Supplemental Table 5. Retinitis pigmentosa (RP) probands with SNV detected near the novel exon on *IMPDH1.*

PROBAND	PHENO	CHROM	POS	REF	ALT	GENO
NEI_392	RP	chr7	128033767	G	C	heterozygous
NEI_408	RP	chr7	128033796	C	T	heterozygous
NEI_413	RP	chr7	128033796	C	T	heterozygous

The genomic coordinates of the variants are based on the human genome hg19.