CORRIGENDUM

Exome sequencing identifies a *de novo* mutation in HDAC8 associated with Cornelia de Lange syndrome

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mutation ‘c.586A>T; p.M196K’ should have been ‘c.587A>T; p.M196K’.

The authors apologize for any inconvenience caused.

Since the publication of this article, the authors of the above paper have noticed an error in the description of mutation. The *de novo*