

## Corrigendum: Prolyl 3-hydroxylase 1 deficiency causes a recessive metabolic bone disorder resembling lethal/severe osteogenesis imperfecta

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In the version of this article initially published, the nucleotide positions of the mutations in the *LEPRE1* cDNA and genomic DNA sequence in Table 1, Supplementary Table 1 and Supplementary Figure 1 were incorrectly numbered relative to the first nucleotide of exon 1 rather than the first nucleotide of the *LEPRE1* start codon. These errors have been corrected in the HTML and PDF versions of the article.

## Corrigendum: Hypomorphic mutations in syndromic encephalocele genes are associated with Bardet-Biedl syndrome

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In the version of this article initially published, the name of the seventh author was misspelled. The correct author name is Majid Alfadhel. The error has been corrected in the HTML and PDF versions of the article.

## Corrigendum: Mutations in 15-hydroxyprostaglandin dehydrogenase cause primary hypertrophic osteoarthropathy

Sandeep Uppal, Christine P Diggle, Ian M Carr, Colin W G Fishwick, Mushtaq Ahmed, Gamal H Ibrahim, Philip S Helliwell, Anna Latos-Bieleńska, Simon E V Phillips, Alexander F Markham, Christopher P Bennett & David T Bonthron  
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In the version of this article initially published, Figure 3a–d and Supplementary Figure 2a–c show 16-OH PGE<sub>2</sub>, not 15-OH PGE<sub>2</sub>. The errors have been corrected in the HTML and PDF versions of the article.

