

Joubert Syndrome 2 (JBTS2) in Ashkenazi Jews Is Associated with a *TMEM216* Mutation

Simon Edvardson,^{1,9} Avraham Shaag,^{2,9} Shamir Zenvirt,³ Yaniv Erlich,^{5,6} Gregory J. Hannon,^{5,6} Alan L. Shanske,⁸ John Moshe Gomori,⁴ Joseph Ekstein,⁷ and Orly Elpeleg^{2,3,*}

(The American Journal of Human Genetics 86, 93–97; January 8, 2010)

In this paper, Alan L. Shanske's name was mistakenly omitted from the author list. Above is the corrected author list, and the affiliations are shown below. The authors regret this error.

¹Pediatric Neurology Unit, ²Metabolic Disease Unit, ³Monique and Jacques Roboh Department of Genetic Research, ⁴Department of Radiology, Hadassah, Hebrew University Medical Center, 91120 Jerusalem, Israel; ⁵Watson School of Biological Sciences, ⁶Howard Hughes Medical Institute, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY 11724, USA; ⁷Dor Yeshorim, the Committee for Prevention of Jewish Genetic Diseases, Brooklyn, NY 11219, USA, and 97774 Jerusalem, Israel; ⁸Center for Craniofacial Disorders, Children's Hospital at Montefiore, Albert Einstein College of Medicine, Bronx, NY 10461, USA

⁹These authors contributed equally to this work

*Correspondence: elpeleg@cc.huji.ac.il

DOI 10.1016/j.ajhg.2010.01.022. ©2010 by The American Society of Human Genetics. All rights reserved.