

Table S2: Top 100 signals in the YRI population using the  $T_1$  test statistic.

Rank	Gene name	Chromosome	$T_1$	Rank	Gene name	Chromosome	$T_1$
1	<b><i>HLA-A</i></b> <sup>74,75</sup>	6	307.4	51	<i>WWTR1</i>	3	119.9
2	<i>FANK1</i>	10	302.2	52	<i>WDR75</i>	2	119.4
3	<b><i>HLA-B</i></b> <sup>15,74,75</sup>	6	272.3	53	<i>C1orf130</i>	1	118.2
4	<i>TEKT4</i>	2	271.7	54	<i>FOPNL</i>	16	118.1
5	<b><i>HLA-DPA1</i></b> <sup>76</sup>	6	267.1	55	<i>SGCG</i>	13	117.6
6	<b><i>HLA-C</i></b> <sup>75</sup>	6	262.7	56	<i>ZNF568</i>	19	117.0
7	<i>MLL3</i>	7	240.3	57	<i>FRAS1</i>	4	116.1
8	<i>HLA-DRB5</i>	6	227.2	58	<i>SLC39A12</i>	10	116.0
9	<i>OR4C3</i>	11	209.3	59	<i>KANK1</i>	9	115.5
10	<b><i>HLA-DQA1</i></b> <sup>14,75</sup>	6	205.4	60	<i>EMID2</i>	7	114.7
11	<i>FRG2C</i>	3	189.5	61	<i>MYRIP</i>	3	114.7
12	<b><i>HLA-DQB1</i></b> <sup>14,75</sup>	6	186.7	62	<b><i>PRIM2</i></b> <sup>79</sup>	6	114.4
13	<b><i>HLA-DPB1</i></b> <sup>14</sup>	6	180.2	63	<i>ASB18</i>	2	114.3
14	<i>SNTG2</i>	2	170.8	64	<b><i>HLA-DRA</i></b>	6	114.2
15	<i>ZNF717</i>	3	170.3	65	<i>ARHGAP24</i>	4	113.8
16	<i>POLR1E</i>	9	167.8	66	<i>KCNAB1</i>	3	113.8
17	<i>DMBT1</i>	10	166.4	67	<b><i>TRIM40</i></b> <sup>78</sup>	6	111.9
18	<i>CPE</i>	4	166.0	68	<i>LUZP2</i>	11	111.8
19	<i>RNF144B</i>	6	163.2	69	<i>ABCD4</i>	14	111.7
20	<i>CTNNA3</i>	10	160.1	70	<b><i>LGALS8</i></b> <sup>15</sup>	1	111.1
21	<b><i>HLA-DRB1</i></b> <sup>14,75</sup>	6	157.2	71	<i>BNC2</i>	9	109.4
22	<i>OR4C45</i>	11	156.4	72	<i>PTPRB</i>	12	109.3
23	<i>AXDND1</i>	1	151.8	73	<i>RBFOX1</i>	16	109.1
24	<b><i>SLC2A9</i></b> <sup>15</sup>	4	150.8	74	<i>KL</i>	13	106.6
25	<i>ZNF85</i>	19	150.4	75	<i>HEATR1</i>	1	106.3
26	<i>MAP2K3</i>	17	150.3	76	<i>POLN</i>	4	105.5
27	<i>KCNJ12</i>	17	150.1	77	<i>PDE11A</i>	2	105.1
28	<i>SORD</i>	15	148.5	78	<i>VWDE</i>	7	103.9
29	<i>ARHGAP42</i>	11	146.6	79	<i>SPATA13</i>	13	103.7
30	<i>KCNJ18</i>	17	146.3	80	<i>SPATA16</i>	3	103.6
31	<i>ARPC5</i>	1	143.7	81	<b><i>APBB1IP</i></b>	10	103.0
32	<i>MCM9</i>	6	141.1	82	<i>MPHOSPH6</i>	16	102.0
33	<i>C18orf1</i>	18	139.5	83	<i>CES5A</i>	16	101.9
34	<b><i>SLC38A9</i></b>	5	132.0	84	<i>PARP4</i>	13	100.9
35	<i>GRIN2A</i>	16	131.9	85	<b><i>KRT83</i></b>	12	100.9
36	<i>MYOM2</i>	8	131.7	86	<i>MOSC2</i>	1	100.9
37	<i>GBA3</i>	4	127.8	87	<i>PACRG</i>	6	99.1
38	<i>STK32A</i>	5	125.2	88	<b><i>RCBTB1</i></b> <sup>15</sup>	13	97.8
39	<i>CSMD1</i>	8	125.2	89	<i>FAM55A</i>	11	96.5
40	<b><i>FHIT</i></b> <sup>77</sup>	3	125.0	90	<i>GALC</i>	14	96.0
41	<i>ART3</i>	4	124.5	91	<i>LRPPRC</i>	2	95.6
42	<i>RGS6</i>	14	123.7	92	<i>SLC24A4</i>	14	95.3
43	<i>CEP112</i>	17	123.7	93	<i>ADCY5</i>	3	95.0
44	<i>IGSF5</i>	21	123.6	94	<i>ULK4</i>	3	94.9
45	<i>GRHL1</i>	2	122.8	95	<i>FYB</i>	5	94.3
46	<i>RGL1</i>	1	121.9	96	<i>KIAA0748</i>	12	93.8
47	<i>SNX19</i>	11	121.8	97	<i>COL4A3</i>	2	93.7
48	<i>SPEF2</i>	5	121.2	98	<i>CCDC169</i>	13	93.6
49	<i>FXN</i>	9	121.0	99	<i>CCDC169-SOHLH2</i>	13	93.6
50	<i>PTCHD3</i>	10	120.3	100	<i>PLEKHG1</i>	6	93.3

