Impact of natural selection on global patterns of genetic variation and association with clinical phenotypes at genes involved in SARS-CoV-2 infection

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Human genomic diversity has been shaped by both ancient and ongoing challenges from viruses. The current coronavirus disease 2019 (COVID-19) pandemic caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has had a devastating impact on population health. However, genetic diversity and evolutionary forces impacting host genes related to SARS-CoV-2 infection are not well understood. We investigated global patterns of genetic variation and signatures of natural selection at host genes relevant to SARS-CoV-2 infection (angiotensin converting enzyme 2 [ACE2], transmembrane protease serine 2 [TMPRSS2], dipeptidyl peptidase 4 [DPP4], and lymphocyte antigen 6 complex locus E [LY6E]). We analyzed data from 2,012 ethnically diverse Africans and 15,977 individuals of European and African ancestry with electronic health records and integrated with global data from the 1000 Genomes Project. At ACE2, we identified 41 nonsynonymous variants that were rare in most populations, several of which impact protein function. However, three nonsynonymous variants (rs138390800, rs147311723, and rs145437639) were common among central African hunter-gatherers from Cameroon (minor allele frequency 0.083 to 0.164) and are on haplotypes that exhibit signatures of positive selection. We identify signatures of selection impacting variation at regulatory regions influencing ACE2 expression in multiple African populations. At TMPRSS2, we identified 13 amino acid changes that are adaptive and specific to the human lineage compared with the chimpanzee genome. Genetic variants that are targets of natural selection are associated with clinical phenotypes common in patients with COVID-19. Our study provides insights into global variation at host genes related to SARS-CoV-2 infection, which have been shaped by natural selection in some populations, possibly due to prior viral infections.

SARS-CoV-2/COVID-19 | genetic variation | phenotype association | natural selection | African diversity

Coronavirus disease 2019 (COVID-19) is caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). Coronaviruses are enveloped, positive-sense, and single-stranded RNA viruses, many of which are zoonotic pathogens that crossed over into humans. Seven coronavirus species, including SARS-CoV-2, have been discovered that, depending on the virus and host physiological condition, may cause mild or lethal respiratory disease. There is considerable variation in disease prevalence and severity across populations and communities. Importantly, minority populations in the United States appear to have been disproportionally affected by COVID-19 (1, 2). For example, in Chicago, more than 50% of COVID-19 cases and nearly 70% of COVID-19 deaths are in African Americans (who make up 30% of the population of Chicago) (1). While social and economic factors are largely responsible for driving COVID-19 health disparities, investigating genetic diversity at host genes related to SARS-CoV-2 infection could help identify functionally important variation, which may play a role in individual risk for severe COVID-19 infection.

In this study, we focused on four key genes playing a role in SARS-CoV-2 infection (3). The ACE2 gene, encoding the angiotensin-converting enzyme-2 protein, was reported to be a main binding site for severe acute respiratory syndrome coronavirus (SARS-CoV) during an outbreak in 2003, and evidence showed stronger binding affinity to SARS-CoV-2, which enters the target cells via ACE2 receptors (3, 4). The ACE2 gene is located on the X chromosome (chrX); its expression level varies among populations (5); and it is ubiquitously expressed in the lung, blood vessels, gut, kidney, testis, and brain, all organs that appear to be affected as part of the COVID-19 clinical

Significance

Viruses are strong sources of natural selection pressure during human evolutionary history. Investigating genetic diversity and detecting signatures of natural selection at host genes related to severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection help to identify functionally important variation. We conducted a large study of global genomic variation at host genes that play a role in SARS-CoV-2 infection with a focus on underrepresented African populations. We identified nonsynonymous and regulatory variants at ACE2 that appear to be targets of recent natural selection in some African populations. We detected evidence of ancient adaptive evolution at TMPRSS2 in the human lineage. Genetic variants that are targets of natural selection are associated with clinical phenotypes common in patients with coronavirus disease 2019.

The authors declare no competing interest.

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SARS-CoV-2 infects cells through a membrane fusion mechanism, which in the case of SARS-CoV, is known to induce down-regulation of ACE2 (7). Such down-regulation has been shown to cause inefficient counteraction of angiotensin II effects, leading to enhanced pulmonary inflammation and intravascular coagulation (7). Additionally, altered expression of ACE2 has been associated with cardiovascular and cerebrovascular disease, which is highly relevant to COVID-19 as several cardiovascular conditions are associated with severe disease. TMPRSS2, located on the outer membrane of host target cells, binds to and cleaves ACE2, resulting in activation of spike proteins on the viral envelope and facilitating membrane fusion and endocytosis (8). Two additional genes, DPP4 and LY6E, have been shown to play an important role in the entry of SARS-CoV-2 virus into host cells. DPP4 is a known functional receptor for the Middle East respiratory syndrome coronavirus (MERS-CoV), causing a severe respiratory illness with high mortality (9, 10). LY6E encodes a glycosylphosphatidylinositol-anchored cell surface protein, which is a critical antiviral immune effector that controls coronavirus infection and pathogenesis (11). Mice lacking LY6E in hematopoietic cells were susceptible to murine coronavirus infection (11).

Previous studies of genetic diversity at ACE2 and TMPRSS2 in global human populations did not include an extensive set of African populations (5, 12–14). No common coding variants (defined here as minor allele frequency [MAF] > 0.05) at ACE2 were identified in any prior population studies. However, few studies included diverse indigenous African populations whose genomes harbor the greatest diversity among humans. This leads to a substantial disparity in the representation of African ancestries in human genetic studies of COVID-19, impeding health equity as the transferability of findings based on non-African ancestries to African populations can be low (15). Including more African populations in studying the genetic diversity of genes involved in SARS-CoV-2 infection is extremely necessary. Additionally, the evolutionary forces underlying global patterns of genetic diversity at host genes related to SARS-CoV-2 infection are not well understood. Using methods to detect natural selection signatures at host genes related to viral infections helps identify putatively functional variants that could play a role in disease risk.

