

Variant Confirmation Report

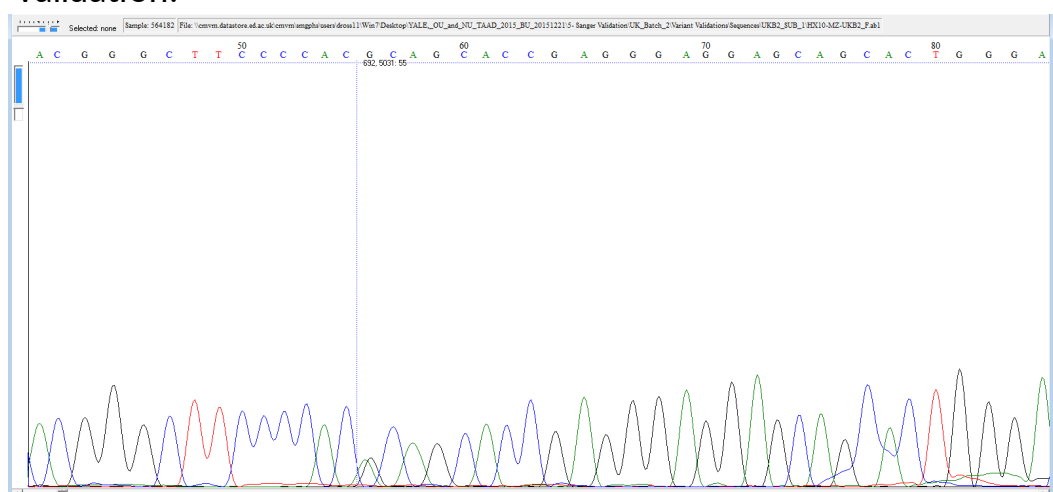
Sample ID:

21MZ0003

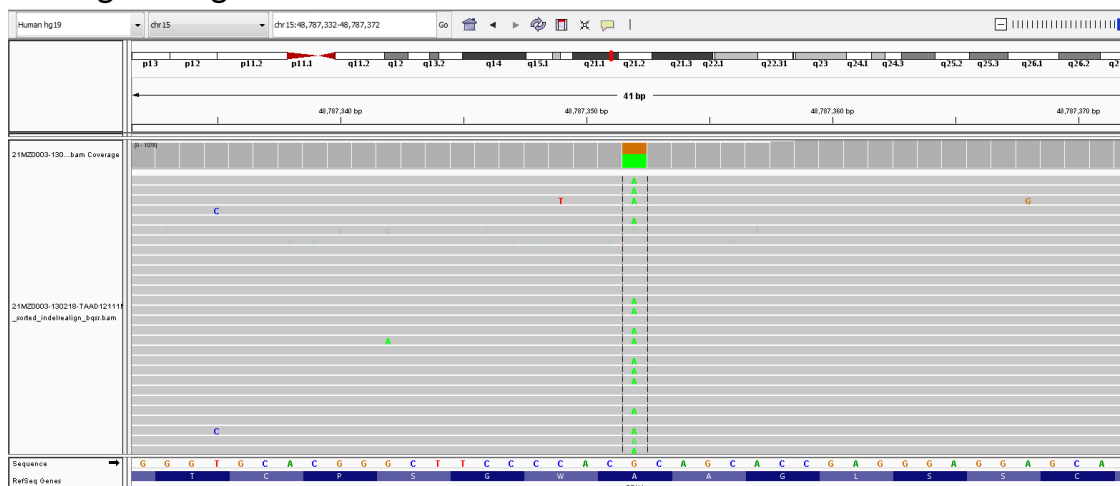
Description:

Gene	FBN1
Exon	22/66
Sequence Variant	ENST00000316623.5:c.2645C>T ENSP00000325527.5:p.Ala882Val
Variant Location (GRCh37)	15:48787352
Allele Balance	0.53
Allele Depth (REF,ALT)	116,131
Allele Frequency (ESP,ExAC,dbSNP)	- - -
Variant Found	Y
Comment	<p>This mutation has been asserted as a disease-causing mutation in the HGMD database</p> <p>HGMD Accession: CM042039 HGMD Classification: DM Loeys (2004) Hum Mutat 24: 140 PubMed: 15241795</p>

Validation:



Fluidigm Image:



The following information is for research purpose only. Any decisions made on the information should be made by an appropriate responsible clinician who may require further confirmation within a clinical laboratory.

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