European Journal of Human Genetics (2012) 20, 714

www.nature.com/ejhg

CORRIGENDUM

Imaging genetics of *FOXP2* in dyslexia

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European Journal of Human Genetics (2012) **20,** 714; doi:10.1038/ejhg.2012.31

Correction to: *European Journal of Human Genetics* (2012) **20,** 224–229; doi:10.1038/ejhg.2011.160; published online 7 September 2011

The authors would like to make the following amendments:

The following sentence in the Materials and Methods section: 'Furthermore, we genotyped the mutation R553H, previously found in a large Pakistani family with severe speech and language disorder' should read: 'Furthermore, we genotyped the mutation R553H, previously found in a large English Caucasian family with severe speech and language disorder.'

Also, in the Discussion section, the sentence: 'We selected FOXP2-tagging SNPs and the coding-variant R553H, a mutation initially found to be associated with severe speech and language disorder in a Pakistani family by Lai *et al*.' should read: 'We selected FOXP2-tagging SNPs and the coding-variant R553H, a mutation initially found to be associated with severe speech and language disorder in an English Caucasian family by Lai *et al*.'

The authors would like to sincerely apologise for this error and to thank Simon Fisher for drawing their attention to it.