

SNiPACard

Block annotations

Block info


genomic range	chr5:96,123,055-96,151,968 <i>el</i>
block size	28,914 bp
variant count	25 variants

Basic features


Conservation/deleteriousness		Linked genes	
phyloP	$\mu = -0.605$ [-2.841 – 0.73]	gene(s) hit or close-by	AC008906.1 <i>el</i> , CTD-2260A17.1 <i>el</i> , CTD-2260A17.2 <i>el</i> , CTD-2260A17.3 <i>el</i> , ERAP1 <i>el</i>
phastCons	$\mu = 0.067$ [0 – 0.895]	eQTL gene(s)	CTD-2260A17.1 <i>el</i> , CTD-2260A17.3 <i>el</i> , ERAP1 <i>el</i> , ERAP2 <i>el</i> , LNPEP <i>el</i>
GERP++	$\mu = -0.570$ [-10.1 – 2.31]	potentially regulated gene(s)	-
CADD score	$\mu = 2.945$ [0.098 – 9.842]	disease gene(s)	-

Trait annotations

Variant association



trait	min(p-value)	source DB	source entry/link	variant(s)
Ankylosing spondylitis	<5.00×10 ⁻¹²	GWAS Catalog	20062062 	1

Variant annotation

trait	type	source DB	source entry/link	Variant(s)
?	HGMD curated	HGMD	CM0911244 	1

Direct effect on regulation

cis-eQTL

gene	transcript	probe	tissue	min(statistic) (type)	source	variant(s)
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	pancreas	4.21×10 ⁻²³ (p-value)	GTEx Portal V6 	25
CTD-2260A17.1 <i>el</i>	?	ENSG00000248734 <i>el</i>	pancreas	4.98×10 ⁻⁶ (p-value)	GTEx Portal V6 	3
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	muscularis mucosae	2.83×10 ⁻⁸ (p-value)	GTEx Portal V6 	24
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	lung	2.55×10 ⁻³³ (p-value)	GTEx Portal V6 	25
CTD-2260A17.1 <i>el</i>	?	ENSG00000248734 <i>el</i>	lung	8.96×10 ⁻⁹ (p-value)	GTEx Portal V6 	24
CTD-2260A17.3 <i>el</i>	?	ENSG00000272109 <i>el</i>	lung	1.00×10 ⁻¹³ (p-value)	GTEx Portal V6 	25
CTD-2260A17.3 <i>el</i>	?	ENSG00000272109 <i>el</i>	atrial appendage	2.69×10 ⁻⁶ (p-value)	GTEx Portal V6 	6
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	atrial appendage	1.23×10 ⁻⁶ (p-value)	GTEx Portal V6 	5
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	transformed fibroblasts	1.87×10 ⁻³⁸ (p-value)	GTEx Portal V6 	25
CTD-2260A17.1 <i>el</i>	?	ENSG00000248734 <i>el</i>	transformed fibroblasts	2.19×10 ⁻²⁸ (p-value)	GTEx Portal V6 	25
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	blood	1.62×10 ⁻⁴⁵ (p-value)	GTEx Portal V6 	25
CTD-2260A17.1 <i>el</i>	?	ENSG00000248734 <i>el</i>	blood	3.22×10 ⁻¹⁷ (p-value)	GTEx Portal V6 	25
CTD-2260A17.3 <i>el</i>	?	ENSG00000272109 <i>el</i>	blood	4.44×10 ⁻¹¹ (p-value)	GTEx Portal V6 	25
ERAP1 <i>el</i>	?	ENSG00000164307 <i>el</i>	breast	2.06×10 ⁻¹⁰ (p-value)	GTEx Portal V6 	24
CTD-2260A17.3 <i>el</i>	?	ENSG00000272109 <i>el</i>	breast	7.84×10 ⁻⁷ (p-value)	GTEx Portal V6 	21

