Erratum to: *PIGN* mutations cause congenital anomalies, developmental delay, hypotonia, epilepsy, and progressive cerebellar atrophy

Chihiro Ohba · Nobuhiko Okamoto · Yoshiko Murakami · Yasuhiro Suzuki ·
Yoshinori Tsurusaki · Mitsuko Nakashima · Noriko Miyake · Fumiaki Tanaka ·
Taroh Kinoshita · Naomichi Matsumoto · Hirotomo Saitsu

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In abstract and Figure 1 of our paper “*PIGN* mutations cause congenital anomalies, developmental delay, hypotonia, epilepsy, and progressive cerebellar atrophy (DOI: [10.1007/s10048-013-0384-7](https://doi.org/10.1007/s10048-013-0384-7))” we erroneously described the mutation as c.808C>T, instead of c.808T>C. We apologize for this mistake.

The online version of the original article can be found at <http://dx.doi.org/10.1007/s10048-013-0384-7>.

C. Ohba · Y. Tsurusaki · M. Nakashima · N. Miyake ·
N. Matsumoto · H. Saitsu (✉)
Department of Human Genetics, Graduate School of Medicine,
Yokohama City University, Yokohama 236-0004, Japan
e-mail: hsaitsu@yokohama-cu.ac.jp

C. Ohba · F. Tanaka
Department of Clinical Neurology and Stroke Medicine, Yokohama
City University, Yokohama 236-0004, Japan

N. Okamoto
Department of Medical Genetics, Osaka Medical Center and
Research Institute for Maternal and Child Health, Izumi 594-1101,
Japan

Y. Murakami · T. Kinoshita
Department of Immunoregulation, Research Institute for Microbial
Diseases, Osaka University, Osaka 565-0871, Japan

Y. Murakami · T. Kinoshita
World Premier International Immunology Frontier Research Center,
Osaka University, Osaka 565-0871, Japan

Y. Suzuki
Department of Pediatric Neurology, Osaka Medical Center and
Research Institute for Maternal and Child Health, Osaka, Japan