDiff-associated SNP was significantly greater than expected by chance, we used a simple permutation scheme. From the set of 2,191 total GO classified genes that were co-localized with at least one phenotype-associated SNP, we randomly selected the same number of total genes that were co-localized with one or more SexDiff-associated SNPs at the FDR cutoff being considered (e.g. 135 genes at FDR<0.001) and counted the number of GO:0007548 genes represented. We repeated this process 10,000 times for each FDR cutoff and compared the resulting distributions to the actual number of GO:0007548 genes co-localized with at least one SexDiff-associated SNP to calculate empirical P-values. The observed proportion of GO:0007548 genes is very unlikely to be explained by chance at the most stringent FDR<0.001 threshold (P=0.0045; **Figure 2B**), with

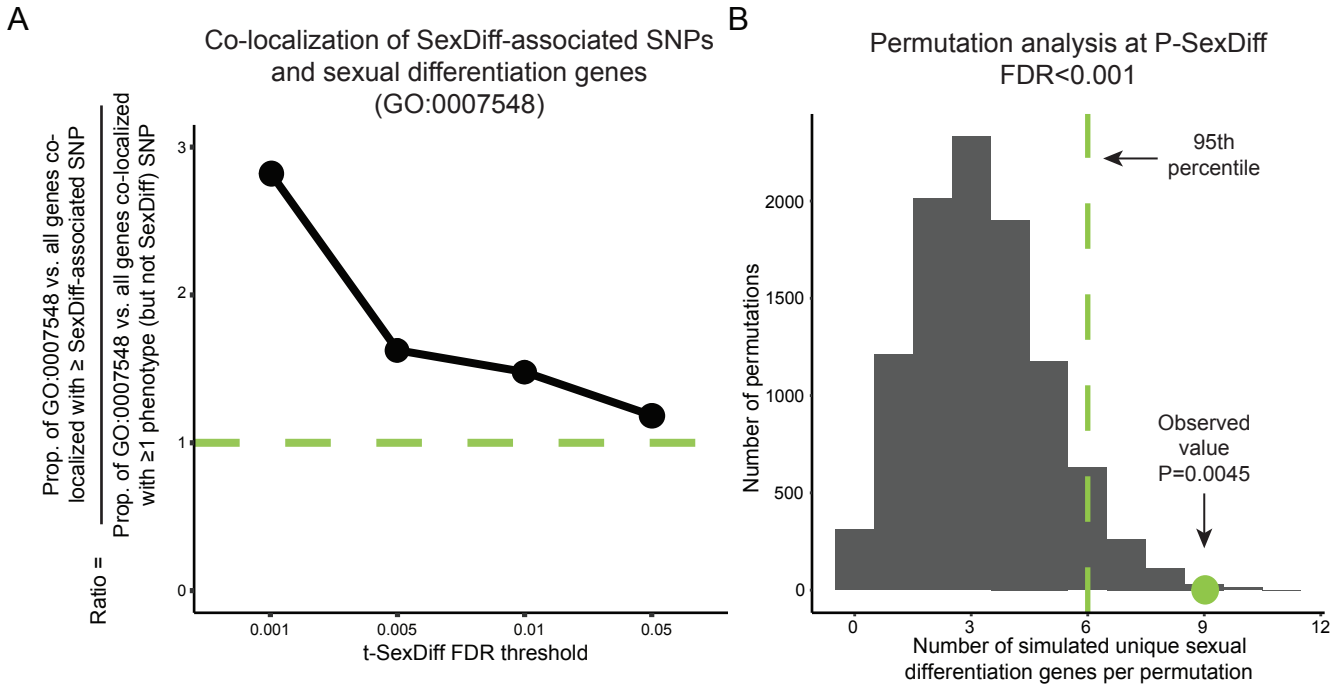

Figure 2. Genes involved in sexual differentiation (GO:0007548) are enriched for SexDiff-associated SNPs. (A) For each t-Sexdiff FDR threshold, we computed the proportion of the number of genes in the "Sexual Differentiation" Gene Ontology category (GO:0007548) to the number of all Gene Ontology genes with at least one co-localized SexDiff-associated SNP (+/- 10,000 base pairs). We also computed the same proportion for the set of SNPs significant associated with our studied phenotypes in general but *not* with sexual differentiation. Values shown are ratios of these two proportions at each t-Sexdiff FDR threshold. The green line indicates the 1:1 ratio that would be expected in the absence of any disproportionate co-localization between SexDiff-associated SNPs and GO:00007548 genes. (B) Permutation analysis of the number of genes involved in sexual differentiation co-localized with at least one SexDiff-associated SNPs at our most stringent FDR threshold (q<0.001). From the set of 2,191 total GO-classified genes that were co-localized with at least one phenotype-associated SNP, we randomly selected 135 genes, the number of total genes that were co-localized with one or more SexDiff-associated SNPs at the FDR<0.001 cutoff. Of these 135 genes, we counted and recorded the number of GO:0007548 genes represented. We repeated this process 10,000 times and computed an empirical p-value (P=0.0045) as the proportion of permutations with a greater than or equal number of GO:0007548 genes as the observed value for FDR=<.001 SexDiff-associated SNPs (9 genes). Results for similar analyses based on t-SexDiff FDR thresholds 0.005, 0.01, and 0.05 are shown in Supplementary Figure 1.

