Incompletely penetrant *PKD1* alleles suggest a role for gene dosage in cyst initiation in polycystic kidney disease


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There was a mistake with Figure 4, panel b. On the second row under 1, it should read 'XY'. The correct figure is shown below.

Figure 4 | Pedigrees of three families with *in utero* ADPKD presentations that have inherited a truncating and a hypomorphic ADPKD allele. Each family has an *in utero* (IU) case, and renal phenotypes of other family members are shown: multiple cysts (MC); negative ultrasound (−ve U/S), or ESRD. Genotypes of the *PKD1* variants: a, Q2158X and R3227C; b, Y3819X and R2765C; and c, 7915dup20 and R2765C, are shown below each pedigree. The etiology of cysts in P118 II is unclear but could potentially represent a low level of mosaicism not detected by sequence analysis.