Previously-hypothesized genes are indicated in bold. Genes overlapping with Table S1 are highlighted in gray.

- 74 P. W. Hedrick, T. S. Whittam, and P. Parham, "Heterozygosity at individual amino acid sites: extremely high levels for *HLA-A* and *-B* genes," *Proc. Natl. Acad. Sci. USA*, vol. 88, pp. 58975901, 1991.
- 75 A. Sánchez-Mazas, "An apportionment of human HLA diversity," *Tissue Antigens*, vol. 69, pp. 198202, 2005.
- 76 P. G. Bronson, S. J. Mack, H. A. Erlich, and M. Slatkin, "A sequence-based approach demonstrates that balancing selection in classical human leukocyte antigen (HLA) loci is asymmetric," *Hum. Mol. Genet.*, vol. 22, pp. 252261, 2013.
- 77 Y. Ding, G. Larson, G. Rivas, C. Lundberg, L. Geller, C. Ouyang, J. Weitzel, J. Ar-  
chambeau, J. Slater, M. B. Daly, A. B. Benson, J. M. Kirkwood, P. J. ODwyer, R.  
Sutphen, J. A. Stewart, D. Johnson, M. Nordborg, and T. G. Krontiris, "Strong sig-  
nature of natural selection within an *FHIT* intron implicated in prostate cancer risk,"  
*PLoS ONE*, vol. 3, p. e3533, 2008.
- 78 R. Cagliani, S. Riva, U. Pozzoli, M. Fumagalli, G. P. Comi, N. Bresolin, M. Clerici,  
and M. Sironi, "Balancing selection is common in the extended MHC region but most  
alleles with opposite risk profile for autoimmune diseases are neutrally evolving," *BMC  
Evol. Biol.*, vol. 1, p. 171, 2011.
- 79 A. Hodgkinson and A. Eyre-Walker, "The genomic distribution and local context of  
coincident SNPs in human and chimpanzee," *Genome Biol. Evol.*, vol. 2, pp. 547557,  
2010.