

# SNiPAcord

## Block annotations

### Block info

genomic range	chr2:85,933,003-85,938,407 <i>e!</i>
block size	5,405 bp
variant count	6 variants

### Basic features

Conservation/deleteriousness		Linked genes	
phyloP	$\mu = -0.852$ [-2.998 – 0.359]	gene(s) hit or close-by	-
phastCons	$\mu = 0.020$ [0 – 0.116]	eQTL gene(s)	AC012454.4 <i>e!</i> , ATOH8 <i>e!</i> , GGCX <i>e!</i> , GNLY <i>e!</i> , MLH3 <i>e!</i> , VAMP8 <i>e!</i>
GERP++	$\mu = -1.305$ [-5.72 – 2.01]	potentially regulated gene(s)	AC105053.3 <i>e!</i> , CAPG <i>e!</i> , IMMT <i>e!</i> , MAT2A <i>e!</i> , MRPL35 <i>e!</i> , POLR1A <i>e!</i> , PTCO3 <i>e!</i> , SFTPB <i>e!</i> , SH2D6 <i>e!</i> , TGOLN2 <i>e!</i> , TMEM150A <i>e!</i> , USP39 <i>e!</i> , VAMP5 <i>e!</i> , VAMP8 <i>e!</i>
CADD score	$\mu = 2.968$ [0.191 – 6.026]	disease gene(s)	GGCX <i>e!</i> , MLH3 <i>e!</i> , SFTPB <i>e!</i>

## Trait annotations










### Disease gene annotation

gene	trait	source DB	source entry/link
GGCX <i>e!</i>	VITAMIN K-DEPENDENT CLOTTING FACTORS, COMBINED DEFICIENCY OF, [...]	OMIM	MIM:277450 <i>OMIM</i> <sup>®</sup>
GGCX <i>e!</i>	PSEUDOXANTHOMA ELASTICUM-LIKE DISORDER WITH MULTIPLE COAGULATION [...]	OMIM	MIM:610842 <i>OMIM</i> <sup>®</sup>
MLH3 <i>e!</i>	COLORECTAL CANCER, HEREDITARY NONPOLYPOSIS, TYPE 7	OMIM	MIM:614385 <i>OMIM</i> <sup>®</sup>
MLH3 <i>e!</i>	COLORECTAL CANCER	OMIM	MIM:114500 <i>OMIM</i> <sup>®</sup>
SFTPB <i>e!</i>	SURFACTANT METABOLISM DYSFUNCTION, PULMONARY, 1	OMIM	MIM:265120 <i>OMIM</i> <sup>®</sup>
SFTPB <i>e!</i>	RESPIRATORY DISTRESS SYNDROME IN PREMATURE INFANTS	OMIM	MIM:267450 <i>OMIM</i> <sup>®</sup>
GGCX <i>e!</i>	Body skin hyperlaxity due to vitamin K-dependent coagulation factor deficiency	OrphaNet	OrphaNet:91135 <i>orphanet</i>
GGCX <i>e!</i>	Hereditary combined deficiency of vitamin K-dependent clotting factors	OrphaNet	OrphaNet:98434 <i>orphanet</i>
MLH3 <i>e!</i>	Hereditary nonpolyposis colon cancer	OrphaNet	OrphaNet:144 <i>orphanet</i>
SFTPB <i>e!</i>	Congenital pulmonary alveolar proteinosis	OrphaNet	OrphaNet:264675 <i>orphanet</i>
SFTPB <i>e!</i>	Infant acute respiratory distress syndrome	OrphaNet	OrphaNet:70587 <i>orphanet</i>
SFTPB <i>e!</i>	Neonatal acute respiratory distress with surfactant metabolism deficiency	OrphaNet	OrphaNet:217563 <i>orphanet</i>

## Direct effect on regulation

### cis-eQTL

gene	transcript	probe	tissue	min(statistic) (type)	source	variant(s)
GNLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	pancreas	8.36×10 <sup>-9</sup> (p-value)	GTE Portal V6 <i>mq</i>	6
GNLY <i>e!</i>	ENST00000489214 <i>e!</i>	ILMN_1708779 <i>e!</i>	adipocyte	6.83×10 <sup>-14</sup> (p-value)	MuTHER consortium <i>mq</i>	5
GNLY <i>e!</i>	ENST00000409696 <i>e!</i>					
GNLY <i>e!</i>	ENST00000534351 <i>e!</i>					
GNLY <i>e!</i>	ENST00000489980 <i>e!</i>					
GNLY <i>e!</i>	ENST00000533041 <i>e!</i>					
GNLY <i>e!</i>	ENST00000482900 <i>e!</i>					
GNLY <i>e!</i>	ENST00000470974 <i>e!</i>					
GNLY <i>e!</i>	ENST00000464298 <i>e!</i>					
GNLY <i>e!</i>	ENST00000263863 <i>e!</i>					
GNLY <i>e!</i>	ENST00000524600 <i>e!</i>					
GNLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	muscularis mucosae	1.78×10 <sup>-14</sup> (p-value)	GTE Portal V6 <i>mq</i>	6

GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	lung	1.37×10 <sup>-11</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	atrial appendage	6.52×10 <sup>-7</sup> (p-value)	GTEX Portal V6 	5
ATOH8 <i>e!</i>	?	ENSG00000168874 <i>e!</i>	blood	5.22×10 <sup>-20</sup> (p-value)	GTEX Portal V6 	6
AC012454.4 <i>e!</i>	?	ENSG00000203363 <i>e!</i>	blood	1.46×10 <sup>-12</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	blood	9.39×10 <sup>-24</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	thyroid	6.13×10 <sup>-11</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	skeletal muscle	4.22×10 <sup>-6</sup> (p-value)	GTEX Portal V6 	5
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	prostate	9.77×10 <sup>-9</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	transverse colon	1.42×10 <sup>-8</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	sun exposed skin	7.34×10 <sup>-7</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	aorta	2.64×10 <sup>-8</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	subcutaneous adipocytes	5.31×10 <sup>-12</sup> (p-value)	GTEX Portal V6 	6
?	?	ILMN_2256295 <i>e!</i>	monocyte	7.03×10 <sup>-22</sup> (p-value)	Zeller et al. 	2
?	?	ILMN_1790692 <i>e!</i>	monocyte	1.26×10 <sup>-24</sup> (p-value)	Zeller et al. 	2
ATOH8 <i>e!</i>	ENST00000469442 <i>e!</i>	ILMN_1654065 <i>e!</i>	monocyte	1.32×10 <sup>-157</sup> (p-value)	Zeller et al. 	2
ATOH8 <i>e!</i>	ENST00000306279 <i>e!</i>					
ATOH8 <i>e!</i>	ENST00000489682 <i>e!</i>					
GPLY <i>e!</i>	ENST00000489214 <i>e!</i>	ILMN_1708779 <i>e!</i>	monocyte	2.86×10 <sup>-19</sup> (p-value)	Zeller et al. 	2
GPLY <i>e!</i>	ENST00000409696 <i>e!</i>					
GPLY <i>e!</i>	ENST00000534351 <i>e!</i>					
GPLY <i>e!</i>	ENST00000489980 <i>e!</i>					
GPLY <i>e!</i>	ENST00000533041 <i>e!</i>					
GPLY <i>e!</i>	ENST00000482900 <i>e!</i>					
GPLY <i>e!</i>	ENST00000470974 <i>e!</i>					
GPLY <i>e!</i>	ENST00000464298 <i>e!</i>					
GPLY <i>e!</i>	ENST00000263863 <i>e!</i>					
GPLY <i>e!</i>	ENST00000524600 <i>e!</i>					
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	visceral adipocytes	1.30×10 <sup>-7</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	stomach	9.04×10 <sup>-10</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	spleen	3.83×10 <sup>-8</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	sigmoid colon	3.04×10 <sup>-6</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	coronary artery	3.38×10 <sup>-6</sup> (p-value)	GTEX Portal V6 	5
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	tibial nerve	1.39×10 <sup>-6</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	esophagus mucosa	2.80×10 <sup>-16</sup> (p-value)	GTEX Portal V6 	6
GPLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	gastroesophageal junction	2.82×10 <sup>-7</sup> (p-value)	GTEX Portal V6 	6
GGCX <i>e!</i>	ENST00000233838 <i>e!</i>	ILMN_1758232 <i>e!</i>	blood	1.22×10 <sup>-4</sup> (p-value)	Westra et al. 	1
VAMP8 <i>e!</i>	ENST00000263864 <i>e!</i>	ILMN_2190084 <i>e!</i>	blood	9.93×10 <sup>-6</sup> (p-value)	Westra et al. 	5
VAMP8 <i>e!</i>	ENST00000432071 <i>e!</i>					
VAMP8 <i>e!</i>	ENST00000409760 <i>e!</i>					

ATOH8 <i>e!</i>	ENST00000306279 <i>e!</i>	ILMN_1654065 <i>e!</i>	monocyte	2.74×10 <sup>-25</sup> (p-value)	Fairfax et al. <i>!</i>	1
ATOH8 <i>e!</i>	ENST00000469442 <i>e!</i>					
ATOH8 <i>e!</i>	ENST00000489682 <i>e!</i>					
GNLY <i>e!</i>	?	ENSG00000115523 <i>e!</i>	terminal ileum	1.09×10 <sup>-6</sup> (p-value)	GTEx Portal V6 <i>!</i>	1

### trans-eQTL

gene	transcript	probe	chromosome	tissue	min(statistic) (type)	source	variant(s)
MLH3 <i>e!</i>	ENST00000555144 <i>e!</i>	ILMN_2282282 <i>e!</i>	chr14	monocyte	2.52×10 <sup>-18</sup> (p-value)	Zeller et al. <i>!</i>	2
MLH3 <i>e!</i>	ENST00000555499 <i>e!</i>						
MLH3 <i>e!</i>	ENST00000553713 <i>e!</i>						
MLH3 <i>e!</i>	ENST00000556740 <i>e!</i>						
MLH3 <i>e!</i>	ENST00000355774 <i>e!</i>						