We characterized genetic variation and studied natural selection signatures at ACE2, TMPRSS2, DPP4, and LY6E in ethnically diverse human populations by analyzing 2,012 genomes from ethnically diverse Africans (referred to as the “African diversity dataset”), 2,504 genomes from the 1000 Genomes Project (1KG), and whole-exome sequencing of 15,977 individuals from Africa (KAHG), including its receptor binding domain (RBD) region, which binds to the SARS-CoV-2 spike protein, dimerization interface, and transmembrane helix. In particular, two nonsynonymous variants Gly354Asp (chrX:15581230 C>T) and Ser434Asn (chrX:15600784 C>T) are both found directly in the RBD binding region of ACE2 (Fig. 1D and Dataset S1); the former is only found in low frequency in one population, the Fulani from Cameroon (MAF = 0.004), and the latter is also an African-specific variant that is at low frequency in only three East African populations, two of which are Afroasiatic-speaking populations from Kenya (MAF = 0.018) and Ethiopia (MAF = 0.008) (Dataset S1). The variant Arg708Trp (rs776995986) occurs in the region identified as the TMPRSS2 cleavage site in ACE2 (20) and is found only in the Afroasiatic-speaking populations from Ethiopia (MAF = 0.002). The presence of arginine residues has been shown to be important in “multibasic” cleavage sites (3). Therefore, due to the drastic change in physiochemical properties of the residue, this variation could be expected to interfere in TMPRSS2 cleavage efficiency, although it warrants experimental validation. Finally, two variants are located at glycosylation sites. Variant Asn546Ser (rs756905974), which causes the loss of a conserved glycosylation site on the ACE2 protein, is found only in the South Asian (SAS) populations (MAF = 0.001). Variant Lys26Arg (rs4646116), found in individuals from the European (EUR; MAF = 0.005), African (AFR; MAF = 0.001), and SAS (MAF = 0.002) populations from the 1KG dataset (Dataset S1), occurs near both the conserved ACE2
glycosylation site Asn90 and the RBD binding site. The modification to a similarly positively charged positive residue could suggest a role for electrostatic interactions, although no direct interference with RBD binding could be deduced without further studies.

**Regulatory Variation at ACE2 among Global Populations.** In contrast to coding variants, which have direct effects on protein structure in all cells expressing a gene, the effects of regulatory genetic variants are relatively difficult to determine (21). We first extracted 2,053 expression quantitative trait loci (eQTLs) significantly associated with ACE2 gene expression (P < 0.001) from the Genotype-Tissue Expression (GTEx) project database (6) (Dataset S1). To narrow down candidate functional variants, we focus on the eQTLs located in the promoter regions of target genes or in enhancers supported by chromatin interaction data (22) (Dataset S1).

We identified six eQTLs (rs4830977, rs4830978, rs5936010, rs4830979, rs4830980, and rs5934263) located in a strong deoxyribonuclelease (DNase) peak 73.3 kb upstream of ACE2 that have direct interactions with ACE2 based on RNA Pol2 chromatin interaction analysis by paired-end tag sequencing (ChIA-PET) data (Fig. 1E, Dataset S1, and SI Appendix, Fig. S2). All six single nucleotide polymorphisms (SNPs) are eQTLs of ACE2, and all of them have positive normalized effect sizes (NESs; NES > 0.2) and significant P values (P < 0.00008) in brain, bilateral nerve, bilateral artery, pituitary, and prostate cells (Dataset S1 and SI Appendix, Fig. S3). In non-African populations, these six eQTLs are in high linkage disequilibrium (LD; R² = 0.91 to 1.0) (SI Appendix, Fig. S4), and thus, there are two common haplotypes: “CCGGAT” and “ATCATC.” The frequency for the ATCATC haplotype ranges from 0.31 to 0.47 in all populations except the East Asian population, which has a frequency of 0.068 at all six SNPs (Fig. 1F). In African populations, LD is lower (R² > 0.5) (SI Appendix, Fig. S4), and there are three common haplotypes: CCGGAT (0.564), ATCATC (0.308), and “CCGGAC” (0.116). Of note, every allele in the haplotype CCGGAT is correlated with higher expression of ACE2 in the cortex of the brain, while alleles in haplotype ATCATC are correlated with lower expression of ACE2; other haplotypes have alleles with both positive and negative effect sizes in different tissues (Dataset S1). Haplotype CCGGAC is only present in populations with African ancestry, and its frequency is highest in the Botswana Khoesan (0.38) and Cameroon CAHG (0.38) hunter-gatherer populations. We also identified one variant (rs186029035) located in a strong transcription factor (TF) binding and DNase region (based on the Encyclopedia of DNA Elements, ENCODE) in the 16th intron of ACE2. This variant is only common in the Cameroon CAHG population, and therefore, there are no eQTL data for this SNP in the GTEx database (MAF = 0.153) (Dataset S1).
Signatures of Natural Selection at ACE2. As indicated above, most of the nonsynonymous variants at ACE2 are rare in global populations, and many of them are predicted to be deleterious, indicating that this gene is under strong purifying selection. To formally test for signatures of natural selection at ACE2, we first examined the ratio of nonsynonymous and synonymous variants at each gene using the the ratio of nonsynonymous to synonymous substitutions (dN/dS) test (23) (Materials and Methods). The dN/dS for all pooled samples was 0.77, indicating that ACE2 is under moderate purifying selection globally (Dataset S2 and SI Appendix, Fig. S5). However, in the East Asian population, we observed seven nonsynonymous variants (all of them are rare) and only one synonymous variant, and the dN/dS value is 1.85, indicating an excess of nonsynonymous variation. In other populations, the dN/dS ratio ranges from 0 to 0.79 (Dataset S2 and SI Appendix, Fig. S5). Thus, ACE2 appears to be under strong purifying selection in most populations but may be under weak purifying selection in the East Asian population. We next applied the McDonald–Kreitman (MK) test (24), which compares the ratio of fixed nonsynonymous sites between humans and chimpanzee (Dn = 8) and fixed synonymous sites (Ds = 6) with the ratio of polymorphic nonsynonymous sites among populations (Pn = 41) relative to polymorphic synonymous sites (Ps = 14), and found that it is not significant (odds ratio [OR] = 0.45, P = 0.94, two-sided Fisher’s exact test) (Dataset S3 and SI Appendix, Figs. S6 and S7).