increasing probability for the FDR 0.005, 0.01, and 0.5 thresholds (P=0.0426, P=0.0647, and P=0.1713, respectively; Supplementary Figure 1).

Altogether, these results suggest that the anthropometric trait SexDiff-associated SNPs we identified – especially those discovered with the most stringent FDR<0.001 threshold – are at least enriched for SNPs that do underlie (or are linked to those that do) sexual differentiation variation. We thus proceeded to study the evolutionary histories of these SNPs. All subsequent analyses were limited to SNPs classified based on the FDR<0.001 analysis.

Signatures of positive selection on SexDiff-associated alleles

We next tested whether the SexDiff-associated SNPs identified in our analyses are significantly enriched for signatures of recent (~3,000 years) positive selection, using the Singleton Density Score statistic (SDS) (32). Briefly, alleles affected by recent positive selection are predicted to be found on haplotypes with relatively fewer singleton mutations; the SDS quantifies this pattern.

In turn, the trait-SDS statistic reflects directionality with respect to an associated phenotype. A positive trait-SDS value reflects a recent increase in the frequency of a phenotype-increasing allele, while a negative trait-SDS value reflects an increase in the frequency of a phenotype-decreasing allele (32). We performed our analyses with previously-published SDS data computed from the whole genome sequences of 3,195 British individuals from the UK10K project (20, 33). This dataset is ideal for integration with the UK BioBank genotype-phenotype association data, given study population similarity.

To examine the evolutionary histories of alleles associated with anthropometric trait sex differences, for each phenotype we first divided our SexDiff-associated SNPs into those that are disproportionately associated with trait variation in females vs. those disproportionately associated with trait variation in males. To avoid linkage disequilibrium (LD)-based result inflation, we pruned each of these sets of SNPs to include only the one SNP (if any) with the lowest t-SexDiff FDR value located within each of 1,703 approximately LD-independent blocks of the human genome (34). For comparison to the disproportionate female and male SexDiff-associated SNPs, we prepared similar LD-pruned sets of all SNPs significantly associated with the phenotype for each of the five traits (to be included in Dryad Digital Repository deposition).

