

SNiPAcad

Block annotations

Block info



genomic range	chr3:42,906,116-42,906,116 <i>e!</i>
block size	1 bp
variant count	1 variants

Basic features

Conservation/deleteriousness		Linked genes	
phyloP	2.284	gene(s) hit or close-by	ACKR2 <i>e!</i> , CYP8B1 <i>e!</i> , KRBOX1 <i>e!</i> , RP11-136C24.3 <i>e!</i> , RP11-141M3.6 <i>e!</i>
phastCons	0.985	eQTL gene(s)	-
GERP++	5.52	potentially regulated gene(s)	-
CADD score	10.24	disease gene(s)	-

Trait annotations

Variant association

trait	min(p-value)	source DB	source entry/link	variant(s)
Cerebrospinal fluid levels of Alzheimer's disease-related proteins	<4.00×10 ⁻¹⁸	GWAS Catalog	25340798 	1
Monocyte count	<2.00×10 ⁻⁷	GWAS Catalog	23314186 	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)
ACKR2 <i>e!</i>	missense variant	ENST00000493193 <i>e!</i>	?	ENSP00000476581	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000442925 <i>e!</i>	?	ENSP00000396150	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000497921 <i>e!</i>	?	ENSP00000476901	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000494619 <i>e!</i>	?	ENSP00000477157	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000422265 <i>e!</i>	NM_001296.4	ENSP00000416996	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000273145 <i>e!</i>	?	ENSP00000273145	A/V	gCg/gTg	?	?	1
ACKR2 <i>e!</i>	missense variant	ENST00000492609 <i>e!</i>	?	ENSP00000477475	A/V	gCg/gTg	?	?	1

Putative effect on regulation

Variation proximal to gene

gene	variant type	min(distance)	transcript	RefSeq id	protein	variant(s)
ACKR2 <i>e!</i>	downstream gene variant	90	ENST00000463699 <i>e!</i>	?	?	1

Putative effect on transcript

Intron variant

gene	affected transcript	RefSeq id	protein	variant(s)
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CYP8B1 <i>e!</i>	ENST00000437102 <i>e!</i>	?	ENSP00000404499 <i>e!</i>	1
KRBOX1 <i>e!</i>	ENST00000426937 <i>e!</i>	?	ENSP00000413859 <i>e!</i>	1
RP11-136C24.3 <i>e!</i>	ENST00000451200 <i>e!</i>	?	?	1
RP11-141M3.6 <i>e!</i>	ENST00000487368 <i>e!</i>	?	?	1
RP11-141M3.6 <i>e!</i>	ENST00000471537 <i>e!</i>	?	?	1
RP11-141M3.6 <i>e!</i>	ENST00000496604 <i>e!</i>	?	?	1

Non-coding exon variant

gene	affected transcript		RefSeq id	variant(s)
ACKR2 <i>e!</i>	ENST00000460855 <i>e!</i>	?		1
ACKR2 <i>e!</i>	ENST00000498111 <i>e!</i>	?		1