Because the above-mentioned methods are more suitable for detecting signals of natural selection acting over long timescales (25, 26), we then tested for signatures of recent positive selection at ACE2 in global populations using the integrated haplotype score (iHS) test (27) to detect extended haplotype homozygosity (EHH) (28), which identifies regions of extended LD surrounding a positively selected locus. We first focused on the three common nonsynonymous variants in the CAHG population from Cameroon (rs138390800, rs147311723, and rs145437639; MAF = 0.083 to 0.164) and a common putative regulatory variant (rs186029035, located in TF and DNase regions in the 16th intron of ACE2; MAF = 0.153). The derived alleles of these variants exist on three different haplotype backgrounds; rs147311723 and rs145437639 are on the same haplotype backgrounds, while rs138390800 and rs186029035 are on similar but distinct haplotype backgrounds (Fig. 2A). The derived alleles of the corresponding SNPs on each haplotype background show EHH extending longer than 2 Mb, while the ancestral alleles of these SNPs harbor haplotypes extending less than 0.3 Mb (Fig. 2B). We then calculated the iHS of each of these variants to determine whether these extended haplotypes are unusually long compared with other SNPs with a similar allele frequency; the iHS values were not significant for any of these variants (iHS values for rs138390800, rs147311723, and rs145437639 in CAHG were 0.5, −0.08, and −0.55, respectively) (Dataset S4 and SI Appendix, Fig. S8). However, if selection was acting on multiple haplotypes simultaneously, the EHH and iHS tests would not be well powered to detect selection (29). We also used the d2 statistic (30) to measure if allele frequencies at these candidate SNPs were significantly differentiated between Cameroon CAHG and other populations. The d2 values of SNPs rs138390800 and rs186029035 were in the top 1.4% and 1.7%, respectively, of d2 values for all SNPs examined, indicating that that allele frequencies at these variants are among the most highly differentiated in the CAHG population, consistent with local adaptation. However, it should be noted that in the CAHG, these four variants (rs138390800, rs147311723, rs145437639, and rs186029035) are in complete LD based on D’ (D’ = 1) with the six eQTLs described above (SI Appendix, Figs. S9 and S10), indicating that the alleles are on the same haplotype background. Thus, it is not possible to distinguish if the nonsynonymous variants are targets of selection or if they are “hitchhiking” to high frequency due to selection on flanking regulatory variants. Given the high LD in the region, it is possible that multiple functional variants on the same haplotype backgrounds have been under selection.

We then investigated signatures of recent positive selection at candidate regulatory variants near ACE2 in the global data-sets. In total, there are 234 variants that had high iHS scores (|iHS| > 2) in at least one population extending over an ~200-kb region (Dataset S4), and 48% (n = 113) of these variants are either eQTLs or located at DNase hypersensitive regions, which are in high LD based on D’ (Dataset S4 and SI Appendix, Figs. S5 and S6).
Among the region near the transcription start site (TSS) (<10 kb from ACE2), there are two variants in high LD (D' = 1) that had high iHS scores in the San population from Botswana (Fig. 3A); rs150147953 is located in a DNase peak in multiple tissues, including lung, intestine, and heart, and rs2097723 is an eQTL of ACE2 in the brain (Fig. 1E, Dataset S4, and SI Appendix, Fig. S11). We also identified high iHS signals at the region 50 to 120 kb upstream of ACE2 (chrX:15650000-15720000) in the AFR 1KG population, the San from Botswana, and Niger–Congo-speaking populations from Cameroon as well as Afroasiatic- and Nilo-Saharan-speaking populations from Kenya (Fig. 3A and SI Appendix, Fig. S7). Two SNPs in this region (rs5936010 and rs5934263) have elevated iHS scores ([iHS] > 2) in the San population from Botswana and the Afroasiatic population from Kenya (Fig. 3A) and are part of the six eQTLs described above, located within a strong enhancer interacting with the promoter of ACE2 (Fig. 1F). Two additional eQTLs that are in complete LD with the six eQTLs (D' = 1) (SI Appendix, Fig. S10), rs4830984 and rs4830986, had high iHS scores in four of the five African populations listed above (all but Kenya Afroasiatic) (Fig. 3A).

We performed haplotype network analysis to examine phylogenetic relationships among haplotypes at ACE2 in global

![Diagram of haplotype network analysis](https://example.com/hapNetwork.png)

**Fig. 3.** Natural selection signatures at the upstream region of ACE2 in African populations. (A) iHS signals at the upstream region of ACE2 (chrX:15650000-15720000) in African populations. Each dot represents a SNP. Red dots denote SNPs that are significant (iHS > 2). The gray solid lines denote the gene body region of ACE2. Putatively causal tag SNPs are annotated in the plots. (B) Haplotype network over 150 kb flanking ACE2 in diverse ethnic populations. The network was constructed with SNPs that showed iHS signals in all populations and overlapped with DNase regions or eQTLs. The four functional candidates identified in Cameroonian CAHG were also included in the networks. Each pie represents a haplotype, each color represents a geographical population, and the size of the pie is proportional to that haplotype frequency. The dashed line denotes the boundary of clade 1 and clade 2. Black ovals denote haplotypes containing the corresponding variants. (C) Haplotypes containing variants rs5936010, rs5934263, rs4830984, and rs4830986 are highlighted. Red pies denote haplotypes containing the derived allele of the corresponding variants, while green pies denote haplotypes containing the ancestral allele of the corresponding variants.
populations for SNPs showing signatures of natural selection (Fig. 3 B and C). We identified two haplotype clades; one (clade 1) is nearly specific to Africans, and the other (clade 2) encompasses global populations (Fig. 3B). In the CAHG, haplotypes containing the rs138390800 (Lys341Arg) nonsynonymous variant and the rs186029035 regulatory variant are in clade 1, whereas haplotypes containing the rs147311723 (Leu731Phe) and rs145437639 (Asp597Glu) nonsynonymous variants are located in clade 2 (Fig. 3B). Haplotypes containing the two regulatory variants (rs5936010 and rs5934263) located 50 to 120 kb upstream of ACE2 are shared in global populations, and the nearby regulatory variants rs4830984 and rs4830986 are sublineages on those haplotype backgrounds (Fig. 3 B and C).

