Supplementary material to article by Z. Zhang et al. "Premature Aging Syndrome, Penttinen Type: Report of a Chinese Case with a PDGFRB Mutation"

Fig. S2. (a) Exome sequencing identified a PDGFRB (c.1994T>C, p.Val665Ala) variant (red frame). (b) Sanger sequencing confirmed the PDGFRB variant in the proband. (c–e) Sanger sequencing showed no PDGFRB variants in the proband’s father, mother, sister, respectively.