We then tested whether the average trait-SDS values for the pruned female or male SexDiff-associated SNPs were significantly different than the value for the corresponding pruned set of all phenotype-associated SNPs (**Figure 3**), using a permutation analysis. For example, there were n=21 pruned SexDiff-associated SNPs disproportionately associated with female height and n=532 pruned SNPs associated with height generally (but not necessarily with sex differences). We randomly selected 21 of the 532 pruned height-associated SNPs and calculated the average trait-SDS score. We repeated this process 10,000 times and calculated an empirical P-value based on the proportion (multiplied by two; see Methods) of permuted observations with equal or more extreme average trait-SDS values than the observed average trait-SDS for the actual female

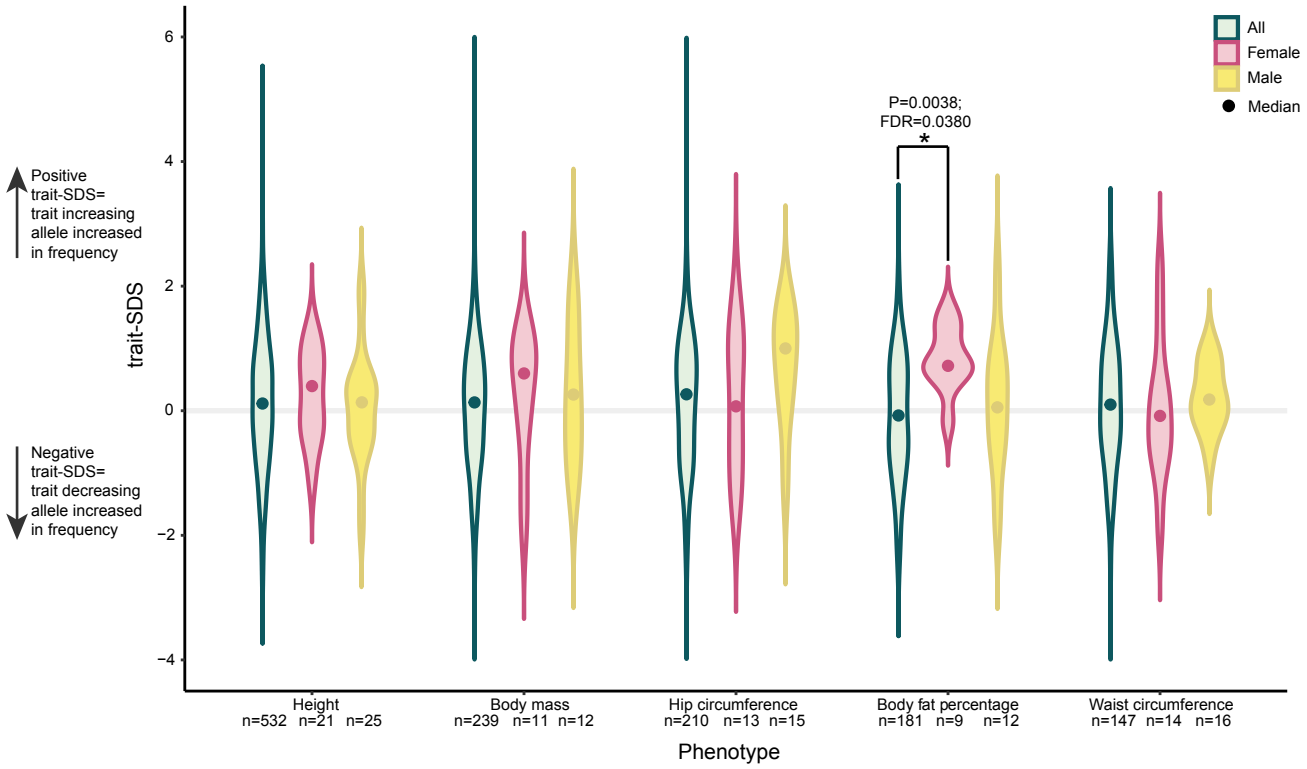

Figure 3. Distributions of trait-SDS scores for anthropometric trait-associated SNPs. For each anthropometric trait, a maximum of one SNP per each of 1,703 approximately LD-independent blocks of the human genome was included (within each block, the SNP with the strongest statistical significance) from the sets of i) all SNPs associated with trait variation, and the subsets of those SNPs disproportionately associated with variation in ii) females and iii) males (SexDiff-associated SNPs at FDR<0.001). Violin plots of trait-SDS values for these sets of SNPs are shown. Positive trait-SDS values reflects recent increases in the frequencies of phenotype-increasing alleles, while negative trait-SDS values reflect increases in the frequencies of phenotype-decreasing alleles. The trait-SDS distributions for each set of female and male SexDiff pruned SNPs were compared to that for the corresponding phenotype-association set (Supplementary Table 4).