Genetic Variation at TMPRSS2 among Global Populations. The TMPRSS2 protein enhances the spike protein–driven viral entry of SARS-CoV-2 into cells (3). At this gene, we identified 48 nonsynonymous variants. Among the nonsynonymous variants, only two (rs12329760 [Val197Met] and rs75603675 [Gly8Val]) have high MAF (>0.05) in the pooled global dataset (Fig. 4A and Dataset S1). While rs75603675 is highly variable in non–East Asian populations (AFR = 0.3, AMR = 0.27, EUR = 0.4, and SAS = 0.2), it is not highly variable in East Asians (MAF = 0.02) (Fig. 4 B and C and Dataset S1). In addition, some nonsynonymous variants were common and specific to African populations. Notably, the nonsynonymous variant rs61735795 (Pro375Ser) had a high MAF in the Khoesan-speaking population from Botswana (MAF = 0.18). This variant is present at low frequency in populations from Cameroon (MAF < 0.01) and Ethiopia (MAF < 0.03) and was absent in non-African populations. The nonsynonymous variant rs367866934 (Leu403Phe) is common in the Cameroon CAHG population (MAF = 0.15) and has low frequency (MAF = 0.02) in other populations from Cameroon, but it is absent from non-Cameroonian populations (Fig. 4B and Dataset S1). Another nonsynonymous variant rs61735790 (His18Arg) is common in the CAHG populations from Cameroon (MAF = 0.12) and the Nilo-Saharan populations from Ethiopia (MAF = 0.12) but is rare in other populations (Fig. 4B and Dataset S1).

We identified two regulatory SNPs (rs76833541 and rs4283504) in the promoter region of the TMPRSS2 gene that have been identified as eQTLs of TMPRSS2 in tests (4, Dataset S1, and SI Appendix, Fig. S12). The MAF of rs76833541 is higher in EUR (MAF = 0.16) than other populations (EAS = 0.002, AFR = 0.006, AMR = 0.06, and SAS = 0.05), and the MAF of rs4283504 is more common in EAS (MAF = 0.21) than other populations (EUR = 0.11, AFR = 0.04, AMR = 0.12, and SAS = 0.14) (Dataset S1 and SI Appendix, Fig. S13).

Signatures of Natural Selection at TMPRSS2. We applied the MK test at TMPRSS2 and observed that Dn/Ds (13/2) is significantly larger than Pn/Ps (48/45) among pooled human samples (OR = 6.1, P value = 0.009, Fisher’s exact test) (Fig. 5A and Dataset S3) as well as in individual ethnic groups (OR ranged from 5.0 to 17), indicating positive selection in the hominin lineage after divergence from chimpanzee. Notably, there are 13 nonsynonymous and 2 synonymous variants at TMPRSS2 (ENST00000398585.7) (SI Appendix, Fig. S14 shows ENST00000332149.10) that were fixed in human populations. The nonsynonymous variants are located in different structural domains of TMPRSS2. Amino acids A3P, N10S, and S189I are located in the transmembrane region, N144K is located in the extracellular region, S165N and S178G are located in the low-density lipoprotein (LDL) receptor class A domain, E441Q and T515M are located in the serine peptidase (Peptidase S1) domain that is involved in the interaction with the SARS-CoV-2 spike protein (3), and S529G is located in the last amino acid position of the protein (Fig. 5B). In contrast to the MK test, the dN/dS ratio test was not significant in any population, indicating no excess of non-synonymous to synonymous variation within populations (Dataset S2 and SI Appendix, Fig. S5).

We also tested for recent positive selection at TMPRSS2 in all ethnic groups using iHS (Dataset S4 and SI Appendix, Fig. S15). We found many SNPs (n = 153) with high iHS scores ([iHS > 2]) in different ethnic groups in a 78-kb region encompassing the TMPRSS2 gene that show high levels of LD (chrX:...
41454000-41541000) (SI Appendix, Figs. S16 and S17). We identified a nonsynonymous variant (rs150969307) that shows a signature of positive selection (iHS = 2.01) and is common only in the Chabu hunter-gatherer population from Ethiopia (MAF = 0.079) (Dataset S4). We found that more than one-third of SNPs with |iHS| > 2.0 (62 of 153) are located in putative regulatory regions (Dataset S4 and SI Appendix, Fig. S18).

**Genetic Variation and Signatures of Natural Selection at DPP4 and LY6E.** DPP4 is a receptor for MERS-CoV and was reported to interact with SARS-CoV-2 (10). At this gene, we identified 47 nonsynonymous variants and 1 loss-of-function variant (Dataset S1). There were no common nonsynonymous variants in the pooled global dataset (SI Appendix, Fig. S19A), suggesting that this gene is extremely conserved during human evolutionary history. Only one nonsynonymous variant (rs1129599, Ser43Thr) was common in the Fulani pastoralists from Cameroon (MAF = 0.081), was present at low frequency in other African populations, and was absent in non-African populations (SI Appendix, Fig. S19 B and C). In addition to the nonsynonymous variants, one loss-of-function variant was identified at DPP4. The variant rs149291595 (Q170*) has low MAF in some African populations (MAF < 0.05) but is absent in non-African populations.

We identified four eQTLs (rs1861978, rs35128070, rs17574, and rs13015258) in the promoter region of the DPP4 gene (SI Appendix, Figs. S19D and S20). Three of the variants (rs1861978, rs35128070, and rs17574) are significant eQTLs in the transverse colon, and rs13015258 is an eQTL in the lung (P < 5.9e-6) (Dataset S1 and SI Appendix, Fig. S19). The minor alleles of these three variants are rare in EAS (MAF < 0.05) but common in all other populations (MAF > 0.15) (Dataset S1 and SI Appendix, Fig. S21). The fourth SNP, rs13015258, resides in the center of a cluster of DNase peaks identified in ENCODE (SI Appendix, Fig. S19D), with MAF ranging from 0.38 in the AMR population to 0.6 in other populations (Dataset S1 and SI Appendix, Fig. S21).