### Putative effect on regulation

#### ENCODE promoter-associated distal DHS (Enhancer)

SNiPA enhancer id	variant(s)	associated SNiPA promoter id	associated gene(s)
ENCE00000253865 <i>e!</i>	1	ENCP00000026860	USP39 <i>e!</i>
		ENCP00000026814	TGOLN2 <i>e!</i>
		ENCP00000026873	AC105053.3 <i>e!</i>
		ENCP00000026830	CAPG <i>e!</i>
		ENCP00000026897	MRPL35 <i>e!</i>
		ENCP00000026853	TMEM150A <i>e!</i>
		ENCP00000026896	IMMT <i>e!</i>
ENCE00000253866 <i>e!</i>	2	ENCP00000026854	USP39 <i>e!</i>
		ENCP00000026837	SH2D6 <i>e!</i>
		ENCP00000026840	MAT2A <i>e!</i>
		ENCP00000026849	TMEM150A <i>e!</i>
		ENCP00000026888	PTCD3 <i>e!</i>
		ENCP00000026853	TMEM150A <i>e!</i>
		ENCP00000026863	SFTPB <i>e!</i>
		ENCP00000026834	SH2D6 <i>e!</i>
		ENCP00000026841	MAT2A <i>e!</i>
		ENCP00000026864	SFTPB <i>e!</i>
		ENCP00000026883	POLR1A <i>e!</i>
		ENCP00000026845	VAMP8 <i>e!</i>
		ENCP00000026858	USP39 <i>e!</i>
		ENCP00000026825	CAPG <i>e!</i>
		ENCP00000026814	TGOLN2 <i>e!</i>
		ENCP00000026828	CAPG <i>e!</i>
		ENCP00000026830	CAPG <i>e!</i>
ENCP00000026847	VAMP5 <i>e!</i>		

#### Regulatory feature cluster

element id	variant(s)	tissue/cell	factors
ENSR00000593772 <i>e!</i> (promoter flanking region)	3	embryonic stem cell (H1ESC)	DNase1, H3K27me3, Rad21, CTCF
		HSMMtube	DNase1, CTCF
		blood (K562)	H3K27me3, DNase1, Rad21, CTCF
		skin (NHDF-AD)	CTCF, DNase1
		muscle (HSMM)	CTCF, H3K27ac, DNase1
		liver (HepG2)	Rad21, CTCF, H3K27me3
		lung (IMR90)	DNase1, CTCF
		blood (GM12878)	H3K4me2, CTCF, H3K27ac, Rad21, H3K4me3, PU1, Srf, BATF, H3K4me1, DNase1

		nervous (NH-A)	CTCF, DNase1
		skin (NHEK)	DNase1, H3K4me1, CTCF
		NHLF	CTCF, DNase1
		Osteobl	CTCF, H3K27ac
		blood (DND-41)	CTCF, H3K27me3
		breast (HMEC)	DNase1, H3K4me1, CTCF
		cervix (HeLa-S3)	Jund, H3K27ac, H3K4me1, CTCF, DNase1
		monocytes (Monocytes-CD14+)	CTCF, H3K27ac, H3K9ac, H3K4me1, DNase1
		endothelium (HUVEC)	DNase1, CTCF
		A549	H3K4me3, CTCF, DNase1
ENSR00000676793 <i>e!</i>	4	embryonic stem cell (H1ESC)	Rad21, CTCF, DNase1
(CTCF binding site)		HSMMtube	DNase1, CTCF, H3K27me3
		blood (K562)	H3K27me3, DNase1, Rad21, CTCF
		skin (NHDF-AD)	CTCF, DNase1
		muscle (HSMM)	CTCF, DNase1
		liver (HepG2)	Rad21, CTCF, H3K27me3
		lung (IMR90)	DNase1, CTCF
		blood (GM12878)	DNase1, Rad21, CTCF
		nervous (NH-A)	CTCF, H3K27me3
		skin (NHEK)	DNase1, CTCF
		NHLF	CTCF
		Osteobl	H3K27me3, CTCF
		blood (DND-41)	CTCF, H3K27me3
		breast (HMEC)	CTCF
		cervix (HeLa-S3)	DNase1, CTCF
		monocytes (Monocytes-CD14+)	H3K9ac, DNase1, H3K4me1, CTCF, H3K27ac
		endothelium (HUVEC)	DNase1, CTCF
		A549	H3K4me3, CTCF, DNase1
ENSR00001544078 <i>e!</i>	1	HSMMtube	H3K27me3
(enhancer)		blood (DND-41)	H3K27me3
		blood (K562)	H3K27me3
		muscle (HSMM)	DNase1
		monocytes (Monocytes-CD14+)	H3K4me1, H3K27ac
		liver (HepG2)	H3K27me3
		nervous (NH-A)	H3K27me3
		skin (NHEK)	H3K27me3

