Erratum

In the April 2000 issue of the *Journal,* in the article "Mutational and Haplotype Analyses of Families with Familial Partial Lipodystrophy (Dunnigan Variety) Reveal Recurrent Missense Mutations in the Globular C-Terminal Domain of Lamin A/C," by Speckman et al. (66:1192–1198), family F600 was erroneously reported, both in the "Mutational Analysis" subsection of the Results section and in table 1, as having mutation R482W; in fact, no mutation was detected in this family. The "Mutational Analysis" subsection also contained two additional errors: the variant seen in CEPH individual 1701 was R644C (not R633C), and the polymorphism in exon 5 was A287A (not A297A). The authors regret these errors.