

# SNiPacard

## Block annotations

### Block info

genomic range	chr7:143,092,918-143,108,081 <i>e!</i>
block size	15,164 bp
variant count	10 variants

### Basic features

Conservation/deleteriousness		Linked genes	
phyloP	$\mu = -0.574$ [-1.838 – 0.994]	gene(s) hit or close-by	EPHA1 <i>e!</i> , EPHA1-AS1 <i>e!</i> , ZYX <i>e!</i>
phastCons	$\mu = 0.026$ [0 – 0.13]	eQTL gene(s)	EPHA1 <i>e!</i> , EPHA1-AS1 <i>e!</i> , ZYX <i>e!</i>
GERP++	$\mu = 0.477$ [-1.57 – 3.88]	potentially regulated gene(s)	-
CADD score	$\mu = 5.976$ [2.371 – 12.8]	disease gene(s)	-

## Direct effect on transcript

### Amino acid sequence alteration

gene	effect type	affected transcript	RefSeq id	protein	exchanged AA's	exchanged codons	SIFT prediction	PolyPhen prediction	variant(s)
EPHA1 <i>e!</i>	missense variant	ENST00000275815 <i>e!</i>	NM_005232.4	ENSP00000275815	V/A	gTg/gCg	tolerated	benign	1

## Direct effect on regulation

### cis-eQTL

gene	transcript	probe	tissue	min(statistic) (type)	source	variant(s)
EPHA1-AS1 <i>e!</i>	?	ENSG00000229153 <i>e!</i>	blood	3.02×10 <sup>-8</sup> (p-value)	GTEx Portal V6 <i>!M</i>	2
ZYX <i>e!</i>	?	ENSG00000159840 <i>e!</i>	blood	2.45×10 <sup>-8</sup> (p-value)	GTEx Portal V6 <i>!M</i>	3
EPHA1 <i>e!</i>	?	ENSG00000146904 <i>e!</i>	sun exposed skin	2.87×10 <sup>-7</sup> (p-value)	GTEx Portal V6 <i>!M</i>	8
EPHA1 <i>e!</i>	?	ENSG00000146904 <i>e!</i>	thyroid	6.98×10 <sup>-7</sup> (p-value)	GTEx Portal V6 <i>!M</i>	8

## Putative effect on regulation

### Regulatory feature cluster

element id	variant(s)	tissue/cell	factors
ENSR00001564739 <i>e!</i> (enhancer)	2	embryonic stem cell (H1ESC) blood (K562) skin (NHDF-AD) breast (HMEC) cervix (HeLa-S3) endothelium (HUVEC) liver (HepG2) lung (IMR90) skin (NHEK)	H3K36me3 CTCF, H3K27me3 DNase1, CTCF H3K36me3 DNase1 DNase1, H3K4me1, PolII H3K4me1, H3K4me2, HNF4A, DNase1, RXRA H3K36me3 H3K36me3
ENSR00000069717 <i>e!</i> (CTCF binding site)	2	embryonic stem cell (H1ESC) blood (K562) skin (NHDF-AD) breast (HMEC) endothelium (HUVEC)	CTCF, Rad21, H3K36me3 CTCF, H3K27me3 CTCF H3K36me3 DNase1, PolII, H3K4me1

		liver (HepG2)	RXRA, H3K4me1, H3K4me2, HNF4A, DNase1
		lung (IMR90)	H3K36me3
		skin (NHEK)	H3K36me3
ENSR00000635676 <i>e!</i>	2	embryonic stem cell (H1ESC)	H3K36me3, DNase1
(open chromatin region)		lung (IMR90)	H3K27me3
		A549	H3K27me3
		blood (K562)	H3K27me3
		skin (NHEK)	H3K36me3
ENSR00001718722 <i>e!</i>	2	embryonic stem cell (H1ESC)	DNase1, H3K4me3, PolII, Rad21, Egr1, Nrsf, CTCF, TAF1, SP1
(promoter flanking region)		HSMMtube	H3K27me3
		blood (K562)	H3K27me3, Egr1, CTCF, DNase1
		breast (HMEC)	CTCF, H3K4me3
		cervix (HeLa-S3)	CTCF
		monocytes (Monocytes-CD14+)	DNase1, H3K4me1, H3K27ac, H3K4me3
		endothelium (HUVEC)	CTCF
		liver (HepG2)	H3K4me3, H3K4me2, DNase1
		blood (GM12878)	ZEB1
		lung (IMR90)	CTCF
		nervous (NH-A)	H3K27me3
		A549	H3K4me3, H3K4me2, H3K27me3
		skin (NHEK)	DNase1, CTCF, H3K4me3
ENSR00000635678 <i>e!</i>	2	monocytes (Monocytes-CD14+)	H3K4me3, H3K27ac, H3K4me1, DNase1
(promoter flanking region)		embryonic stem cell (H1ESC)	H3K4me2, DNase1
		Osteobl	H3K4me2
		blood (K562)	H3K27me3
		skin (NHDF-AD)	DNase1

### Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
EPHA1 <i>e!</i>	upstream gene variant	4434	ENST00000458129 ? <i>e!</i>		?	2
EPHA1 <i>e!</i>	downstream gene variant, upstream gene variant	819	ENST00000497891 ? <i>e!</i>		?	6
EPHA1 <i>e!</i>	downstream gene variant, upstream gene variant	1267	ENST00000479459 ? <i>e!</i>		?	4
EPHA1 <i>e!</i>	upstream gene variant	224	ENST00000465208 ? <i>e!</i>		?	3
EPHA1 <i>e!</i>	upstream gene variant	1041	ENST00000275815 ? <i>e!</i>	NM_005232.4	ENSP00000275815 <i>e!</i>	4
EPHA1 <i>e!</i>	upstream gene variant	1128	ENST00000488068 ? <i>e!</i>		?	4
EPHA1 <i>e!</i>	upstream gene variant	3326	ENST00000494989 ? <i>e!</i>		?	2
EPHA1-AS1 <i>e!</i>	upstream gene variant	1526	ENST00000429289 ? <i>e!</i>		?	2
EPHA1-AS1 <i>e!</i>	upstream gene variant	1546	ENST00000421648 ? <i>e!</i>		?	2
ZYX <i>e!</i>	downstream gene variant	4716	ENST00000354434 ? <i>e!</i>		ENSP00000346417 <i>e!</i>	1
ZYX <i>e!</i>	downstream gene variant	4714	ENST00000322764 ? <i>e!</i>	NM_003461.4, NM_001010972.1	ENSP00000324422 <i>e!</i>	1
ZYX <i>e!</i>	downstream gene variant	4716	ENST00000392910 ? <i>e!</i>		ENSP00000376642 <i>e!</i>	1

### Putative effect on transcript

#### Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
EPHA1 <i>e!</i>	ENST00000497891 <i>e!</i>	?	?	3
EPHA1 <i>e!</i>	ENST00000488068 <i>e!</i>	?	?	5
EPHA1 <i>e!</i>	ENST00000275815 <i>e!</i>	NM_005232.4	ENSP00000275815 <i>e!</i>	5
EPHA1 <i>e!</i>	ENST00000494989 <i>e!</i>	?	?	2

EPHA1-AS1 <i>e!</i>	ENST00000421648 <i>e!</i>	?	?	4
EPHA1-AS1 <i>e!</i>	ENST00000429289 <i>e!</i>	?	?	4

### Non-coding exon variant

gene	affected transcript	RefSeq id	variant(s)
EPHA1 <i>e!</i>	ENST00000497891 <i>e!</i>	?	1
EPHA1 <i>e!</i>	ENST00000488068 <i>e!</i>	?	2

