## Correction: Amelogenesis imperfecta in familial hypomagnesaemia and hypercalciuria with nephrocalcinosis caused by CLDN19 gene mutations

Yamaguti PM, Neves FDAR, Hotton D, *et al.* Amelogenesis imperfecta in familial hypomagnesaemia and hypercalciuria with nephrocalcinosis caused by *CLDN19* gene mutations. *J Med Genet* 2017;54:26–37.

One of the author names is spelled incorrectly. 'Pascal Houiller' should be 'Pascal Houillier'.



**Open Access** This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

© Article author(s) (or their employer(s) unless otherwise stated in the text of the article) 2017. All rights reserved. No commercial use is permitted unless otherwise expressly granted.

J Med Genet 2017; 0:1. doi:10.1136/jmedgenet-2016-103956corr1