Gene	ENSG ID	Tissue	p-value	Source	Count
ERAP2	ENSG00000164308	breast	6.95×10 ⁻⁶ (p-value)	GTEX Portal V6	1
ERAP1	ENSG00000164307	tibial artery	6.71×10 ⁻⁸ (p-value)	GTEX Portal V6	24
CTD-2260A17.3	ENSG00000272109	tibial artery	1.01×10 ⁻⁶ (p-value)	GTEX Portal V6	21
ERAP1	ENSG00000164307	thyroid	3.20×10 ⁻¹⁰ (p-value)	GTEX Portal V6	24
CTD-2260A17.3	ENSG00000272109	thyroid	9.13×10 ⁻⁶ (p-value)	GTEX Portal V6	5
ERAP1	ENSG00000164307	skeletal muscle	9.90×10 ⁻²² (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	skeletal muscle	9.45×10 ⁻¹⁵ (p-value)	GTEX Portal V6	25
CTD-2260A17.3	ENSG00000272109	skeletal muscle	1.31×10 ⁻¹¹ (p-value)	GTEX Portal V6	25
ERAP2	ENSG00000164308	skeletal muscle	7.63×10 ⁻⁷ (p-value)	GTEX Portal V6	4
ERAP1	ENSG00000164307	putamen	8.32×10 ⁻⁷ (p-value)	GTEX Portal V6	20
ERAP1	ENSG00000164307	EBV lymphocytes	2.08×10 ⁻¹⁵ (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	EBV lymphocytes	2.47×10 ⁻⁸ (p-value)	GTEX Portal V6	24
ERAP1	ENSG00000164307	transverse colon	1.92×10 ⁻¹⁸ (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	transverse colon	3.63×10 ⁻⁷ (p-value)	GTEX Portal V6	5
ERAP1	ENSG00000164307	liver	1.32×10 ⁻⁸ (p-value)	GTEX Portal V6	25
ERAP1	ENSG00000164307	sun exposed skin	8.36×10 ⁻²⁵ (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	sun exposed skin	5.11×10 ⁻⁸ (p-value)	GTEX Portal V6	24
CTD-2260A17.3	ENSG00000272109	sun exposed skin	4.99×10 ⁻⁸ (p-value)	GTEX Portal V6	24
ERAP1	ENSG00000164307	unexposed skin	1.54×10 ⁻¹⁸ (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	unexposed skin	3.03×10 ⁻⁷ (p-value)	GTEX Portal V6	23
CTD-2260A17.3	ENSG00000272109	unexposed skin	1.23×10 ⁻⁸ (p-value)	GTEX Portal V6	25
ERAP2	ENSG00000164308	unexposed skin	1.00×10 ⁻⁵ (p-value)	GTEX Portal V6	1
ERAP1	ENSG00000164307	left ventricle	4.06×10 ⁻¹¹ (p-value)	GTEX Portal V6	25
ERAP1	ENSG00000164307	subcutaneous adipocytes	4.70×10 ⁻¹⁸ (p-value)	GTEX Portal V6	25
CTD-2260A17.1	ENSG00000248734	subcutaneous adipocytes	5.21×10 ⁻⁸ (p-value)	GTEX Portal V6	24
CTD-2260A17.3	ENSG00000272109	subcutaneous adipocytes	5.24×10 ⁻¹⁰ (p-value)	GTEX Portal V6	25
ERAP1	ENSG00000164307	caudate basal ganglia	8.05×10 ⁻⁷ (p-value)	GTEX Portal V6	20
ERAP1	ENSG00000164307	stomach	2.22×10 ⁻¹⁵ (p-value)	GTEX Portal V6	24
ERAP1	ENSG00000164307	visceral adipocytes	2.96×10 ⁻¹⁰ (p-value)	GTEX Portal V6	25
ERAP2	ENSG00000164308	visceral adipocytes	1.33×10 ⁻⁹ (p-value)	GTEX Portal V6	24
CTD-2260A17.3	ENSG00000272109	visceral adipocytes	3.80×10 ⁻⁸ (p-value)	GTEX Portal V6	25
ERAP1	ENSG00000164307	spleen	2.61×10 ⁻⁷ (p-value)	GTEX Portal V6	16
ERAP1	ENSG00000164307	coronary artery	1.88×10 ⁻⁷ (p-value)	GTEX Portal V6	23
ERAP1	ENSG00000164307	sigmoid colon	7.72×10 ⁻¹³ (p-value)	GTEX Portal V6	25
ERAP1	ENSG00000164307	tibial nerve	3.34×10 ⁻²⁰ (p-value)	GTEX Portal V6	25