The MK test result of DPP4 was not significant in either the pooled samples (Dn = 3, Ds = 5, Pn = 45, Ps = 33; OR = 0.44, P = 0.9, two-sided Fisher’s exact test) or each population separately (Dataset S3 and SI Appendix, Fig. S7). For the dN/dS test, we observed ratios ranging from 0 to 0.52 in individual populations, indicating that DPP4 is highly conserved (Dataset S2 and SI Appendix, Fig. S5) within human populations. Using the iHS test, we identified eight SNPs that had extremely high iHS scores (|iHS| > 2) in the Khoesan populations from Botswana (Dataset S4 and SI Appendix, Fig. S22). Five of these SNPs (rs10166124, rs2284872, rs2284870, rs7600879, and rs2160927) are in LD (D’ > 0.95) with each other (SI Appendix, Fig. S23). The SNP rs2284870 is located in a strong DNase peak in heart tissue (Dataset S4 and SI Appendix, Fig. S24).

Studies show that mice lacking LY6E were highly susceptible to a usually nonlethal mouse coronavirus (11). At LY6E, we observed 28 nonsynonymous variants, and all of them, except rs11547127 (MAF = 0.057), have MAFs that are rare in the pooled global dataset (Dataset S1 and SI Appendix, Fig. S25A). However, some nonsynonymous variants are common in specific populations (SI Appendix, Fig. S25B). For instance, the nonsynonymous variant rs111560737 (Asp104Asn) was common in the southern African Khoesan population from Botswana (MAF = 0.36) and the Chabu population from Ethiopia (MAF = 0.17) (SI Appendix, Fig. S25C). Three loss-of-function variants (rs200177123 [stop gained, Ser59*], chr8:143020941, and chr8:143020946) were also identified at LY6E, and all of them are rare.

We identified three regulatory eQTLs (rs13252864, rs17061979, and rs114909654) located within 2 kb of the transcription start site of LY6E (SI Appendix, Fig. S25D), all of which are significant in esophageal mucosa (P < 1e-5) (Dataset S1 and SI Appendix, Fig. S26), which has a high expression level of LY6E (transcript per million, TPM = 108, GTEx). The minor alleles of rs13252864 and rs114909654 are common in African populations (MAF > 0.15) while very rare in other populations (MAF < 0.02) (SI Appendix, Fig. S27), whereas the MAF of rs17061979 is relatively high in EAS (0.18) and SAS (0.13) and rare in other populations (MAF < 0.05) (SI Appendix, Fig. S27).

The MK test for LY6E was not significant in either the pooled samples (Dn = 0, Ds = 4, Pn = 9, Ps = 9; OR = 0, P = 0.9, two-sided Fisher’s exact test) or each population separately (OR ranging from 0 to 0.52) (Dataset S3 and SI Appendix, Fig. S7). For the dN/dS test, we observed ratios ranging from 0 to 0.68 in individual populations, indicating that LY6E is highly conserved (Dataset S2 and SI Appendix, Fig. S5). We identified 19 variants that had extreme high iHS scores (|iHS| > 2) (Dataset S4 and SI Appendix, Fig. S28), some of which are in LD in specific populations (SI Appendix, Fig. S29). One variant (rs867069115) shows an extreme iHS score in the Hadza hunter-gatherer population from Tanzania (iHS = −2.94). This variant is located in a regulatory region ∼1.9 kb downstream of LY6E within DNase and TF peaks in the lung, intestine, kidney, heart, stomach, pancreas, and skeletal muscle from ENCODE (SI Appendix, Fig. S30); is common only in the Hadza population (MAF = 0.14); is rare in other African populations (MAF < 0.05); and is absent in all non-African populations (Dataset S1). SNP rs10283236, which shows an extreme iHS value in the CEU population, is an eQTL of LY6E located within DNase and TF clusters identified in ENCODE (~4.14 kb downstream of LY6E) active in many tissues, including lung, kidney, and small intestine.

**Associations between Genetic Variation in Host Genes and Clinical Disease Phenotypes.** We examined associations of genetic variation at four host genes relevant to SARS-CoV-2 infection with clinical phenotypes using the PMBB cohort that...
consists of exome-sequencing data from 15,977 participants between the ages of 19 and 89 years (52% female) with extensive clinical data available through their EHRs. Of these, 7,061 individuals were of European ancestry (EA) (42%), and 8,916 were of African ancestry (AA) (55%) (SI Appendix, Table S1).

**Gene burden phenotype-association study with clinical phenotypes.** To test for the association between rare coding variants and clinical phenotypes, we applied a gene-based approach (32, 33) as well as single-variant analysis. First, we performed a gene-based analysis by collapsing the coding region variants with MAF < 0.01 that are annotated as nonsynonymous or putative loss-of-function (pLOF) variants. We examined ∼1,800 phenodes derived from the EHR and performed a phenotype-wide association study (PheWAS) with individual SNPs (32–34). After multiple testing correction, we identified one association in the AA population and five associations in the EA population reaching study-wide significance ($P = 6.6 \times 10^{-6}$) ([0.05/(1,866 codes × 4 genes)]) (Dataset S5). Myocarditis, a rare cardiovascular disease caused by viral infection, was the top PheWAS association with $ACE2$ in the AA population but was not significant in the EA population. Although the population difference for this specific association is unclear, recent studies have reported that COVID-19 patients have a 16 times higher risk of myocarditis (35). Furthermore, $ACE2$ is expressed in heart tissue and its upregulation in cardiomyocytes has an important role in both dilated cardiomyopathy and hypertrophic cardiomyopathy (36–38).

**Gene burden association analyses with 12 COVID-19–relevant organ dysfunctions.** We tested for the association of rare coding variants with 12 phenotypes, encompassing COVID-19–relevant disease classes affecting different organ systems, defined by EHR-based diagnosis codes (Dataset S5). In the AA population, the phenotypes with the most significant associations with $ACE2$ were hepatic encephalopathy and respiratory failure (Fig. 6A and Table 1). The association with respiratory failure is interesting as it is one of the key severe clinical features reported for COVID-19 (39–43). However, the same association was not significant in the EA population, which could be explained by lack of power due to a lower number of coding variants at $ACE2$ in EA. Within the EA population, the most significant associations with $ACE2$ included hepatic coma, respiratory syncytial virus infectious disease, and cirrhosis of the liver (Table 1). In the gene-based analysis of variants in $DPP4$, we identified significant associations (only in the sequence kernel association test [SKAT] model) with respiratory syncytial virus infectious disease and upper respiratory tract disease in the AA population (Fig. 6A, Table 1, and Dataset S5); this observation was not observed in the EA population. None of the associations with $DPP4$, $TMPRSS2$, and $LY6E$ reach statistical significance in gene burden analysis.

