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Erratum

Bardet-Biedl syndrome: A rare genetic disease

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When this article was originally published, some parts of the text appeared incorrect. The corrections are provided below. The original article can be viewed here: http://iospress.metapress.com/content/y604415l68408742/?p=7e8a20cfdf3e44bf99676469a543b026&pi=2.

1 The correct order of authors should be: Sheila Castro-Sánchez, María Álvarez-Satta and Diana Valverde. The first two authors contributed equally to this work.

2 In the abstract and keywords, the following terms should be written in italics: *BBS* genes *and BBS1-BBS17*.

3 Table 1 appeared incorrect. The correct version is presented below.

Table 1
Clinical diagnostic criteria for Bardet-Biedl syndrome [9]

Primary clinical features
Retinitis pigmentosa (rod-cone dystrophy)
Obesity
Polydactyly
Hypogonadism
Intellectual disability/cognitive impairment
Renal abnormalities
Secondary clinical features
Diabetes mellitus type II
Cardiovascular problems
Hearing loss
Speech deficiency
Behavioral problems
Craniofacial dysmorphism
Short stature
Hepatic involvement
Eye abnormalities
Ataxia
Dental and palatal abnormalities
Anosmia
Hirschsprung disease