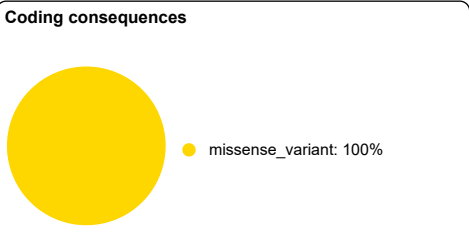
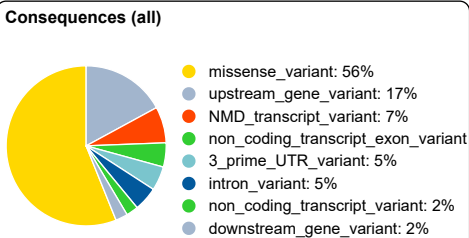


Variant Effect Predictor results

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)

Page: 1 of 1 | Show: 1 All variants

Filters

Uploaded variant is defined

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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
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000344610.12</a>	protein_coding
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000349533.11</a>	protein_coding
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000371058.1</a>	protein_coding
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000371059.7</a>	protein_coding
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000371060.7</a>	protein_coding
.	<a href="#">1:65570758-65570758</a>	G	non_coding_transcript_exon_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000462765.5</a>	protein_coding_CDS_not_defined
.	<a href="#">1:65570758-65570758</a>	G	missense_variant	LEPR	<a href="#">ENSG00000116678</a>	Transcript	<a href="#">ENST00000616738.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000269305.9</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000359597.8</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000413465.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000420246.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000445888.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000455263.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense_variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000503591.2</a>	protein_coding

Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000504290.5</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000504937.5</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	non coding transcript exon variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000505014.5</a>	retained_intron
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000508793.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	intron variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000509690.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000510385.5</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	intron variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000514944.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000574684.1</a>	protein_coding_CDS_not_defined
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000576024.2</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000604348.6</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000610292.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000610538.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000610623.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	upstream gene variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000618944.4</a>	protein_coding
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.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000619485.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000620739.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000622645.4</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant, NMD transcript variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000635293.1</a>	nonsense_mediated_decay
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000714356.1</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000714357.1</a>	protein_coding
.	<a href="#">17:7676154-7676154</a>	A	3 prime UTR variant, NMD transcript variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000714358.1</a>	nonsense_mediated_decay

Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
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	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000714408.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000714409.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000905353.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000923566.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000923567.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000923568.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000923569.1</a>	protein_coding
	<a href="#">17:7676154-7676154</a>	A	missense variant	TP53	<a href="#">ENSG00000141510</a>	Transcript	<a href="#">ENST00000949117.1</a>	protein_coding
	<a href="#">18:60371535-60371535</a>	T	missense variant	MC4R	<a href="#">ENSG00000166603</a>	Transcript	<a href="#">ENST00000299766.5</a>	protein_coding
	<a href="#">18:60371535-60371535</a>	T	intron variant, non coding transcript variant	-	<a href="#">ENSG00000285681</a>	Transcript	<a href="#">ENST00000650201.1</a>	lncRNA
	<a href="#">18:60371535-60371535</a>	T	intron variant, non coding transcript variant	-	<a href="#">ENSG00000285681</a>	Transcript	<a href="#">ENST00000658928.1</a>	lncRNA
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000333646.11</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000374080.8</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000374102.6</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	downstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000429213.3</a>	nonsense_mediated_decay
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000462984.2</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	3 prime UTR variant, NMD transcript variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000686548.1</a>	nonsense_mediated_decay
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000687382.1</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000688079.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	missense variant, NMD transcript variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000688663.1</a>	nonsense_mediated_decay
	<a href="#">X:71121046-71121046</a>	T	non coding transcript exon variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000688718.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	3 prime UTR variant, NMD transcript variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000689008.1</a>	nonsense_mediated_decay
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000689768.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000690145.1</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000690242.1</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	non coding transcript exon variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000690828.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000691113.1</a>	nonsense_mediated_decay
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000691385.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	upstream gene variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000691426.1</a>	retained_intron
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000691468.1</a>	protein_coding
	<a href="#">X:71121046-71121046</a>	T	missense variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000692304.1</a>	protein_coding

Uploaded variant	Location	Allele	Consequence	Symbol	Gene	Feature type	Feature	Biotype
.	<a href="#">X:71121046-71121046</a>	T	3_prime_UTR_variant, NMD_transcript_variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000692864.1</a>	nonsense_mediated_decay
.	<a href="#">X:71121046-71121046</a>	T	missense_variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000693324.1</a>	protein_coding
.	<a href="#">X:71121046-71121046</a>	T	upstream_gene_variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000693391.1</a>	protein_coding
.	<a href="#">X:71121046-71121046</a>	T	downstream_gene_variant	-	<a href="#">ENSG00000228427</a>	Transcript	<a href="#">ENST00000740246.1</a>	lncRNA
.	<a href="#">X:71121046-71121046</a>	T	missense_variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000938012.1</a>	protein_coding
.	<a href="#">X:71121046-71121046</a>	T	missense_variant	MED12	<a href="#">ENSG00000184634</a>	Transcript	<a href="#">ENST00000938013.1</a>	protein_coding

Page: 1 of 1 | Show: [1](#) [All](#) variants

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