Review article
Clinical features, molecular genetics, and pathophysiology of dominant optic atrophy M Votruba, A T Moore, S S Bhattacharya

Original articles
Localisation of a gene for non-specific X linked mental retardation (MRX46) to Xq25-q26
H G Yntema, B C J Hamel, A P T Smits, T van Roosmalen, B van den Helm, H Kremer, H-H Ropers, D F C M Smeets, H van Bokhoven

The prevalence of PAX2 mutations in patients with isolated colobomas or colobomas associated with urogenital anomalies
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Neurofibromatosis type 1 (NF1): a protein truncation assay yielding identification of mutations in 73% of patients
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Parental origin effects in human trisomy for chromosome 14q: implications for genomic imprinting
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A common DLX3 gene mutation is responsible for tricho-dento-osseous syndrome in Virginia and North Carolina families
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