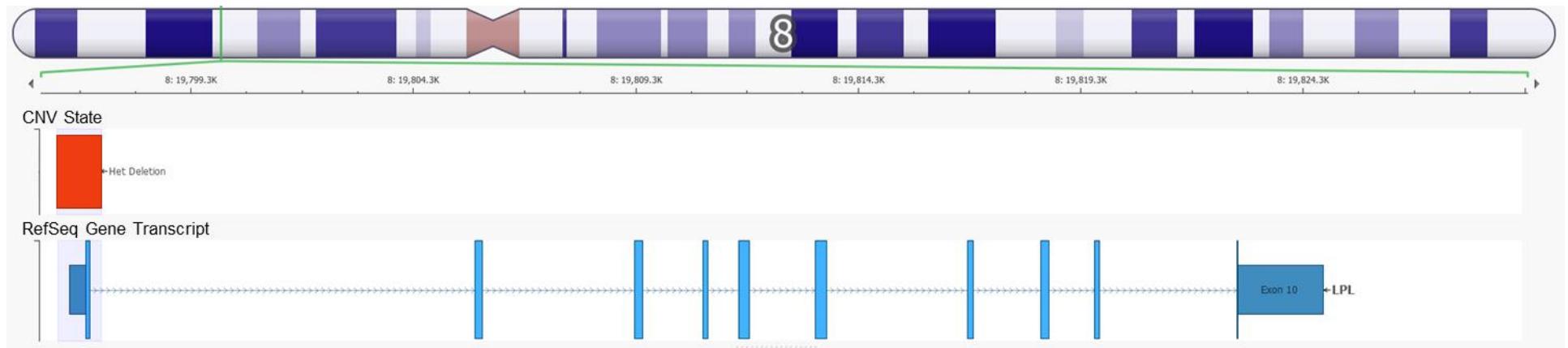
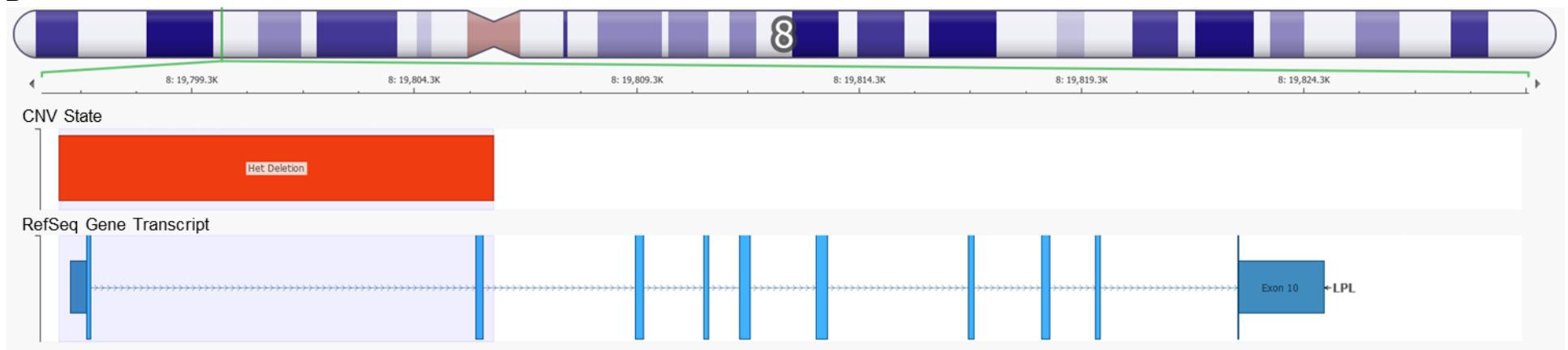


Supplemental Table S1: Screening primers for *LPL* CNVs.

CNV	Breakpoint	Primer direction	Primer sequence (5' to 3')	Annealing temp (°C)	Primer labels in Figure 1 and 2
5'UTR – exon 1	Upstream	F	TTGTAGGTTAGAGTGAACGTGCACAG	60	P2
		R	CATTATGCTGATGCTGCACAACTCTG	60	P3
	Downstream	F	TTCACACTTGATGGTCTCATTCACTGG	60	P4
		R	GATCAGACTGAATTGATTGGTCTGTTCA	60	P5
5' UTR – exon 2	Upstream	F	CTCTATTGGACGTGCTAATGGCACAG	60	P1
		R	CATTATGCTGATGCTGCACAACTCTG	60	P3
	Downstream	F	ACTGACATGCTGACATGCCAGATG	60	P6
		R	CATCTGTGTGAATTCTGTTAGTAG	60	P7
<p>The primers listed were designed to flank the two breakpoints for each CNV. The “Breakpoint” listed is relative to the deleted section of the gene. The sequence orientation for P1-P7 are relative to <i>LPL</i>. Highlighted primer sequences are the same. Abbreviations: CNV = copy-number variant; F = forward; R = reverse; UTR = untranslated region.</p>					

A**B**

Supplemental Figure S1. Identification of *LPL* CNVs using the VarSeq-CNV® caller algorithm on targeted sequencing data. Chr8:19,795,931-19,829,369 (hg19 genome build) is the region visualized in each panel. **A)** Subject 1, carrier of a heterozygous deletion spanning the 5'UTR and exon 1 of *LPL*. **B)** Subject 2 to 4, carriers of a heterozygous deletion spanning the 5'UTR, exon 1 and exon 2 of *LPL*. Abbreviations: chr = chromosome; CNV = copy-number variant; UTR = untranslated region.