

SNiPACard

Block annotations

Block info

genomic range	chr4:74,701,512-74,703,999 <i>e!</i>
block size	2,488 bp
variant count	2 variants

Basic features

Conservation/deleteriousness		Linked genes	
phyloP	$\mu = 0.118$ [0.075 – 0.161]	gene(s) hit or close-by	CXCL6 <i>e!</i>
phastCons	$\mu = 0.519$ [0.04 – 0.999]	eQTL gene(s)	CXCL6 <i>e!</i> , PF4V1 <i>e!</i>
GERP++	$\mu = 1.190$ [0 – 2.38]	potentially regulated gene(s)	-
CADD score	$\mu = 7.585$ [1.391 – 13.78]	disease gene(s)	-

Direct effect on regulation

cis-eQTL

gene	transcript	probe	tissue	min(statistic) (type)	source	variant(s)
CXCL6 <i>e!</i>	?	ENSG00000124875 <i>e!</i>	tibial nerve	2.03×10 ⁻⁷ (p-value)	GTEx Portal V6 <i>!m</i>	2
CXCL6 <i>e!</i>	?	ENSG00000124875 <i>e!</i>	pancreas	5.79×10 ⁻⁷ (p-value)	GTEx Portal V6 <i>!m</i>	1
PF4V1 <i>e!</i>	ENST00000226524 <i>e!</i>	ILMN_1745522 <i>e!</i>	monocyte	1.91×10 ⁻¹⁷ (p-value)	Zeller et al. <i>!m</i>	1

Putative effect on regulation

Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
CXCL6 <i>e!</i>	upstream gene variant	1392	ENST00000503446 <i>e!</i>	?	?	1
CXCL6 <i>e!</i>	upstream gene variant, downstream gene variant	845	ENST00000515050 <i>e!</i>	?	ENSP00000424819 <i>e!</i>	2
CXCL6 <i>e!</i>	upstream gene variant	702	ENST00000226317 <i>e!</i>	NM_002993.3	ENSP00000226317 <i>e!</i>	1

Putative effect on transcript

Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
CXCL6 <i>e!</i>	ENST00000503446 <i>e!</i>	?	?	1

3'-UTR variant

gene	affected transcript	RefSeq id	protein	variant(s)
CXCL6 <i>e!</i>	ENST00000226317 <i>e!</i>	NM_002993.3	ENSP00000226317 <i>e!</i>	1

