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CORRIGENDA

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

Terry Vrijenhoek, Ken Kraaijeveld, Martin Elferink, Joep de Ligt, Elcke Kranendonk, Gijs Santen, Isaac J Nijman, Derek Butler, Godelieve Claes, Adalberto Costessi, Wim Dorlijn, Winfried van Eyndhoven, Dicky JJ Halley, Mirjam CGN van den Hout, Steven van Hove, Lennart F Johansson, Jan DH Jongbloed, Rick Kamps, Christel EM Kockx, Bart de Koning, Marjolein Kriek, Ronald Lekanne dit Deprez, Hans Lunstroo, Marcel Mannens, Olaf R Mook, Marcel Nelen, Corrette Ploem, Marco Rijnen, Jasper J Saris, Richard Sinke, Erik Sistermans, Marjon van Slegtenhorst, Frank Sleutels, Nienke van der Stoep, Marianne van Tienhoven, Martijn Vermaat, Maartje Vogel, Quinten Waisfisz, Janneke Marjan Weiss, Arthur van den Wijngaard, Wilbert van Workum, Helger Ijntema, Bert van der Zwaag, Wilfred FJ van IJcken, Johan T den Dunnen, Joris A Veltman, Raoul Hennekam and Edwin Cuppen

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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.