Journal of Medical Genetics May 1997 Vol 34 No 5





Original articles

Epigenetic modification and uniparental inheritance of H19 in Beckwith-Wiedemann syndrome D Catchpoole, W W K Lam, D Valler, I K Temple, J A Joyce, W Reik, P N Schofield, E R Maher	353
The genetics of primary nocturnal enuresis: inheritance and suggestion of a second major gene	
on chromosome 12q H Arnell, K Hjälmås, M Jägervall, G Läckgren, A Stenberg, B Bengtsson, C Wassén, T Emahazion, G Annerén, U Pettersson, M Sundvall, N Dahl	360
Further delineation of Nevo syndrome L I Al-Gazali, D Bakalinova, E Varady, J Scorer, M Nork	366
Analysis of the 5' upstream sequence of the Huntington's disease (HD) gene shows six new rare alleles which are unrelated to the age at onset of HD R Coles, J Leggo, D C Rubinsztein	371
Genotypic diagnosis of familial Mediterranean fever (FMF) using new microsatellite markers: example of two extensive non-Ashkenazi Jewish pedigrees M Dupont, C Dross, N Smaoui, B Nedelec, G Grateau, C Clépet, I Gourdier, I Koné-Paut, M Delpech, J Demaille, I Touitou	375
Psychological distress in applicants for predictive DNA testing for autosomal dominant, heritable, late onset disorders A C DudokdeWit, A Tibben, H J Duivenvoorden, P G Frets, M W Zoeteweij, M Losekoot, A van Haeringen, M F Niermeijer, J Passchier, and the other members of the Rotterdam/Leiden Genetics Workgroup	382
Homozygosity by descent for a rare mutation in the myophosphorylase gene is associated with variable phenotypes in a Druze family with McArdle disease S lyengar, H Kalinsky, S Weiss, M Korostishevsky, M Sadeh, Y Zhao, K K Kidd, B Bonne-Tamir	391
Trisomy 15 rescue with jumping translocation of distal 15q in Prader-Willi syndrome K Devriendt, P Petit, G Matthijs, J R Vermeesch, M Holvoet, A De Muelenaere, P Marynen, J-J Cassiman, J-P Fryns	395
Syndrome of the month	
Ehlers-Danlos syndrome has varied molecular mechanisms F M Pope, N P Burrows	400
Short reports	
Meiotic instability associated with the CAGR1 trinucleotide repeat at 13q13 NT Potter	411
Chromosome 18q22.2→qter deletion and a congenital anomaly syndrome with multiple vertebral segmentation defects SB Dowton, AV Hing, V Sheen-Kaniecki, MS Watson	414
Autosomal dominant inheritance of Weaver syndrome A Fryer, C Smith, L Rosenbloom, T Cole	418
A novel mutation (a886g) in exon 5 of FGFR2 in members of a family with Crouzon phenotype and plagiocephaly D Steinberger, H Collmann, B Schmalenberger, U Müller	420
Chromosome 22q11 deletion presenting as the Potter sequence K Devriendt, P Moerman, D Van Schoubroeck, K Vandenberghe, J-P Fryns	423
Clinical features in four patients with Angelman syndrome resulting from paternal uniparental disomy A Smith, R Marks, E Haan, J Dixon, R J Trent	426
Interstitial deletion of band 3q25 A M Slavotinek, S M Huson, M Fitchett	430
Wiedemann-Rautenstrauch neonatal progeroid syndrome: report of three new patients H Arboleda, L Quintero, E Yunis	433
Book reviews	438
Notices	440