

## Block annotations

### Block info

genomic range	chr3:186,335,941-186,341,830 <i>e!</i>
block size	5,890 bp
variant count	6 variants

### Basic features

Conservation/deleteriousness		Linked genes	
phyloP	$\mu = -0.386$ [-1.316 – 0.255]	gene(s) hit or close-by	AHSG <i>e!</i> , RP11-573D15.8 <i>e!</i>
phastCons	$\mu = 0.000$ [0 – 0.001]	eQTL gene(s)	-
GERP++	$\mu = 0.464$ [-3.04 – 3.81]	potentially regulated gene(s)	-
CADD score	$\mu = 3.943$ [0.153 – 11.52]	disease gene(s)	-

## Trait annotations

### Variant annotation

trait	type	source DB	source entry/link	Variant(s)
Calcium oxalate urolithiasis	association	ClinVar	RCV000128583.1 <i>ClinVar</i>	1

## Putative effect on regulation

### Regulatory feature cluster

element id	variant(s)	tissue/cell	factors
ENSR00001486982 <i>e!</i> (promoter flanking region)	2	embryonic stem cell (H1ESC)	H3K27me3, Rad21, CTCF, Yy1, DNase1
		HSMMtube	DNase1, H3K27me3, CTCF
		blood (K562)	CTCF, Rad21, H3K27me3
		skin (NHDF-AD)	H3K4me2, DNase1, H3K4me3, CTCF
		muscle (HSMM)	H3K4me2, CTCF, H2AZ, DNase1
		liver (HepG2)	DNase1, H3K36me3, CTCF, Rad21, PolII, H4K20me1
		blood (GM12878)	DNase1, H2AZ, Rad21, H3K4me2, CTCF
		lung (IMR90)	DNase1, H3K4me2, H3K27me3, CTCF
		nervous (NH-A)	CTCF, H3K4me2, DNase1
		skin (NHEK)	DNase1, CTCF, H3K4me3, H3K4me2
		NHLF	CTCF, DNase1
		Osteobl	H2AZ, H3K4me2, CTCF, H3K27me3
		blood (DND-41)	H3K27me3, CTCF
		breast (HMEC)	CTCF, H3K4me2, DNase1
		cervix (HeLa-S3)	Max, CTCF, H3K27me3, DNase1, H3K4me2
		monocytes (Monocytes-CD14+)	CTCF, H3K27me3
		endothelium (HUVEC)	H3K36me3, CTCF, H3K27me3, DNase1
		A549	H3K27me3, DNase1, CTCF, H3K4me3, H3K4me2

### Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
AHSG <i>e!</i>	downstream gene variant	163	ENST00000478441 <i>e!</i>	?	?	3
AHSG <i>e!</i>	downstream gene variant	2286	ENST00000411641 <i>e!</i>	NM_001622.2	ENSP00000393887 <i>e!</i>	3
AHSG <i>e!</i>	downstream gene variant	2304	ENST00000273784 <i>e!</i>	?	ENSP00000273784 <i>e!</i>	3

## Putative effect on transcript

### Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
AHSG <i>e!</i>	ENST00000273784 <i>e!</i>	?	ENSP00000273784 <i>e!</i>	3
AHSG <i>e!</i>	ENST00000411641 <i>e!</i>	NM_001622.2	ENSP00000393887 <i>e!</i>	3
RP11-573D15.8 <i>e!</i>	ENST00000625386 <i>e!</i>	?	?	6
RP11-573D15.8 <i>e!</i>	ENST00000630178 <i>e!</i>	?	?	6
RP11-573D15.8 <i>e!</i>	ENST00000628505 <i>e!</i>	?	?	6