**PheWAS of eQTLs near COVID-19 host immunity-related genes.** Lastly, we conducted PheWAS of eQTLs identified near the host genes. For the six eQTLs identified near $ACE2$, rs5936010 and rs5934263 (targets of positive selection in both Afroasian populations from Kenya and Khoesan populations from Botswana) were significantly associated with type 2 diabetes ($P = 1.23 \times 10^{-5}$, OR = 1.1) and hypertension ($P = 8.8 \times 10^{-4}$, OR = 1.13), respectively. The association was only observed in the AA population (Fig. 6B and Dataset S6). The PheWAS of the two regulatory eQTLs (rs76833541 and rs4283504) near $TMPRSS2$ described above identified association of rs76833541 with abnormal glucose ($P = 8.9 \times 10^{-4}$, OR = 1.5) in EA and rs4283504 with glucocorticoid deficiency ($P = 0.001$, OR = 2.7) in AA (Fig. 6B). The PheWAS of four regulatory eQTLs near $DPP4$ identified the association with malignant neoplasm of the rectum (commonly referred as colon cancer). The SNP rs17574 was associated with increased risk ($P = 4.49 \times 10^{-9}$, OR = 1.8) of colon cancer; however, the observation was only observed among AA individuals. The association analysis of regulatory variants near $LY6E$ identified significant associations with “severe protein-calorie malnutrition” (rs114909654, $P = 2.35 \times 10^{-6}$, OR = 1.9) and “acute posthemorrhagic anemia” (rs114909654, $P = 6.4 \times 10^{-4}$, OR = 1.6) in the AA population. In the EA population, “chronic ulcer of skin” with rs13252864 ($P = 0.001$, OR = 2.2) was the most significant association (Fig. 6B and Dataset S6). Among EHR-derived phenotypes for respiratory disorders, the rs35128070 eQTL near $DPP4$ was associated with “abnormal results of function study of pulmonary system” ($P = 0.002$, OR = 1.6) in the AA population (Fig. 6B and Dataset S6).

**Discussion**

Investigating global patterns of genetic variation at genes that play a role in SARS-CoV-2 infection could provide insights into potential differences in susceptibility to COVID-19 among diverse human populations. However, African populations are underrepresented in the majority of current genetic studies of COVID-19 susceptibility and severity, despite the fact that they have the highest genetic diversity among human populations (44, 45) and have high burdens of infectious disease (44, 45).

**Three Nonsynonymous Variants That Are Common and Specific to CAHG at ACE2 Are on Haplotypes with Signatures of Positive Selection.** Several studies have investigated patterns of genetic variation at $ACE2$, a receptor of SARS-CoV-2 entry (5, 12–14). None of these studies identified any common coding variation at $ACE2$, suggesting that $ACE2$ is evolutionarily conserved. However, these studies did not include an extensive set of African populations. At $ACE2$, we identified 41 nonsynonymous variants, most of which are rare, suggesting that they are under purifying selection. Tests based on dN/dS indicate that East Asians have an excess of nonsynonymous variation at $ACE2$, indicating that weak purifying selection has influenced patterns of variation in that population. However, we identified three common nonsynonymous variants (rs138390800, rs147311723, and rs145437639) at $ACE2$ with MAF ranging from 0.083 to 0.164 in CAHGs, which were the only common coding variants (defined here as MAF > 0.05) found in global populations studied here and by others (5, 12–14). We observed that the derived alleles of the common nonsynonymous SNPs (rs138390800, rs147311723, rs145437639) and one putative regulatory variant (rs186029035) at $ACE2$ in CAHG show evidence of EHH, with the extended haplotypes extending longer than 2 Mb, although they did not show deviation from neutrality based on the iHS test. However, we do not have much power to detect a selection signal using this test because the SNPs are on three different haplotype backgrounds in CAHG, possibly due to selection on existing variation (e.g., “soft selection”), which decreases the power to detect significant iHS scores. Moreover, each haplotype is at a relatively low frequency (0.083 to 0.164), which further reduces the power of the iHS test. Allele frequencies at two of the putative functional variants are among the most highly differentiated between the CAHG population and other populations, consistent with local adaptation, as indicated by the $d_i$ values of SNPs rs138390800 and rs186029035, which were in the top 1.4% and 1.7%, respectively, of $d_i$ values for all SNPs examined. The CAHGs are traditionally hunter-gatherers living in a rainforest ecosystem who consume wild
animals. They have high exposure to animal viruses and were reported to have relative resistance to viral infection (46). Thus, it is possible that this locus is adaptive for protection from infectious diseases in this population. Future in vitro or in vivo studies will be needed to determine the functional significance of these variants.

**TMPRSS2 Shows Adaptive Evolution in the Human Lineage after Divergence from Chimpanzee.** At TMPRSS2, we identified 48 nonsynonymous variants, only 2 of which had a high MAF (>0.05) in the pooled global dataset (rs12329760 and rs75603675). However, some variants have high MAF in two African hunter-gatherer populations. Notably, the nonsynonymous variant rs61735795 (Pro375Ser) is only common in the Khoesan-speaking San population from Botswana (MAF = 0.18), and the nonsynonymous variant rs367866934 (Leu403-Arg) is only common in the Cameroonian CAHG populations (MAF = 0.15). At TMPRSS2, we observed a signature of adaptive evolution in the human lineage after divergence from chimpanzee ~6 Mya (47). In total, 13 nonsynonymous variants located on different structural domains of TMPRSS2 were fixed in human populations. Among them, E441Q and T515M are located in the Peptidase S1 domain that plays an important role in acute respiratory syndrome–like (SARS) coronavirus (SARS-CoV-2) infection (48), and six (A3P, N10S, T46P, A70V, R103C, and M104T) are at the cytoplasmic amino terminal domains of TMPRSS2, which plays an important role in signal transduction. By contrast, the coding regions of DPP4 and LY6E are evolutionarily conserved.