CTD-2260A17.3 <i>e!</i> ?	ENSG00000272109 <i>e!</i>	tibial nerve	1.80×10 ⁻⁶ (p-value)	GTEx Portal V6 <i>!M</i>	23	
ERAP1 <i>e!</i> ?	ENSG00000164307 <i>e!</i>	esophagus mucosa	2.25×10 ⁻²⁵ (p-value)	GTEx Portal V6 <i>!M</i>	25	
CTD-2260A17.1 <i>e!</i> ?	ENSG00000248734 <i>e!</i>	esophagus mucosa	2.32×10 ⁻⁷ (p-value)	GTEx Portal V6 <i>!M</i>	24	
CTD-2260A17.3 <i>e!</i> ?	ENSG00000272109 <i>e!</i>	esophagus mucosa	1.50×10 ⁻¹⁰ (p-value)	GTEx Portal V6 <i>!M</i>	25	
ERAP1 <i>e!</i> ?	ENSG00000164307 <i>e!</i>	gastroesophageal junction	6.19×10 ⁻⁷ (p-value)	GTEx Portal V6 <i>!M</i>	23	
ERAP1 <i>e!</i> ?	ENSG00000164307 <i>e!</i>	adrenal gland	4.65×10 ⁻¹³ (p-value)	GTEx Portal V6 <i>!M</i>	25	
CTD-2260A17.1 <i>e!</i> ?	ENSG00000248734 <i>e!</i>	adrenal gland	1.73×10 ⁻⁶ (p-value)	GTEx Portal V6 <i>!M</i>	11	
CTD-2260A17.3 <i>e!</i> ?	ENSG00000272109 <i>e!</i>	testis	1.49×10 ⁻¹⁴ (p-value)	GTEx Portal V6 <i>!M</i>	25	
ERAP1 <i>e!</i> ?	ENSG00000164307 <i>e!</i>	frontal cortex	1.45×10 ⁻⁷ (p-value)	GTEx Portal V6 <i>!M</i>	23	
ERAP2 <i>e!</i>	ENST00000513084 <i>e!</i>	ILMN_1743145 <i>e!</i>	skin	1.67×10 ⁻⁹ (p-value)	MuTHER consortium <i>!M</i>	9
ERAP2 <i>e!</i>	ENST00000512869 <i>e!</i>		blood	7.98×10 ⁻¹¹ (p-value)	MuTHER consortium <i>!M</i>	9
ERAP2 <i>e!</i>	ENST00000379904 <i>e!</i>		adipocyte	1.05×10 ⁻⁹ (p-value)	MuTHER consortium <i>!M</i>	9
ERAP2 <i>e!</i>	ENST00000437043 <i>e!</i>					
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>	ILMN_1752145 <i>e!</i>	blood	1.25×10 ⁻⁵ (p-value)	MuTHER consortium <i>!M</i>	1
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>	214012_at <i>e!</i>	blood	6.80×10 ⁻⁸ (p-value)	Dixon et al. <i>!M</i>	2
ERAP1 <i>e!</i>	ENST00000443439 <i>e!</i>	209788_s_at <i>e!</i>	blood	2.70×10 ⁻³² (p-value)	Dixon et al. <i>!M</i>	2
ERAP2 <i>e!</i>	ENST00000513084 <i>e!</i>	ILMN_1743145 <i>e!</i>	monocyte	3.89×10 ⁻⁷ (p-value)	Fairfax et al. <i>!M</i>	2
ERAP2 <i>e!</i>	ENST00000512869 <i>e!</i>		b-cell	1.82×10 ⁻⁷ (p-value)	Fairfax et al. <i>!M</i>	2
ERAP2 <i>e!</i>	ENST00000379904 <i>e!</i>					
ERAP2 <i>e!</i>	ENST00000437043 <i>e!</i>					
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>	ILMN_1752145 <i>e!</i>	monocyte	3.32×10 ⁻⁶ (p-value)	Fairfax et al. <i>!M</i>	2
ERAP2 <i>e!</i> ?	ENSG00000164308 <i>e!</i>	blood	2.05×10 ⁻⁴ (q-value)	SeeQTL DB (HapMap) <i>!M</i>	3	
LNPEP <i>e!</i>	ENST00000395770 <i>e!</i>	ILMN_1814737 <i>e!</i>	blood	7.60×10 ⁻⁶ (p-value)	Westra et al. <i>!M</i>	9
LNPEP <i>e!</i>	ENST00000231368 <i>e!</i>					
CTD-2260A17.3 <i>e!</i> ?	ENSG00000272109 <i>e!</i>	aorta	1.22×10 ⁻⁵ (p-value)	GTEx Portal V6 <i>!M</i>	2	
ERAP2 <i>e!</i>	ENST00000513084 <i>e!</i>	ILMN_1743145 <i>e!</i>	monocyte	6.82×10 ⁻²³ (p-value)	Zeller et al. <i>!M</i>	2
ERAP2 <i>e!</i>	ENST00000512869 <i>e!</i>					
ERAP2 <i>e!</i>	ENST00000379904 <i>e!</i>					
ERAP2 <i>e!</i>	ENST00000437043 <i>e!</i>					
ERAP1 <i>e!</i>	ENST00000443439 <i>e!</i>	ILMN_2336220 <i>e!</i>	monocyte	1.80×10 ⁻²³ (p-value)	Zeller et al. <i>!M</i>	2
ERAP1 <i>e!</i>	ENST00000514604 <i>e!</i>					
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>					

