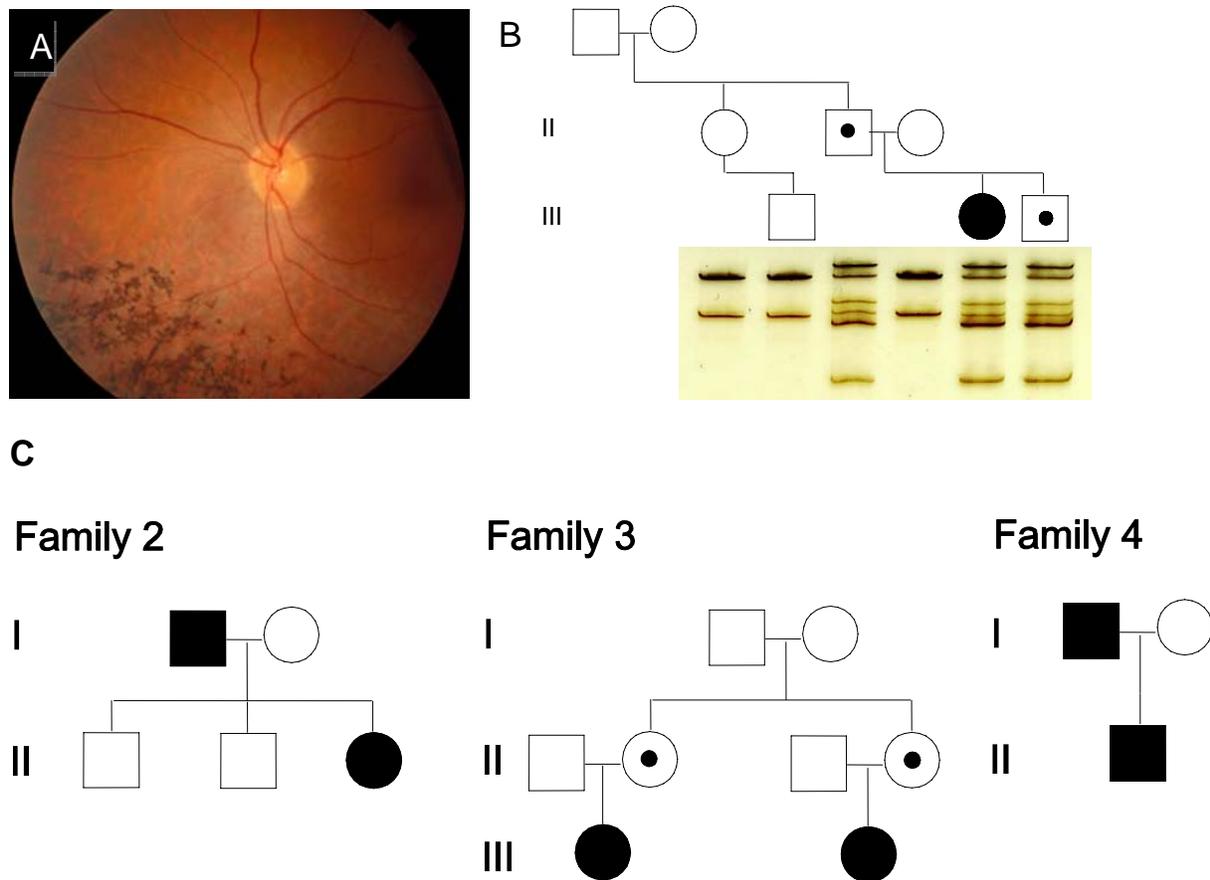


Mutations in *TSPAN12* Cause Autosomal-Dominant

Familial Exudative Vitreoretinopathy

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Figure S1. Pedigrees of FEVR Families and Additional Clinical Information



A, Unusual fundus appearance seen in both eyes of individual II:2 from Family 1 who carries a *TSPAN12* mutation. Fundus photograph shows localised bone spicule pigmentation, reminiscent in appearance to that seen in retinitis pigmentosa. B, Mutation segregating with FEVR in Family 1. Unaffected individuals (unblackened); affected individuals (blackened); individuals asymptomatic

for FEVR (blackened circle in square). Segregation illustrated by SSCP trace of exon 4. C, Pedigrees of FEVR families reported in this study.

Table S1. *TSPAN12* Primers

Exon	Primer sequences (5'-3')	Size (bp)
2	F: ATGTCCCGTGTTCCTCTCTCC	382
	R: CCAGGGGTGGATTTCTTTGT	
3	F: TGGTAATTGGGAAAGATATTATGTAAC	291
	R: CCAAAGATCAAGGAAGAGCA	
4	F: TGAGGCATCATGATTGAAAGAA	346
	R: GCTATCACTGCTCCCTAATCTTGT	
5	F: GGTCCCCTTTCTTGGAGAAC	947
	R: TGGAAATGTGCTTTAGACACA5GA	
6	F: GTACAAAATACCTCTTCATTTATCACA	529
	R: GAAGAAAAGCAGGCCATGAA	
7	F: TGATGACAGATATAGCTCTGGGT	376
	R: TTTTAAGGCCTTTTACATTTAGACA	
8	F: GCTTCCCTGAGAACCACTG	605
	R: CCATCCTCATTTTAAAGCATAGA	