A common mutation in the COG7 gene with a consistent phenotype including microcephaly, adducted thumbs, growth retardation, VSD and episodes of hyperthermia

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Since the publication of the above article, the authors have noticed an error in the legend of Figure 2. The patient mentioned in the figure is Patient 2 and not Patient 3 as previously published. The correct figure legend is shown below.

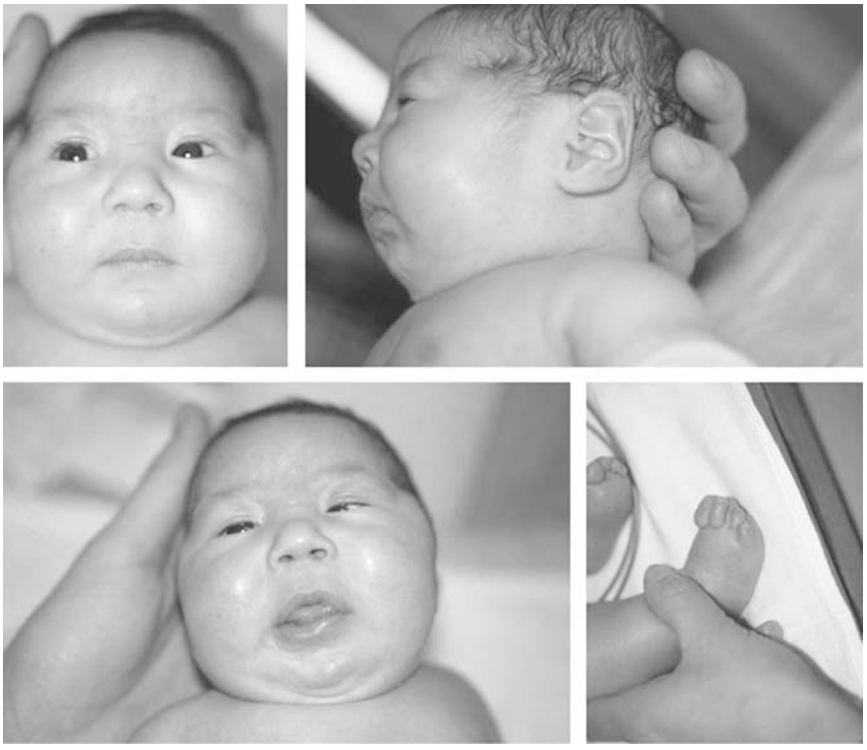

Figure 2 Clinical features of Patient 2 demonstrating microcephaly, flat forehead, dysplastic ears, short nose, retrognathia and irregular insertion of the toes.