

Block annotations

Block info

genomic range	chr17:64,195,431-64,305,051 <i>e!</i>
block size	109,621 bp
variant count	5 variants

Basic features


Conservation/deleteriousness		Linked genes	
phyloP	$\mu = 1.057$ [-1.088 – 6.439]	gene(s) hit or close-by	APOH <i>e!</i> , PRKCA <i>e!</i> , PSMD7P1 <i>e!</i>
phastCons	$\mu = 0.241$ [0 – 1]	eQTL gene(s)	-
GERP++	$\mu = 1.266$ [-2.42 – 5.5]	potentially regulated gene(s)	-
CADD score	$\mu = 8.965$ [0.147 – 26.9]	disease gene(s)	-

Trait annotations

Variant association

trait	min(p-value)	source DB	source entry/link	variant(s)
LDL cholesterol	<1.00×10 ⁻¹¹	GWAS Catalog	24097068 	1

Variant annotation

trait	type	source DB	source entry/link	Variant(s)
?	HGMD curated	HGMD	CM970094 	1

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)
APOH <i>e!</i>	missense variant	ENST00000205948 <i>e!</i>	NM_000042.2	ENSP00000205948	G/C	Ggc/Tgc	?	?	1

Putative effect on regulation

Regulatory feature cluster

element id	variant(s)	tissue/cell	factors
ENSR00001537522 <i>e!</i>	1	embryonic stem cell (H1ESC)	DNase1, H3K27me3
(promoter)		HSMMtube	H3K27me3
		Osteobl	H3K27me3
		blood (K562)	H3K27me3
		blood (DND-41)	H3K27me3
		cervix (HeLa-S3)	DNase1
		monocytes (Monocytes-CD14+)	H3K27me3
		liver (HepG2)	FOSL2, Jund, PolII, H3K79me2, DNase1, H3K4me1, p300, H3K4me2, FOXA1, H3K9ac, H3K27ac, H3K4me3, H3K27me3, H3K36me3
		lung (IMR90)	H3K27me3
		A549	H3K4me3, DNase1

Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
APOH <i>e!</i>	downstream gene variant, upstream gene variant	20	ENST00000581797 <i>e!</i>	?	ENSP00000463553 <i>e!</i>	3

APOH <i>e!</i>	upstream gene variant	3460	ENST00000205948 <i>e!</i>	NM_000042.2	ENSP00000205948 <i>e!</i>	1
PRKCA <i>e!</i>	downstream gene variant	2585	ENST00000583775 <i>e!</i>	?	?	1
PSMD7P1 <i>e!</i>	downstream gene variant	1075	ENST00000579375 <i>e!</i>	?	?	1

Putative effect on transcript

Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
APOH <i>e!</i>	ENST00000205948 <i>e!</i>	NM_000042.2	ENSP00000205948 <i>e!</i>	1
APOH <i>e!</i>	ENST00000577982 <i>e!</i>	?	ENSP00000464301 <i>e!</i>	2
APOH <i>e!</i>	ENST00000585162 <i>e!</i>	?	ENSP00000462260 <i>e!</i>	1
PRKCA <i>e!</i>	ENST00000284384 <i>e!</i>	?	ENSP00000284384 <i>e!</i>	1
PRKCA <i>e!</i>	ENST00000578063 <i>e!</i>	?	ENSP00000462087 <i>e!</i>	1
PRKCA <i>e!</i>	ENST00000413366 <i>e!</i>	NM_002737.2	ENSP00000408695 <i>e!</i>	1
PRKCA <i>e!</i>	ENST00000583361 <i>e!</i>	?	?	1

