

Block annotations

Block info










genomic range	chr6:31,377,642-31,391,257 <i>el</i>
block size	13,616 bp
variant count	13 variants

Basic features

Conservation/deleteriousness		Linked genes	
phyloP	$\mu = -0.284$ [-1.521 – 0.426]	gene(s) hit or close-by	HCP5 <i>el</i> , MICA <i>el</i>
phastCons	$\mu = 0.057$ [0 – 0.409]	eQTL gene(s)	CCHCR1 <i>el</i> , HLA-B <i>el</i> , MICA <i>el</i> , PSORS1C1 <i>el</i> , PSORS1C2 <i>el</i> , Y_RNA <i>el</i>
GERP++	$\mu = -0.195$ [-1.75 – 0.536]	potentially regulated gene(s)	MICA <i>el</i> , MICA <i>el</i> , MICA <i>el</i> , MICA <i>el</i> , MICA <i>el</i> , VARS2 <i>el</i> , VARS2 <i>el</i> , VARS2 <i>el</i> , VARS2 <i>el</i> , VARS2 <i>el</i> , VARS2 <i>el</i>
CADD score	$\mu = 2.861$ [0.826 – 9.209]	disease gene(s)	<i>el</i> , <i>el</i> , <i>el</i> , VARS2 <i>el</i> , HLA-B <i>el</i>












Trait annotations

Disease gene annotation

gene	trait	source DB	source entry/link
<i>el</i>	PSORIASIS 1, SUSCEPTIBILITY TO	OMIM	MIM:177900 
<i>el</i>	PSORIATIC ARTHRITIS, SUSCEPTIBILITY TOPSORIATIC ARTHRITIS, SUSCEPTIBILITY [...]	OMIM	MIM:607507 
<i>el</i>	COMBINED OXIDATIVE PHOSPHORYLATION DEFICIENCY 20	OMIM	MIM:615917 
<i>el</i>	PSORIASIS 1, SUSCEPTIBILITY TO	OMIM	MIM:177900 
<i>el</i>	PSORIATIC ARTHRITIS, SUSCEPTIBILITY TOPSORIATIC ARTHRITIS, SUSCEPTIBILITY [...]	OMIM	MIM:607507 
VARS2 <i>el</i>	COMBINED OXIDATIVE PHOSPHORYLATION DEFICIENCY 20	OMIM	MIM:615917 
HLA-B <i>el</i>	Takayasu arteritis	OrphaNet	OrphaNet:3287 
HLA-B <i>el</i>	Stevens-Johnson syndrome	OrphaNet	OrphaNet:36426 
HLA-B <i>el</i>	Behcet disease	OrphaNet	OrphaNet:117 

Direct effect on regulation

cis-eQTL

gene	transcript	probe	tissue	min(statistic) (type)	source	variant(s)
MICA <i>el</i>	?	ENSG00000204520 <i>el</i>	muscularis mucosae	6.76×10 ⁻⁸ (p-value)	GTEEx Portal V6 	1
MICA <i>el</i>	?	ENSG00000204520 <i>el</i>	lung	6.56×10 ⁻¹² (p-value)	GTEEx Portal V6 	12
PSORS1C2 <i>el</i>	?	ENSG00000204538 <i>el</i>	lung	8.23×10 ⁻⁷ (p-value)	GTEEx Portal V6 	1
PSORS1C1 <i>el</i>	?	ENSG00000204540 <i>el</i>	lung	7.32×10 ⁻⁷ (p-value)	GTEEx Portal V6 	1
Y_RNA <i>el</i>	?	ENSG00000199332 <i>el</i>	lung	2.71×10 ⁻⁸ (p-value)	GTEEx Portal V6 	11
MICA <i>el</i>	?	ENSG00000204520 <i>el</i>	atrial appendage	1.02×10 ⁻⁸ (p-value)	GTEEx Portal V6 	11
MICA <i>el</i>	?	ENSG00000204520 <i>el</i>	transformed fibroblasts	1.36×10 ⁻¹² (p-value)	GTEEx Portal V6 	11
HLA-B <i>el</i>	?	ENSG00000234745 <i>el</i>	transformed fibroblasts	5.82×10 ⁻⁶ (p-value)	GTEEx Portal V6 	1
Y_RNA <i>el</i>	?	ENSG00000199332 <i>el</i>	transformed fibroblasts	1.00×10 ⁻⁹ (p-value)	GTEEx Portal V6 	12
MICA <i>el</i>	?	ENSG00000204520 <i>el</i>	tibial artery	7.30×10 ⁻¹¹ (p-value)	GTEEx Portal V6 	11
Y_RNA <i>el</i>	?	ENSG00000199332 <i>el</i>	tibial artery	2.46×10 ⁻¹² (p-value)	GTEEx Portal V6 	12

MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	breast	2.71×10 ⁻¹⁰ (p-value)	GTEEx Portal V6 <i>!m</i>	10
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	blood	4.42×10 ⁻¹¹ (p-value)	GTEEx Portal V6 <i>!m</i>	11
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	thyroid	5.40×10 ⁻¹² (p-value)	GTEEx Portal V6 <i>!m</i>	12
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	skeletal muscle	1.18×10 ⁻¹² (p-value)	GTEEx Portal V6 <i>!m</i>	12
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	EBV lymphocytes	2.60×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	1
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	EBV lymphocytes	3.18×10 ⁻⁷ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	sun exposed skin	3.28×10 ⁻¹¹ (p-value)	GTEEx Portal V6 <i>!m</i>	11
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	aorta	1.80×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	11
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	aorta	1.92×10 ⁻¹⁰ (p-value)	GTEEx Portal V6 <i>!m</i>	11
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	left ventricle	5.23×10 ⁻⁹ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	subcutaneous adipocytes	3.15×10 ⁻⁹ (p-value)	GTEEx Portal V6 <i>!m</i>	10
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	subcutaneous adipocytes	3.69×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	caudate basal ganglia	5.43×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	visceral adipocytes	2.59×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	tibial nerve	9.72×10 ⁻¹⁴ (p-value)	GTEEx Portal V6 <i>!m</i>	11
CCHCR1 <i>e!</i>	?	ENSG00000204536 <i>e!</i>	tibial nerve	2.40×10 ⁻⁶ (p-value)	GTEEx Portal V6 <i>!m</i>	1
PSORS1C1 <i>e!</i>	?	ENSG00000204540 <i>e!</i>	tibial nerve	9.33×10 ⁻⁶ (p-value)	GTEEx Portal V6 <i>!m</i>	1
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	tibial nerve	1.35×10 ⁻⁸ (p-value)	GTEEx Portal V6 <i>!m</i>	4
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	sigmoid colon	1.96×10 ⁻⁶ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	esophagus mucosa	1.52×10 ⁻¹⁰ (p-value)	GTEEx Portal V6 <i>!m</i>	1
Y_RNA <i>e!</i>	?	ENSG00000199332 <i>e!</i>	esophagus mucosa	3.49×10 ⁻⁶ (p-value)	GTEEx Portal V6 <i>!m</i>	1
MICA <i>e!</i>	?	ENSG00000204520 <i>e!</i>	testis	2.92×10 ⁻⁷ (p-value)	GTEEx Portal V6 <i>!m</i>	1

Putative effect on regulation

ENCODE promoter-associated distal DHS (Enhancer)

SNiPA enhancer id	variant(s)	associated SNiPA promoter id	associated gene(s)
ENCE00000444713 <i>e!</i>	1	ENCP00000048015	MICA <i>e!</i> MICA <i>e!</i> MICA <i>e!</i> MICA <i>e!</i> MICA <i>e!</i>
		ENCP00000047968	VARS2 <i>e!</i> VARS2 <i>e!</i> VARS2 <i>e!</i> VARS2 <i>e!</i> VARS2 <i>e!</i> VARS2 <i>e!</i>

Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
MICA <i>e!</i>	downstream gene variant	39	ENST00000449934 <i>e!</i>	NM_001177519.2	ENSP00000413079 <i>e!</i>	4

MICA <i>e!</i>	downstream gene variant	41	ENST00000616296 <i>e!</i>	NM_001289152.1, NM_001289153.1, NM_001289154.1	ENSP00000482382 <i>e!</i>	4
MICA <i>e!</i>	downstream gene variant	39	ENST00000421350 <i>e!</i>	?	ENSP00000402410 <i>e!</i>	4

Putative effect on transcript

Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
HCP5 <i>e!</i>	ENST00000414046 <i>e!</i>	?	?	13
MICA <i>e!</i>	ENST00000616296 <i>e!</i>	NM_001289152.1, NM_001289153.1, NM_001289154.1	ENSP00000482382 <i>e!</i>	1
MICA <i>e!</i>	ENST00000449934 <i>e!</i>	NM_001177519.2	ENSP00000413079 <i>e!</i>	1
MICA <i>e!</i>	ENST00000421350 <i>e!</i>	?	ENSP00000402410 <i>e!</i>	1

