

## CORRIGENDA

# Next-generation sequencing-based genome diagnostics across clinical genetics centers: implementation choices and their effects

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**Correction to:** *European Journal of Human Genetics* (2015) **23**, 1142–1150; doi:10.1038/ejhg.2014.279; published online 28 January 2015

Post publication, the authors of this paper realised that Olaf R Mook, Department of Human Genetics, Academic Medical Center, University

of Amsterdam, Amsterdam, The Netherlands, was not included in the list of authors. This has now been rectified and the corrected article appears in this issue.

The authors would like to apologise for this omission.

# Further delineation of the KBG syndrome caused by *ANKRD11* aberrations

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Since the publication of this article, the authors have noted that the coordinates and size of the microdeletion in patient 13 were not

mentioned correctly throughout the article. This issue has now been rectified and the corrected article appears in this issue. The HTML and online PDF versions have also been rectified.

The authors would like to apologise for their oversight.