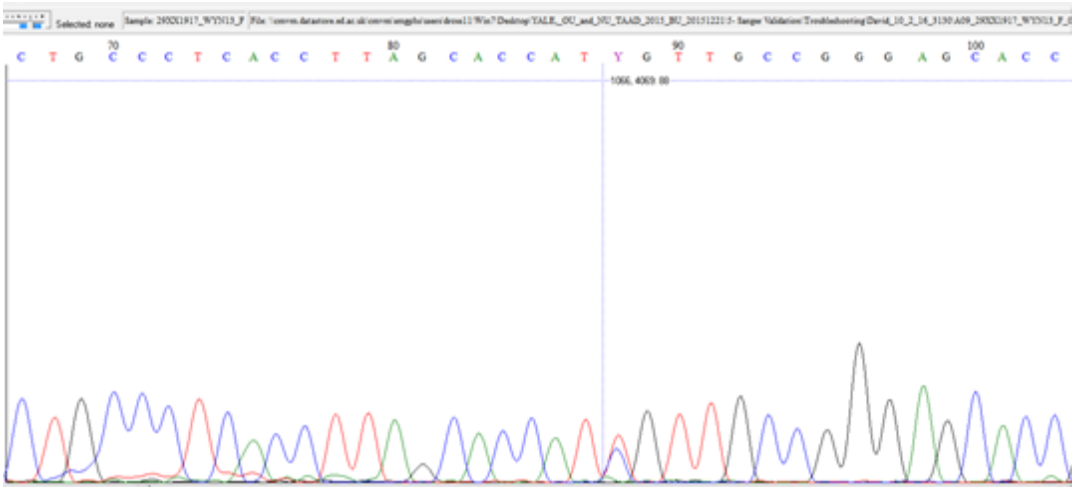
Variant Confirmation Report

Sample ID:	21GC1025

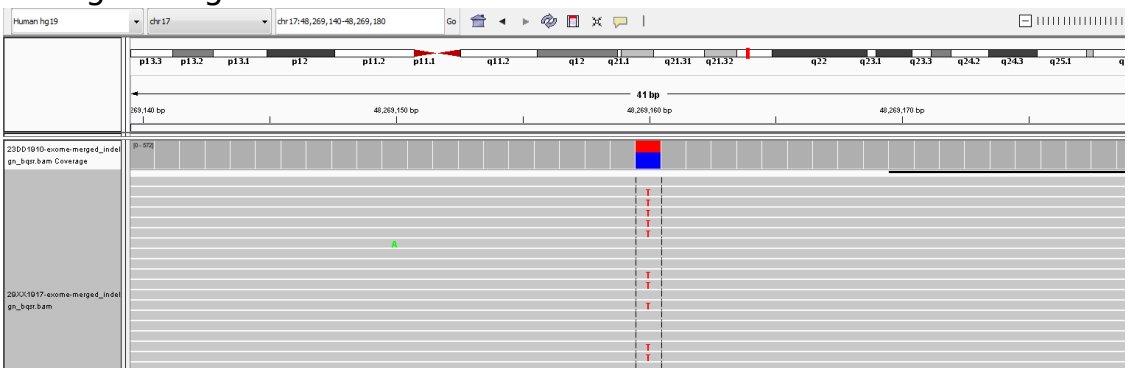
Description:

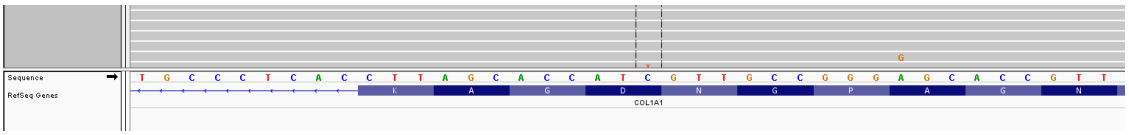
Gene	TGFBR1
Intron	7/8
Sequence Variant	ENST00000374994.4:c.1255+1G>A -
Variant Location (GRCh37)	9:101908892
Allele Balance	0.44
Allele Depth (REF,ALT)	129,102
Allele Frequency (ESP,ExAC,dbSNP)	- - -
Variant Found	Y
Comment	Splicing variant. This will require further investigation

Validation:



Fluidigm Image:





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