eQTLs for Four Genes Vary in Frequency among Populations and Show Signatures of Natural Selection and Associations with Clinical Phenotypes. SARS-CoV replication is significantly reduced in ACE2 knockout mice (49), and cells with low expression of ACE2 were resistant to SARS-CoV-2 infection (50). It has also been shown that both SARS-CoV and SARS-CoV-2 infection could down-regulate ACE2 expression (8, 49, 51). The expression of ACE2 and TMPRSS2 in nasal and bronchial epithelial cells is higher in adults than children and in healthy individuals compared with smokers or patients with chronic obstructive pulmonary disease (51, 52). Therefore, differences in expression levels of ACE2 and TMPRSS2 could influence the susceptibility and host reactions to SARS-CoV-2. Previous studies identified eQTLs influencing ACE2 gene expression, showing differences in allele frequency among different populations (5, 53). For instance, the C allele at the eQTL rs1978124 (53) and the G allele at the eQTL rs4646127 (5), both associated with high ACE2 expression, are close to 100% frequency in East Asians but are <80% frequency in other populations. Recently, a rare eQTL (rs190509934, MAF = 0.3%) has been identified that is associated with decreased ACE2 expression and reduced risk of severe COVID-19 disease (54).

We systematically identified regulatory eQTLs associated with ACE2, TMPRSS2, DPP4, and LY6E gene expression and highlighted the eQTLs showing highly differentiated MAF among populations and/or signatures of natural selection. Regulatory eQTLs that differ in frequency across ethnically diverse populations may play a role in local adaptation and disease susceptibility (55). These eQTLs are located in chromatin immunoprecipitation sequencing (ChiP-seq) and DNase peaks, and they have the potential to influence transcription factor binding and thus, change the promoter or enhancer activities in specific tissues (56, 57). Notably, some of the eQTLs in the upstream regions of ACE2 were under selection in African populations. For example, rs5936010 and rs5934263, located within a
Cirrhosis of liver

Respiratory syncytial virus infectious disease

Table 1. Associations of ACE2, DPP4, TMPRSS2, and LY6E with 12 disease classes derived from EHR data

| Disease phenotype                      | Gene   | Cases | Controls | Carrier controls | Carrier cases | SKAT P  | Burden P | Burden OR | Burden SE | 95% CI   | Dataset |
|----------------------------------------|--------|-------|----------|------------------|---------------|---------|----------|-----------|-----------|----------|---------|
| Hepatic encephalopathy                 | ACE2   | 97    | 8,045    | 441              | 5             | 1.1E-12 | 0.0043   | 5.73      | 0.61      | 0.55–2.94 | AA      |
| Respiratory syncytial virus infectious disease | DPP4   | 56    | 6,392    | 85               | 1             | 6.8E-07 | 0.1221   | 6.06      | 1.17      | -0.48–4.09 | AA      |
| Respiratory failure                    | TMPRSS2| 199   | 6,392    | 11               | 2             | 2.3E-06 | 0.0124   | 7.31      | 0.80      | 0.43–3.55 | AA      |
| Respiratory failure                    | ACE2   | 199   | 6,392    | 351              | 12            | 9.0E-05 | 0.0509   | 3.10      | 0.58      | 0–2.26   | AA      |
| Upper respiratory tract disease        | DPP4   | 144   | 6,392    | 85               | 3             | 2.5E-04 | 0.0978   | 4.16      | 0.86      | -0.26–3.11 | AA      |
| Respiratory syncytial virus infectious disease | TMPRSS2| 56    | 6,392    | 11               | 1             | 3.9E-04 | 0.0217   | 11.63     | 1.07      | 0.36–4.55 | AA      |
| Hepatic coma                           | ACE2   | 16    | 6,817    | 318              | 1             | 4.3E-31 | 0.0019   | 10.45     | 0.76      | 0.87–3.83 | EA      |
| Respiratory syncytial virus infectious disease | ACE2   | 40    | 5,859    | 274              | 3             | 2.3E-07 | 0.1650   | 3.61      | 0.92      | -0.53–3.1 | EA      |
| Cirrhosis of liver                     | ACE2   | 10    | 6,817    | 43               | 1             | 1.8E-04 | 0.0837   | 9.40      | 1.30      | -0.3–4.78 | EA      |

strong enhancer interacting with the promoter of ACE2 as suggested by ChiA-PET, harbored significant iHS scores ([iHS] > 2) in both Afroasian populations from Kenya and the San population from Botswana. Further, PheWAS of these eQTLs in the PMBB populations identified association of eQTLs at ACE2 with type 2 diabetes (rs5936010) and hypertension (rs5934263). These are known preexisting conditions that increase risk of severe illness due to COVID-19 (58–60). Among respiratory diseases, only one eQTL at ACE2 had nominal association (rs4830977) with acute sinusitis. The association was only identified in the AA population and had a protective effect (OR = 0.78 [0.66 to 0.95]). The eQTLs we analyzed are from the GTEx V8 database (61), and 84.6% of the donors are people of European and western Eurasian descent. Therefore, it is possible that we are missing some regulatory variants that are only present in specific ancestry groups due to the lack of sample diversity. Further experimental testing of predicted regulatory variants will provide insights into differences in gene expression regulation at ACE2, TMPRSS2, DPP4, and LY6E among different populations.

