Errata

In the January 2001 issue of the *Journal,* in the article "Hereditary Nonpolyposis Colorectal Cancer in 95 Families: Differences and Similarities between Mutation-Positive and Mutation-Negative Kindreds," by Scott et al. (68:118–127), the second sentence in the second paragraph of the Discussion section should read as follows: "Neither of these mutations led to any unique phenotype but each presumably results in altered hMSH2 function" (rather than as "Neither of these mutations led to any unique phenotype—and, presumably, neither results in altered hMSH2 function").

In this issue of the *Journal,* in the article "Periaxin Mutations Cause Recessive Dejerine-Sottas Neuropathy," by Boerkoel et al. (68:325–333), there should be no footnote designation "a" appended to the multicolumn "Family/Patient" heading; the families and patients listed under this heading are from Boerkoel et al.'s collection in Houston and are not associated with the CMT4F family reported by Delague et al. (2000). The publisher regrets this error.