Editors' Note: WriteClick is restricted to comments about studies published in *Neurology*[®] within the past 8 weeks. The submission below represents an exception because it addresses a genetic sequencing error published in *Neurology* in 2003. In addition, these WriteClick submissions will be available as open access to all readers in an effort to further publicize the Correction.

chorea-acanthocytosis (ChAc) that constituted a significant challenge to the generally accepted recessive inheritance of this disorder.² This, and other articles concerning the same family,^{3–5} led to a debate in the research community.⁶

Tomiyasu et al.⁷ noted that the proband reported by Saiki et al. is indeed compound heterozygous for mutations in the *VPS13A* (*CHAC*) gene and, therefore, a typical recessive ChAc case. Subsequent to the publication of this report, we found the Correction only via PubMed but missed the original Correction in *Neurology*.¹

In their Correction,¹ Saiki et al. mention with regret that “an error in sequencing occurred and the inheritance pattern should have been reported as autosomal recessive,” with no reference to the recent publication of the full details of the mutations in this family.⁷ This Correction effectively changes the impact of the original article and, even more so, its title message. We feel that it would be appropriate to retract the 2003 paper in order to reduce further confusion in the literature.

Alternatively, a more compelling communication from the original authors about the significance of the new data could be offered in a manner readily accessible to the scientific community.

Author Response: Shinji Saiki, Ishikawa, Japan: We regret the error published in our 2003 article.¹ Concerning a pedigree with chorea-acanthocytosis in our previous reports,^{1,3–5,8} we would like to correct

genetics materials associated with our pedigree already revised by Tomiyasu et al.⁷ In addition to a heterozygous splice site mutation (8035G>A, transcript variant A [GenBank NM_033305.2]), which we had reported as 8295G>A (using a different accession number) in the *VPS13A* genes of the patients (III-2 and III-3), we identified a novel nonsense mutation (1305G>A, W435X) in the genes of the unaffected mother (II-4) and patients III-2 and III-3.⁷ According to them, the inheritance pattern of the pedigree is actually autosomal recessive (pseudodominant).

We apologize for the mistake and deeply regret any inconvenience this publication has caused for others.

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1. Saiki S, Sakai K, Kitagawa Y, Saiki M, Kataoka S, Hirose G. Mutation in the *CHAC* gene in a family of autosomal dominant chorea-acanthocytosis [published Correction appears in 2011;77:701]. *Neurology* 2003;61:1614–1616.
2. Dobson-Stone C, Rampoldi L, Bader B, et al. Chorea-acanthocytosis. In: Pagon RA, Bird TD, Dolan CR, Stephens K, eds. GeneReviews [Internet], 2010/03/20. Seattle: University of Washington, Seattle; 1993–2002.
3. Saiki S, Hirose G, Sakai K, et al. Chorea-acanthocytosis associated with Tourettism. *Mov Disord* 2004;19:833–836.
4. Saiki S, Sakai K, Murata KY, et al. Primary skeletal muscle involvement in chorea-acanthocytosis. *Mov Disord* 2007;22:848–852.
5. Ishida C, Makifuchi T, Saiki S, Hirose G, Yamada M. A neuropathological study of autosomal-dominant chorea-acanthocytosis with a mutation of *VPS13A*. *Acta Neuropathol* 2009;117:85–94.
6. Bader B, Velayos-Baeza A, Walker RH, Danek A. Dominant transmission of chorea-acanthocytosis with *VPS13A* mutations remains speculative. *Acta Neuropathol* 2009;117:95–96; author reply 97–98.
7. Tomiyasu A, Nakamura M, Ichiba M, et al. Novel pathogenic mutations and copy number variations in the *VPS13A* gene in patients with chorea-acanthocytosis. *Am J Med Genet B Neuropsychiatr Genet* 2011;156B:620–631.
8. Saiki S, Sakai K, Saiki M, Hirose G. Huntington's disease: like 2 can present as chorea-acanthocytosis. *Neurology* 2004;63:939–940; author reply 939–940.

CORRECTION

Pilomotor seizure: When paroxysmal gooseflesh heralds brain tumor

In the Video *NeuroImage* “Pilomotor seizure: When paroxysmal gooseflesh heralds brain tumor” by L. Fisch et al. (*Neurology*[®] 2012;78:1189), there is an error in the author list. The third author's name should read S. Badoud. The publisher regrets the error.

Neurology®

Pilomotor seizure: When paroxysmal gooseflesh heralds brain tumor

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