height SexDiff-associated SNPs (average trait-SDS=0.240; P=0.66). This test was then repeated for the male SexDiff-associated SNPs for height, and the female and male SexDiff-associated SNPs for each of the remaining four anthropometric traits.

For nine of the comparisons, trait-SDS distributions for the sets of pruned SexDiff-associated SNPs were not significantly different (following FDR adjustment for multiple tests) from those for SNPs associated with the corresponding general phenotype (**Figure 3**; Supplementary Table 3). However, compared to all body fat percentage-associated SNPs (n=181 pruned SNPs; average trait-SDS=-0.124), the average trait-SDS value for the set of female SexDiff-associated SNPs for this phenotype was significantly elevated (n=9 pruned SNPs; average trait-SDS=0.827; P=0.0038; FDR=0.038). In other words, the average frequencies of alleles associated with greater body fat percentage in females have increased in frequency over the past ~3,000 years at a faster rate than expected based on the pattern for SNPs associated with body fat percentage generally, a potential signature of polygenic selection on this sub-trait.

Discussion

Using a sex-specific GWAS framework for five sexually differentiated anthropometric phenotypes, we identified 3,016 SNPs that were disproportionately associated with either female or male trait variation at a low false discovery rate (FDR<0.001). We confirmed the biological plausibility of these results by showing that genes with known roles in sexual differentiation are significantly enriched for SexDiff-associated SNPs. Together, these results confirm the importance of considering sex differences when investigating the genetic structure of human polygenic traits (35). We then used a statistic that quantifies changes in the frequencies of alleles underlying polygenic traits over the past ~3,000 years to identify a signature of recent positive selection on SNPs associated with increased female body fat percentage in the British study population.

We must emphasize that inferring selection signals from GWAS data should be approached with great care, as even subtle uncorrected population structure can impact GWAS and downstream results (36). For example, data from the GIANT consortium were previously used to identify strong signatures of polygenic selection for height across the genome (32). However, subtle population structure in the GIANT sample led to effect-size estimate biases, in turn resulting in false signals of polygenic selection for SNPs not crossing the genome-wide significance threshold and impacting results for significant SNPs as well (36). In contrast, these issues were much less prevalent using GWAS summary statistics from the UK Biobank, in which population structure is minimized (36–38). In light of these considerations, in our study we have i) used UK Biobank GWAS summary statistics only, ii) focused solely on phenotype-associated SNPs below the genome-wide significance threshold, and iii) restricted our evolutionary analyses to direct comparisons between SNPs significantly associated with individual phenotypes and a sub-phenotype (i.e., sex differences).

Our study further demonstrates the value of GWAS-based approaches for testing anthropological hypotheses (39). Concerning the evolution of human body size and shape phenotypes, our results fail to provide support for the prevailing notion of recent (i.e. subsequent to agriculture) adaptive

reductions in levels of sex differences for such traits. Specifically, using large samples of genomes from British individuals we did not observe significant differences in the recent evolutionary trajectories of SNPs disproportionately associated with female or male variation in height, body mass, hip circumference, and waist circumference relative to the trajectories of SNPs associated with these traits generally.