Putative effect on regulation

Regulatory feature cluster

element id	variant(s)	tissue/cell	factors
ENSR00001412052 <i>e!</i> (enhancer)	1	blood (K562)	Max
ENSR00001285989 <i>e!</i> (promoter flanking region)	1	monocytes (Monocytes-CD14+)	H3K4me3
		embryonic stem cell (H1ESC)	DNase1
		blood (GM12878)	H3K27ac, H3K4me2, H3K9ac, H3K4me3, DNase1, PolII, H2AZ, H3K4me1
		blood (K562)	H2AZ
		A549	H3K27me3

blood (DND-41) H3K9ac, H3K27ac, H3K4me1
 skin (NHDF-AD) DNase1

Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
AC008906.1 <i>e!</i>	upstream gene variant, downstream gene variant	7	ENST00000629418 ? <i>e!</i>		?	9
CTD-2260A17.1 <i>e!</i>	downstream gene variant	1353	ENST00000602972 ? <i>e!</i>		?	6
CTD-2260A17.1 <i>e!</i>	downstream gene variant	1353	ENST00000512856 ? <i>e!</i>		?	6
CTD-2260A17.2 <i>e!</i>	downstream gene variant	1954	ENST00000501338 ? <i>e!</i>		?	1
CTD-2260A17.2 <i>e!</i>	downstream gene variant	2100	ENST00000502262 ? <i>e!</i>		?	1
CTD-2260A17.2 <i>e!</i>	downstream gene variant	1958	ENST00000504056 ? <i>e!</i>		?	1
CTD-2260A17.3 <i>e!</i>	upstream gene variant	1195	ENST00000606656 ? <i>e!</i>		?	7
CTD-2260A17.3 <i>e!</i>	upstream gene variant	2509	ENST00000606346 ? <i>e!</i>		?	7
ERAP1 <i>e!</i>	downstream gene variant, upstream gene variant	1217	ENST00000503921 ? <i>e!</i>		ENSP00000427025 <i>e!</i>	6
ERAP1 <i>e!</i>	downstream gene variant, upstream gene variant	2524	ENST00000508227 ? <i>e!</i>		ENSP00000422631 <i>e!</i>	6
ERAP1 <i>e!</i>	upstream gene variant	592	ENST00000514604 ? <i>e!</i>		?	6
ERAP1 <i>e!</i>	upstream gene variant	4165	ENST00000443439 <i>e!</i>	NM_001040458.1, NM_001198541.1	ENSP00000406304 <i>e!</i>	1
ERAP1 <i>e!</i>	upstream gene variant	1576	ENST00000507859 ? <i>e!</i>		?	5
ERAP1 <i>e!</i>	downstream gene variant, upstream gene variant	138	ENST00000503311 ? <i>e!</i>		?	14
ERAP1 <i>e!</i>	downstream gene variant, upstream gene variant	382	ENST00000507154 ? <i>e!</i>		ENSP00000421697 <i>e!</i>	8
ERAP1 <i>e!</i>	upstream gene variant	3974	ENST00000296754 <i>e!</i>	NM_016442.3	ENSP00000296754 <i>e!</i>	1

Putative effect on transcript

Synonymous coding variant

gene	affected transcript	RefSeq id	protein	AA's	exchanged codons	variant(s)
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>	NM_016442.3	ENSP00000296754 <i>e!</i>	A	gcT/gcC	1
ERAP1 <i>e!</i>	ENST00000443439 <i>e!</i>	NM_001040458.1, NM_001198541.1	ENSP00000406304 <i>e!</i>	A	gcT/gcC	1

Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
CTD-2260A17.2 <i>e!</i>	ENST00000502262 <i>e!</i>	?	?	1
CTD-2260A17.2 <i>e!</i>	ENST00000504056 <i>e!</i>	?	?	1
CTD-2260A17.2 <i>e!</i>	ENST00000501338 <i>e!</i>	?	?	1
ERAP1 <i>e!</i>	ENST00000296754 <i>e!</i>	NM_016442.3	ENSP00000296754 <i>e!</i>	22
ERAP1 <i>e!</i>	ENST00000508227 <i>e!</i>	?	ENSP00000422631 <i>e!</i>	7
ERAP1 <i>e!</i>	ENST00000507859 <i>e!</i>	?	?	6
ERAP1 <i>e!</i>	ENST00000503311 <i>e!</i>	?	?	2
ERAP1 <i>e!</i>	ENST00000443439 <i>e!</i>	NM_001040458.1, NM_001198541.1	ENSP00000406304 <i>e!</i>	22
ERAP1 <i>e!</i>	ENST00000503921 <i>e!</i>	?	ENSP00000427025 <i>e!</i>	11

Non-coding exon variant

gene	affected transcript	RefSeq id	variant(s)
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