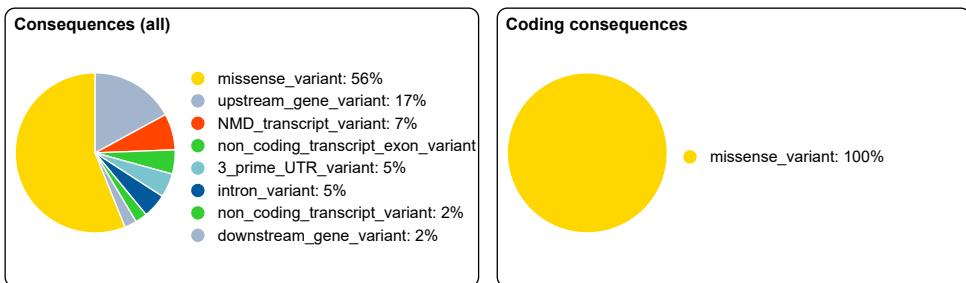


Variant Effect Predictor results

Job details

Summary statistics

Category	Count
Variants processed	4
Variants filtered out	0
Novel / existing variants	0 (0.0) / 4 (100.0)
Overlapped genes	6
Overlapped transcripts	74
Overlapped regulatory features	0



Results preview

Navigation (per variant) Filters Download

Page: 1 of 1 | Show: All variants Uploaded variant Add All: VCF VEP TXT BioMart: Variants Genes

New job

Show/hide columns (14 hidden)

Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000344610.12	protein_coding	
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000349533.11	protein_coding	
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000371058.1	protein_coding	
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000371059.7	protein_coding	
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000371060.7	protein_coding	
1:65570758-65570758	G	non_coding_transcript_exon_variant	LEPR	ENSG00000116678	Transcript	ENST00000462765.5	protein_coding_CDS_not_defined	
1:65570758-65570758	G	missense_variant	LEPR	ENSG00000116678	Transcript	ENST00000616738.4	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000269305.9	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000359597.8	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000413465.6	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000420246.6	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000445888.6	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000455263.6	protein_coding	
17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000503591.2	protein_coding	

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Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000504290.5	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000504937.5	protein_coding
	17:7676154-7676154	A	non_coding_transcript_exon_variant	TP53	ENSG00000141510	Transcript	ENST00000505014.5	retained_intron
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000508793.6	protein_coding
	17:7676154-7676154	A	intron_variant	TP53	ENSG00000141510	Transcript	ENST00000509690.6	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000510385.5	protein_coding
	17:7676154-7676154	A	intron_variant	TP53	ENSG00000141510	Transcript	ENST00000514944.6	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000574684.1	protein_coding_CDS_not_defined
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000576024.2	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000604348.6	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000610292.4	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000610538.4	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000610623.4	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000618944.4	protein_coding
	17:7676154-7676154	A	upstream_gene_variant	TP53	ENSG00000141510	Transcript	ENST00000619186.4	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000619485.4	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000620739.4	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000622645.4	protein_coding
	17:7676154-7676154	A	missense_variant, NMD_transcript_variant	TP53	ENSG00000141510	Transcript	ENST00000635293.1	nonsense-mediated_decay
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000714356.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000714357.1	protein_coding
	17:7676154-7676154	A	3_prime_UTR_variant, NMD_transcript_variant	TP53	ENSG00000141510	Transcript	ENST00000714358.1	nonsense-mediated_decay

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Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000714359.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000714408.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000714409.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000905353.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000923566.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000923567.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000923568.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000923569.1	protein_coding
	17:7676154-7676154	A	missense_variant	TP53	ENSG00000141510	Transcript	ENST00000949117.1	protein_coding
	18:60371535-60371535	T	missense_variant	MC4R	ENSG00000166603	Transcript	ENST00000299766.5	protein_coding
	18:60371535-60371535	T	intron_variant, non_coding_transcript_variant	-	ENSG00000285681	Transcript	ENST00000650201.1	lncRNA
	18:60371535-60371535	T	intron_variant, non_coding_transcript_variant	-	ENSG00000285681	Transcript	ENST00000658928.1	lncRNA
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000333646.11	protein_coding
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000374080.8	protein_coding
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000374102.6	protein_coding
	X:71121046-71121046	T	downstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000429213.3	nonsense-mediated_decay
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000462984.2	retained_intron
	X:71121046-71121046	T	3_prime_UTR_variant, NMD_transcript_variant	MED12	ENSG00000184634	Transcript	ENST00000686548.1	nonsense-mediated_decay
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000687382.1	protein_coding
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000688079.1	retained_intron
	X:71121046-71121046	T	missense_variant, NMD_transcript_variant	MED12	ENSG00000184634	Transcript	ENST00000688663.1	nonsense-mediated_decay
	X:71121046-71121046	T	non_coding_transcript_exon_variant	MED12	ENSG00000184634	Transcript	ENST00000688718.1	retained_intron
	X:71121046-71121046	T	3_prime_UTR_variant, NMD_transcript_variant	MED12	ENSG00000184634	Transcript	ENST00000689008.1	nonsense-mediated_decay
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000689768.1	retained_intron
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000690145.1	protein_coding
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000690242.1	protein_coding
	X:71121046-71121046	T	non_coding_transcript_exon_variant	MED12	ENSG00000184634	Transcript	ENST00000690828.1	retained_intron
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000691113.1	nonsense-mediated_decay
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000691385.1	retained_intron
	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000691426.1	retained_intron
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000691468.1	protein_coding
	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000692304.1	protein_coding

Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
.	X:71121046-71121046	T	3_prime_UTR_variant, NMD_transcript_variant	MED12	ENSG00000184634	Transcript	ENST00000692864.1	nonsense-mediated_decay
.	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000693324.1	protein_coding
.	X:71121046-71121046	T	upstream_gene_variant	MED12	ENSG00000184634	Transcript	ENST00000693391.1	protein_coding
.	X:71121046-71121046	T	downstream_gene_variant	-	ENSG00000228427	Transcript	ENST00000740246.1	lncRNA
.	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000938012.1	protein_coding
.	X:71121046-71121046	T	missense_variant	MED12	ENSG00000184634	Transcript	ENST00000938013.1	protein_coding

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New job

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