Moreover, we found evidence that the average frequencies of alleles disproportionately associated with greater female body fat percentage significantly increased over the past 3,000 years, a pattern consistent with polygenic adaptation. Since SNPs can be pleiotropically associated with multiple phenotypes (34), we cannot definitively conclude that positive selection acted directly on female body fat percentage. Regardless, at the very least there is still no support for the prevailing hypothesis concerning the decrease of sex differences in recent human evolution. That is, given that females have higher average body fat percentages than men in historic and contemporary populations, the direction of polygenic adaptation in the population we studied would actually be *opposite* to expectations under hypotheses of recent adaptive reductions in anthropometric trait sex differences in agricultural societies.

Methods

Subjects and dataset generation

We used genome-wide association study (GWAS) summary statistics generated from analyses of UK Biobank data (18, 22). The original GWAS analyses were restricted to 361,194 unrelated individuals (194,174 females and 167,020 males) of white British ancestry (based on the combination of self-report and genetic ancestry analysis) who did not have sex chromosome aneuploidies. GWAS summary statistics for each phenotype were computed separately for females and males. Because every phenotype was not defined for all individuals, some of the analyses contributing to our study included fewer than 361,194 individuals (Supplementary Table 5). Approximately 40 million SNPs were originally available for analysis, of which only those with minor allele frequencies > 0.001, an INFO score (imputation quality) > 0.8, and a Hardy-Weinberg Equilibrium p-value > $1x10^{-10}$ were retained, resulting in datasets of 13,791,468 SNPs.

We then subsequently filtered these SNPs with minor allele frequencies < 0.05 and those that were not associated with a singleton density score (SDS; for description, see below) (20). These filters resulted in a total genome-wide dataset of ~4.4 million SNPs for each phenotype.

We chose five sexually differentiated anthropometric phenotypes from the GWAS summary statistics for analysis (Supplementary Table 5). To identify phenotype-associated SNPs in female and/or male individuals, we applied the commonly used genome wide significance threshold (i.e., to account for the large number of tested SNPs) of $P=5x10^{-8}$ to the male-specific and female-specific GWAS summary statistics (to be included in Dryad Digital Repository deposition).

Scan for SNPs that are disproportionately associated with male or female trait variation

For each genome-wide SNP significantly associated with a phenotype in females, males, or both sexes, we evaluated whether there was a significant difference in the statistical strengths of

association for the female vs. male-specific GWAS results using the following t-statistic (t-SexDiff) (19).

$$t - SexDiff = \frac{b_{male} - b_{female}}{\sqrt{SE_{male}^2 + SE_{female}^2 - 2r * SE_{male} * SE_{female}}}$$

B_{male} refers to the male-specific beta value and B_{female} refers to the female-specific beta value for each genome-wide SNP. SE_{male} and SE_{female} refer to the male-specific and female-specific standard error for each genome-wide SNP, respectively. The correlation *r* (rho) for each phenotype was calculated as the Spearman rank correlation coefficient between b_{male} and b_{female} using the cor.test() function in R (version 3.5.1). t-SexDiff was converted to a two-sided P-value using the R function pt(). The effects of multiple testing were considered by computing the False Discovery Rate (FDR) for each t-SexDiff P-value using the R function p.adjust (40).

Of the phenotype-associated SNPs that were significantly associated with a given phenotype for females, males, or both sexes, we identified SexDiff-associated SNPs at four different FDR thresholds: <0.05, <0.01, <0.005, and <0.001 (to be included in Dryad Digital Repository deposition).

Assessing the biological plausibility of the SexDiff-associated SNPs

In order to confirm the biological plausibility of our SexDiff-associated SNPs, we tested whether regions of the genome functionally linked to sexual differentiation are more likely than expected by chance to contain one or more of our SexDiff-associated SNPs. Separately for each tSexDiff FDR threshold, we counted the number of unique genes in the Gene Ontology database (30, 31) that contained at least one SexDiff-associated SNP, including within a +/-10,000 base-pair (bp) window around the gene to encompass potential regulatory regions. We then counted the number of these genes with known links to processes of sexual differentiation corresponding to the Gene Ontology (GO) term GO:0007548 and computed the proportion of the numbers of GO:0007548 to all genes co-localized with >= 1 SexDiff-associated SNP. We repeated this analysis for significant phenotype- but not SexDiff-associated SNPs. In the absence of enrichment for SexDiff-associated SNPs, the ratio of these two proportions is expected to equal one.

