CORRECTION

Missense Mutations of the Pro65 Residue of PCGF2 Cause a Recognizable Syndrome Associated with Craniofacial, Neurological, Cardiovascular, and Skeletal Features

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The gender symbol for individual 6 in Figure 1 is incorrect. This individual is female, not male. The corrected figure appears here and online, and the authors regret the error.

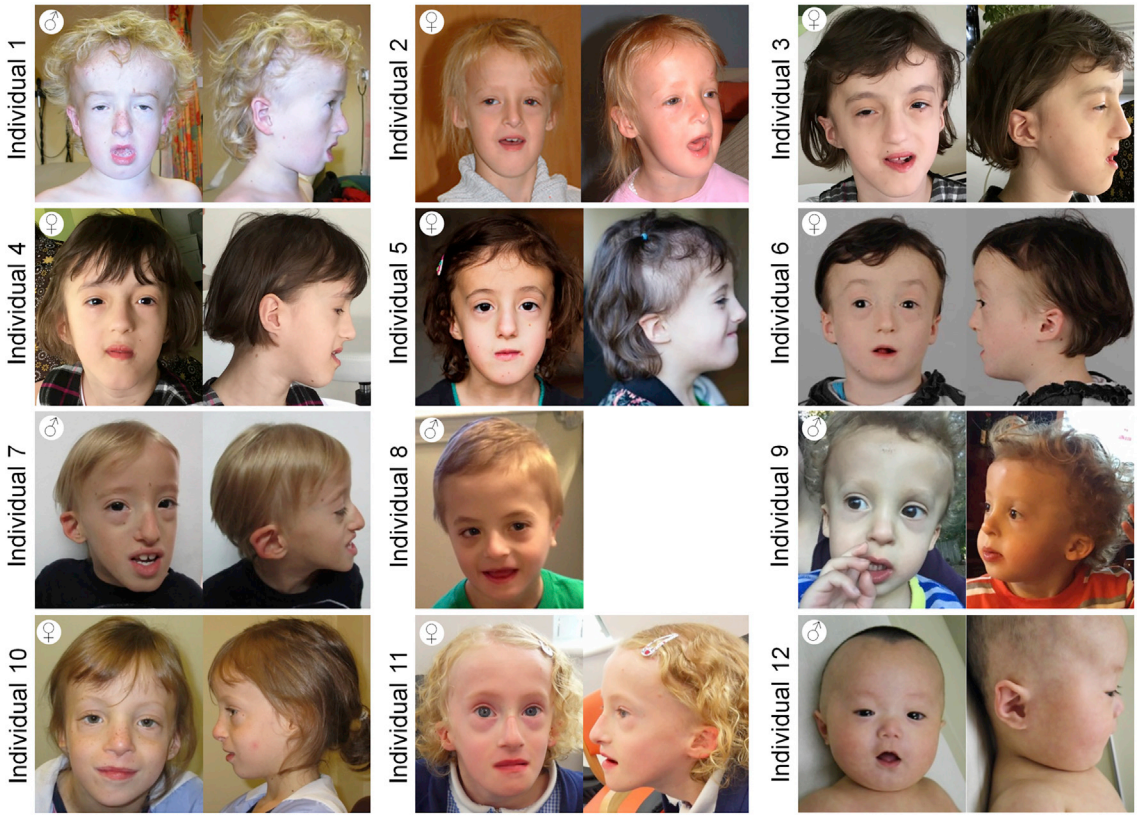

Figure 1. Facial Features of Individuals with PCGF2 Mutations (Corrected)

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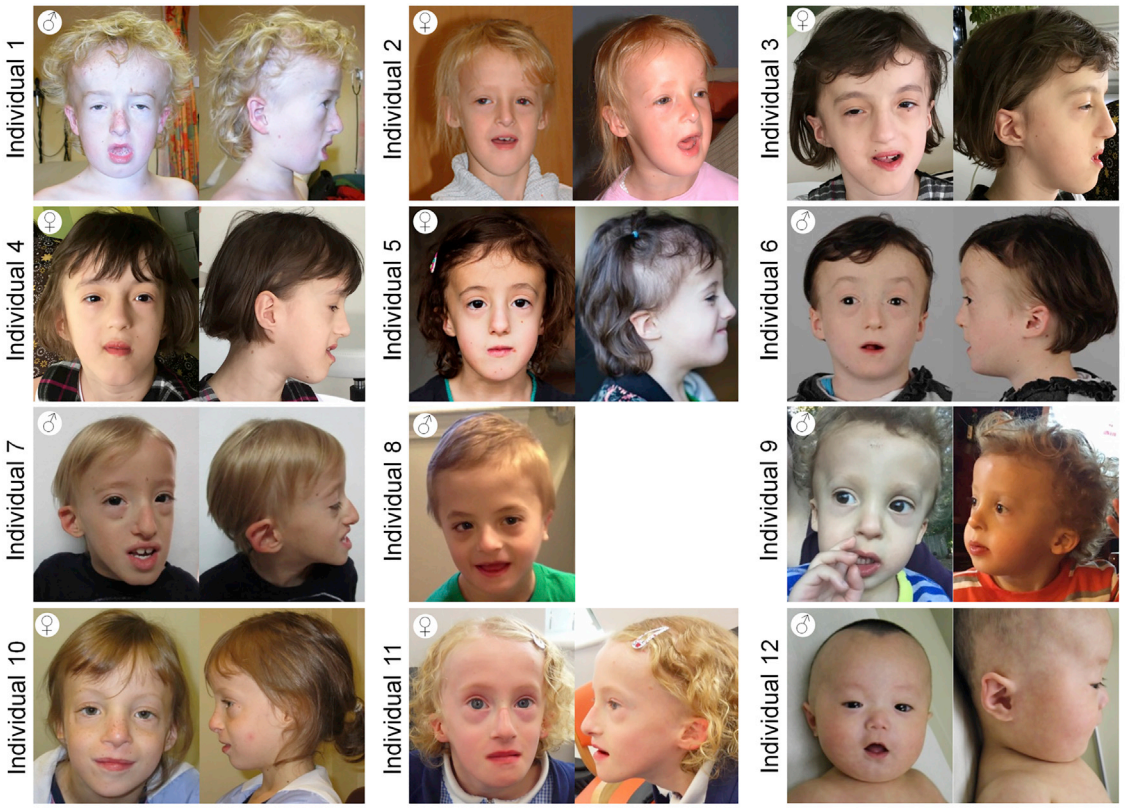

Figure 1. Facial Features of Individuals with PCGF2 Mutations (Original)