Corrigendum: NANS-mediated synthesis of sialic acid is required for brain and skeletal development

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In the version of this article initially published, the name of author Torben Heise was given incorrectly as Thorben Heisse, and the name of author Valérie Cormier-Daire was given incorrectly as Valerie Cormier. The institutional affiliation for Delphine Heron was listed incorrectly as Institut IMAGINE, Hôpital Necker–Enfants Malades, Paris, France, and should have been listed as Département de Génétique Médicale et Centre de Référence Déficiences Intellectuelles, Groupe Hospitalier Pitié-Salpêtrière, Université Pierre et Marie Curie, Paris, France. The errors have been corrected in the HTML and PDF versions of the article.

Corrigendum: Mutations in the histone methyltransferase gene *KMT2B* cause complex early-onset dystonia

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Following publication of this article, the authors were asked to remove a clinical image and some video footage of one of the affected individuals. Although consent was obtained, in keeping with their ethical consent framework, the authors allow for withdrawal of consent and are carrying out the wishes of the research subjects under their consent process. This amendment has been made in the HTML and PDF versions of the article.

Corrigendum: *SMCHD1* mutations associated with a rare muscular dystrophy can also cause isolated arhinia and Bosma arhinia microphthalmia syndrome

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In the version of this article initially published, the legend to Figure 4c stated that only one proband without *SMCHD1* mutation was tested for D4Z4 methylation pattern. However, three probands and one affected family member without *SMCHD1* mutation were tested, as shown in the figure. The error has been corrected in the HTML and PDF versions of the article.