We then used the following permutation scheme for each t-SexDiff FDR cutoff. There was a total of 2,191 GO-classified genes overlapping one or more phenotype-associated SNP (whether SexDiff-associated or not; each t-SexDiff FDR cutoff started with the same genome-wide significant set of phenotype-associated SNPs so the total number of 2,191 co-localized genes applies to each FDR cutoff). Given the number of these genes overlapping >=1 SexDiff-associated SNP for a given FDR cutoff, we randomly selected that number of unique genes from the pool of 2,191 genes and counted the number of GO:0007548 genes. We then repeated this procedure 10,000 times and computed an empirical P-value as the proportion of permuted data sets with an equal or greater to number of sexual differentiation genes when compared to our observation for the associated FDR threshold.

Identification of signatures of positive selection

To test the hypothesis that SexDiff-associated SNPs have been affected by recent (past ~3,000 years) positive selection in recent human evolution, we used the Singleton Density Score (SDS) statistic (32). We used a database of genome-wide SNP SDS scores that were computed (32) using 3,195 whole genome sequences from British individuals the UK10K project (33).

For each phenotype we split our SexDiff-associated SNPs into those that had lower P-values (as identified in the original sex specific GWAS for phenotype-associated SNPs) in females than males (female SexDiff-associated SNPs) and those that had lower p-values in males than females (male SexDiff-associated SNPs). We separately pruned each set of female SexDiff-associated SNPs and male SexDiff-associated SNPs to account for linkage disequilibrium. Specifically, if there was more than one female SexDiff-associated SNP in one of 1,703 approximately LD-independent segments of the genome (34), we only kept the female SexDiff-associated SNP for that segment with the lowest p-value for association with the phenotype in females. We did the same for male SexDiff-associated SNPs in males. After pruning the SNPs, we fixed the sign of SDS scores so that positive values indicate an increased frequency of the trait-increasing allele.

We used permutations to estimate the probability that the average trait-SDS value for each trait and sex could be observed by chance given the distribution trait-SDS scores for SNPs significantly associated with the corresponding phenotype (regardless of SexDiff-association). The set of phenotype-associated SNPs was pruned in a similar fashion to the above, with a maximum of one SNP per each of the 1,703 approximately LD-independent genome segments, chosen as the SNP with the most significant p-value in the segment regardless of whether it was most significant in females or males (to be included in Dryad Digital Repository deposition) from among those below the genome-wide significance threshold. From the pruned set of phenotype-associated SNPs, we then randomly selected a number of SNPs equal to the number observed SexDiff-associated SNPs for that phenotype and the sex and FDR threshold being considered, and we calculated the average trait-SDS score for that set of SNPs. We repeated this procedure 10,000 times and computed the proportion of permuted data sets with equal or more extreme average trait-SDS scores compared to actual result for the observed SexDiff-associated SNPs. This proportion was then multiplied by two to account for the two-tailed nature of this test (i.e. the average trait-SDS values for SexDiff-associated SNPs could have been significantly greater than or less than that for the phenotype-associated SNPs). We computed FDR values to account for multiple testing.

Computational resources

All analyses were conducted in R (version 3.5.1) with Advanced CyberInfrastructure computational resources provided by The Institute for CyberScience at Pennsylvania State University. All scripts are available at https://github.com/audreyarner/dimorphism-evolution.

