CORRECTIONS

Stereotyped Upper Limb Movement in *MECP2* Duplication Syndrome

Neurology® 2022;98:384. doi:10.1212/WNL.0000000000012757

In the Video NeuroImage article "Stereotyped Upper Limb Movement in *MECP2* Duplication Syndrome" by Wakabayashi et al.,[1] the first sentence should read: "A 23-year-old man had epilepsy, intellectual disability, and a stereotyped movement (Figure 1 and Video 1), which was exacerbated by emotion." The authors regret the error.

Reference

1. Wakabayashi T, Fukumura S, Takahashi S, et al. Stereotyped upper limb movement in *MECP2* duplication syndrome. *Neurology*. 2021;97(2):92-94.

Targeted Next-Generation Sequencing Panels in the Diagnosis of Charcot-Marie-Tooth Disease

Neurology® 2022;98:384. doi:10.1212/WNL.0000000000010635

In the article "Targeted Next-Generation Sequencing Panels in the Diagnosis of Charcot-Marie-Tooth Disease" by Cortese et al.,[1] the gene named throughout the article should be *FGD4*. In the amended supplementary table e-2 (doi.org/10.5061/dryad.kp8pb51), the mutation in patient 111 should be *PMP22*: c.68G>C p.(Thr23Arg), and the mutation in patient 21 should be GARS: c.880G>C p.Gly294Arg. The authors regret the errors.

Reference

1. Cortese A, Wilcox JE, Polke JM, et al. Targeted next-generation sequencing panels in the diagnosis of Charcot-Marie-Tooth disease. *Neurology*. 2020;94(1):e51-e61.

CORRECTION & REPLACEMENT

What Is the Role of Stathmin-2 in Axonal Biology and Degeneration?

Neurology® 2022;98:384. doi:10.1212/WNL.0000000000013018

In the Basic Science in the Clinic article "What Is the Role of Stathmin-2 in Axonal Biology and Degeneration?" by Benarroch,[1] a reference to a study by Melamed et al.[2] was inadvertently omitted at the beginning of the article and miscited elsewhere. The updated references and citations have been added in the replacement version of the article.

In addition, statements regarding TDP-43 in the first paragraph under the "Relationship Among STMN2, TDP-43, and Axonal Degeneration" heading have been corrected in the replacement version of the article. The author agrees with a reader who pointed out that the original wording was not substantiated.

The author regrets the omissions and errors. The original version with the changes highlighted is available from a link in the corrected article.

References

1. Benarroch E. What is the role of stathmin-2 in axonal biology and degeneration? *Neurology*. 2021;97(7):330-333.
2. Melamed Z, Lopez-Erauskin J, Baughn MW, et al. Premature polyadenylation-mediated loss of stathmin-2 is a hallmark of TDP-43-dependent neurodegeneration. *Nat Neurosci*. 2019;22(2):180-190.

