For The Blind to support gene mapping of autosomal dominant nystagmus.

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Corrections

In the paper by Richards *et al* on 'Detailed genetic mapping of the von Hippel-Lindau disease tumour suppressor gene' (*J Med Genet* 1993;**30**:104–7), an important collaborator, Dr Per Enblad, was inadvertently omitted from the authorship. The correct authorship is as follows.

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In the paper by Padayachee *et al* on 'Mapping of the X linked form of hyper IgM syndrome (HIGM1)' (*J Med Genet* 1992;**30**:202–5), the primer sequence for DXS102[13] under the heading OLIGONUCLEOTIDE PRIMERS was referenced Luty *et al*. This is incorrect and should be:

Gedeon AK, Holmon K, *et al*. Characterization of new pcr based markers for mapping and diagnosis: AC dinucleotide repeat markers at the DXS237 (GMGX9) and DXS102 (cX38.1) loci. *Am J Med Genet* 1992;**43**:255–60.