Corrigendum: Gain-of-function *SOS1* mutations cause a distinctive form of Noonan syndrome

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In the version of this article initially published online, the labels 'expression' in **Figure 2** panels **a**, **b**, and **d** are incorrect. The correct labels are 'activation'. In addition, author Giuseppe Zampino should have affiliation 8 rather than affiliation 7.
These errors have been corrected for all version of the article.

Corrigendum: A recurrent mutation in the BMP type I receptor ACVR1 causes inherited and sporadic fibrodysplasia ossificans progressiva

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In the version of this article initially published, several contributing authors were listed collectively under the name The FOP International Research Consortium. In order to facilitate the electronic citation of author contributions, the authors have chosen to delete the Consortium name and replace it with the names of the individual consortium authors in alphabetical order. The correct author list and affiliations are as follows:
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Corrigendum: Lamin B1 duplications cause autosomal dominant leukodystrophy

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The construct we described as *moody-GAL4* has a name that was assigned to a different construct in a previously published paper. In this paper, it should be referred to as *SPG-GAL4* (for 'sub-perineural-glia-*GAL4*'). This construct was a gift from R. Bainton (University of California San Francisco School of Medicine). The construction and activity of this promoter will be published elsewhere (R. Bainton, personal communication). The error has been corrected in the PDF version of the article.