ACE2 and TMPRSS2 Are Significantly Associated with Respiratory, Cardiac, and Blood Phenotypes in the PMBB Dataset. The gene-based genetic association analyses of nonsynonymous variants at ACE2, TMPRSS2, DPP4, and LY6E identified several associations with clinical phenotypes. We observed that respiratory failure has significant association with ACE2 and TMPRSS2 among the PMBB AA population. That is a particularly interesting finding as respiratory failure is one of the clinical outcomes observed in some patients with COVID-19 (39–43). However, this association was not significant in the EA population. This observation could be explained by the low number of coding variants and carriers at ACE2 and TMPRSS2 among EA and hence, low power to detect an association. An association with myocarditis, a rare cardiovascular disease caused by viral infection, was also observed in the AA population. Recent studies have reported a link between SARS-CoV-2–induced cardiac injury, such as myocarditis, among COVID-19 patients (62). Further, ACE2 has known expression in heart tissue, and it plays an important role in transcriptional dysregulation in cardiomyocytes–cells that make up cardiac muscles (36–38). We observed an association between ACE2 and myocarditis only in the AA population, but as noted above, we may not have as much power to detect an association in EA. Blood clotting abnormalities in lungs and other organs in COVID-19 patients have been reported by several studies (63). In autopsies of COVID-19 patients, thrombosis was found to be a prominent finding across multiple organs, even despite extensive anticoagulation treatment and regardless of the timing of clinical progression, indicating that thrombosis might be at play in the early stages of disease (63). One hypothesis to explain this observation is that the dysfunction of endothelial cells may play an important role in increased risk of thrombosis (64). We observed associations between the internationalized normalized ratio (INR) derived from the prothrombin time test with ACE2 and LY6E in a gene-based association test. The INR test measures the time it takes blood to clot and is an important measure for individuals with blood clotting disorders or on blood thinners.

Characterizing the genetic variation and clinical phenotype associations at these four genes that play a key role in SARS-CoV-2 infection could be relevant for understanding individual differences in infection susceptibility. We performed evolutionary analyses to dissect the forces underlying global patterns of genetic variation and identified variants that may be targets of selection. It will be important to determine the functional effects of these candidate adaptive variants using in vitro and in vivo approaches in future studies. Additional studies will be needed to investigate the impact of genetic variation in modulating susceptibility/resistance to SARS-CoV-2 infection and other coronaviruses across ethnically diverse populations.

Materials and Methods

Genomic Data and Populations. The genomic data used in this study were from three sources: the Africa 6K project (referred to as the African diversity dataset), which is part of the Trans-Omics in Precision Medicine consortium (65); the 1KG (66); and the PMBB. From the Africa 6K project, a subset of 2,012 high-coverage (>30x) whole-genome sequences of ethnically diverse African populations (SI Appendix, Fig. S1) was included (SI Appendix). Institutional review board (IRB) approval was obtained from the University of Maryland and the University of Pennsylvania. Written informed consent was obtained from all participants, and research/ethics approval and permits were obtained from the following institutions prior to sample collection: Tanzania Commission for Science and Technology,
National Institute for Medical Research, and Muhimbili University of Health and Allied Sciences in Dar es Salaam, Tanzania; the University of Botswana and the Ministry of Health in Gaborone, Botswana; the University of Addis Ababa and the Federal Democratic Republic of Ethiopia Ministry of Science and Technology National Health Research Ethics Review Committee; and the Cameroonian National Ethics Committee and the Cameroonian Ministry of Public Health. The PMBB participants were recruited through the University of Pennsylvania Health System by enrolling at the time of clinic visit. Patients participated by donating either blood or a tissue sample and allowing researchers access to their EHR information, and all participants provided written informed consent. The PMBB is approved under IRB protocol 813913.

Variant Annotation. We used Ensembl Variant Effect Predictor (VEP) for variant annotations (67) (SI Appendix). For gene-based association analysis using the PMBB dataset, we collapsed all the predicted nonsynonymous variants with rare exome variant ensemble (REVEL) score > 0.5 and pLoFs with MAF < 0.01. We assigned variants as pLoFs if the variant was annotated by VEP as start_lost, splice_donor_variant, splice_acceptor_variant, frameshift_variant, stop_gained, and stop_lost. All genome coordinates followed the GRCh38 assembly.

Characterization of Putative Regulatory Variation. We extracted variants located within a ±10 kb distance to their TSS as well as enhancers supported by RNA Pol2 ChIA-PET data from ENCODE (68). These variants were further filtered by overlapping with DNase I hypersensitive sites sequencing (DNase-seq) and ChiP-seq peaks from Roadmap (69), ENCODE (68), and Remap2 (70) or overlapping with significant single-tissue eQTLs (P value < 0.001) from the GTEx V8 database (6).

EHR Phenotypes and Association Testing. We focused on the phenotypes characterized as primary organ dysfunctions in the early studies on COVID-19 (Dataset S5 and SI Appendix). Broadly, we centered our analyses on these four broad clinical conditions/phenotypes: respiratory injury/failure, acute liver injury/failure, acute cardiac injury/failure, and acute kidney injury/failure. These disease classes are well characterized in human disease ontologies, such as Monarch Disease Ontology (SI Appendix).

We used the R SKAT package for conducting a gene-based dispersion test and BioClin for gene burden analysis (32, 71, 72). Here, multiple genetic variations in a gene region were collapsed to generate a gene burden/dispersions score, and regression methods were used to test for association between the genetic score and a phenotype or trait (SI Appendix).

Structural Analysis of Nonsynonymous Variations on the ACE2-S Protein Binding Interface. We determined the three-dimensional protein location of all nonsynonymous coding variants identified in this study using experimentally determined structures of the ACE2 protein complexed with the RBD of SARS-CoV-2 spike glycoprotein based on cryoelectron microscopy available in the Protein Data Bank (PDB code 6M17 (73)). All structural analysis and figures were prepared using VMD (74) (SI Appendix).

Detecting Signatures of Natural Selection. We used two methods [the McDonald-Kreitman test (24) and the dN/dS test (23)] to test for signals of selection acting on the four candidate genes over long timescales and two methods (EH) (28) and iHS (27) to detect recent (e.g., last ~10,000 y before present) signatures of positive selection (SI Appendix). We used d2 statistics to identify SNPs that are highly differentiated in allele frequency between populations based on unbiased estimates of pairwise FST (30) (SI Appendix). Haplotype networks were constructed by PopART (75) using the built-in minimum spanning algorithm.

Description of Supplemental Data. The supplemental data include SI Appendix, SI Methods, Figs. S1–S30, and Table S1, the legends of Datasets S1–S6 and the Regenere Genomic Center authors and contribution statements.

Data Availability. All data are included in the manuscript and/or supporting information.

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