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Figure Captions

Figure 1. SexDiff-associated SNPs for five anthropometric phenotypes. (A) Manhattan plot depicting -$\log_{10}$ P-values for the association of each genome-wide SNP with female height. The black line corresponds to the genome-wide significance threshold of $P=5x10^{-8}$. (B) Manhattan plot for SNP associations with male height. (C) For each of the 67,738 SNPs significantly associated female and/or male height, we used the equation shown to test whether the SNP was disproportionately associated with height between the sexes (height SexDiff-associated SNPs). The plot depicts -$\log_{10}$ P-values for the t-SexDiff statistic. Green bars correspond to four different FDR cutoffs. (D) SexDiff association analyses for significant phenotype-associated SNPs (number of SNPs included in each analysis is shown to the right of each plot) for four additional anthropometric traits: body mass, hip circumference, body fat percentage, and waist circumference.

Figure 2. Genes involved in sexual differentiation (GO:0007548) are enriched for SexDiff-associated SNPs. (A) For each t-SexDiff FDR threshold, we computed the proportion of the number of genes in the "Sexual Differentiation" Gene Ontology category (GO:0007548) to the number of all Gene Ontology genes with at least one co-localized SexDiff-associated SNP (+/- 10,000 base pairs). We also computed the same proportion for the set of SNPs significant associated with our studied phenotypes in general but *not* with sexual differentiation. Values shown are ratios of these two proportions at each t-SexDiff FDR threshold. The green line indicates the 1:1 ratio that would be expected in the absence of any disproportionate co-localization between SexDiff-associated SNPs and GO:00007548 genes. (B) Permutation analysis of the number of genes involved in sexual differentiation co-localized with at least one SexDiff-associated SNPs at our most stringent FDR threshold (q<0.001). From the set of 2,191 total GO-classified genes that were co-localized with at least one phenotype-associated SNP, we randomly selected 135 genes, the number of total genes that were co-localized with one or more SexDiff-associated SNPs at the FDR<0.001 cutoff. Of these 135 genes, we counted and recorded the number of GO:0007548 genes represented. We repeated this process 10,000 times and computed an empirical p-value (P=0.0045) as the proportion of permutations with a greater than or equal number of GO:0007548 genes as the observed value for FDR=<.001 SexDiff-associated SNPs (9 genes). Results for similar analyses based on SexDiff FDR thresholds 0.005, 0.01, and 0.05 are shown in Supplementary Figure 1.

Figure 3. Distributions of trait-SDS scores for anthropometric trait-associated SNPs. For each anthropometric trait, a maximum of one SNP per each of 1,703 approximately LD-independent blocks of the human genome was included (within each block, the SNP with the strongest statistical significance) from the sets of i) all SNPs associated with trait variation, and the subsets of those SNPs disproportionately associated with variation in ii) females and iii) males (SexDiff-associated SNPs at FDR<0.001). Violin plots of trait-SDS values for these sets of SNPs are shown. Positive trait-SDS values reflects recent increases in the frequencies of phenotype-increasing alleles, while negative trait-SDS values reflect increases in the frequencies of phenotype-decreasing alleles. The trait-SDS distributions for each set of female and male SexDiff pruned

SNPs were compared to that for the corresponding phenotype-association set (Supplementary Table 4).

Supplementary Figure 1. Permutation enrichment distribution at each FDR threshold. Permutation analysis of the number of genes involved in sexual differentiation for all anthropometric SNPs at every FDR threshold. Data are the frequency of distribution of our results for 10,000 permuted data sets. The empirical P-value represents the probability that the observed value of sexual differentiation genes from our SexDiff-associated SNP pool is equal to or greater than those from a randomly selected set.

Supplementary Table 1: Observed number of SexDiff-associated SNPs at each FDR threshold for every phenotype.

Supplementary Table 2: Observed unique sexual differentiation genes (SDG) and total number of genes for SexDiff-associated SNPs and Non SexDiff-associated SNPs

Supplementary Table 3: Observed trait-SDS and permutation P-values for each set of Female SexDiff-associated SNPs and Male SexDiff-associated SNPs permutated against phenotype-associated SNPs.

Supplementary Table 4. Observed trait-SDS for each set of pruned phenotype-associated SNP groups.

Supplementary Table 5: Phenotype information