Errata

In the November 2003 issue of the *Journal,* in the article entitled "Localization of a Gene for Migraine without Aura to Chromosome 4q21," by Björnsson et al. (73: 986–993), the list in appendix A is numbered incorrectly. A corrected appendix appears below. The publisher regrets the error.

Appendix A

Migraine without Aura: International Headache Society Criteria

Previously used terms: common migraine, hemicrania simplex.

Diagnostic criteria:

A. At least five attacks fulfilling items B–D.
B. Headache lasting 4–72 hours (untreated or unsuccessfully treated).
C. Headache with at least two of the following characteristics:
   1. Unilateral location.
   2. Pulsating quality.
   3. Moderate or severe intensity (inhibits or prohibits daily activities).
   4. Aggravation by walking stairs or similar routine physical activity.
D. During headache, at least one of the following:
   1. Nausea and/or vomiting.
   2. Photophobia and phonophobia.
E. At least one of the following:
   1. History and physical and neurologic examinations do not suggest one of the disorders listed in groups 5–11 (organic disorders).
   2. History and/or physical and/or neurologic examinations suggest such disorder, but it is ruled out by appropriate investigations.
   3. Such disorder is present, but migraine attacks do not occur for the first time in close temporal relation to the disorder.

In the February 2005 issue of the *Journal,* in the article entitled "Contrasting Effects of Natural Selection on Human and Chimpanzee CC Chemokine Receptor 5," by Wooding et al. (76:291–301), the legend to figure 5 identifies "rejected" and "not rejected" with red and blue areas, respectively, of the graphs in the figure. "Rejected" actually corresponds with light gray, and "not rejected" corresponds with dark gray. The authors and the publisher regret the error.

In the March 2005 issue of the *Journal,* in the article entitled "*PHOX2B* Genotype Allows for Prediction of Tumor Risk in Congenital Central Hypoventilation Syndrome," by Trochet et al. (76:421–426), Dr. Pär-Johan Svensson had an incorrect affiliation listed and Dr. Stephan Niemann was not identified as an author. The correct list of authors is: Delphine Trochet, Louise M. O. Brien, David Gozal, Ha Trang, Agneta Nordenskjöld, Béatrice Laudier, Pär-Johan Svensson, Sabine Uhrig, Trevor Cole, Stephan Niemann, Arnold Munnich, Claude Gaultier, Stanislas Lyonnet, and Jeanne Amiel. Dr. Svensson's affiliation is Department of Molecular Medicine, Karolinska Hospital, Stockholm. Dr. Niemann's affiliation is Institut für Humangenetik, Giessen, Germany. The authors regret the errors.