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## ADDENDUM

During proofreading, there appears a work on the isolation of the gene, *UBE3A*, responsible for Angelman syndrome (Kishino *et al.*, 1997) *UBE3A*-mutations identified in 3 AS patients may cause a frameshift and premature translocation termination of the ubiquitin-mediated protein and may lead to degradation of the protein during brain development in AS.

Kishino T, Lalande M, Wagstaff J (1997): *UBE3A/E6-AP* mutations cause Angelman syndrome. *Nature Genet* (in press).