

Contents

Editorial

- Redrafted Chinese law remains eugenic *M Bobrow* 409
Biographical note: Professor Martin Bobrow 410

Review article

- Nuclear and mitochondrial genetics in Parkinson's disease *A H V Schapira* 411

Original articles

- A clinical and genetic study of campomelic dysplasia *S Mansour, C M Hall, M E Pembrey, I D Young* 415
A large Turkish kindred with syndactyly type II (synpolydactyly). 1 Field investigation, clinical and pedigree data *B S Sayli, A N Akarsu, U Sayli, O Akhan, S Ceylaner, M Sarfarazi* 421
A large Turkish kindred with syndactyly type II (synpolydactyly). 2 Homozygous phenotype? *A N Akarsu, O Akhan, B S Sayli, U Sayli, C Baskaya, M Sarfarazi* 435
DNA microsatellite analysis of families with autosomal dominant polycystic kidney disease types 1 and 2: evaluation of clinical heterogeneity between both forms of the disease *E Coto, S Sanz de Castro, S Aguado, J Alvarez, M Arias, M J Menéndez, C López-Larrea* 442
Allelic associations and homozygosity at loci from HLA-B to D6S299 in genetic haemochromatosis *R Raha-Chowdhury, D J Bowen, A K Burnett, M Worwood* 446
Epidemiology and genetics of microtia-anotia: a registry based study on over one million births *P Mastroiacovo, C Corchia, L D Botto, R Lanni, G Zampino, D Fusco* 453
The dental phenotype in familial adenomatous polyposis: diagnostic application of a weighted scoring system for changes on dental panoramic radiographs *N Thakker, R Davies, K Horner, J Armstrong, T Clancy, S Guy, R Harris, P Sloan, G Evans* 458
Neuroectodermal (CHIME) syndrome: an additional case with long term follow up of all reported cases *V Shashi, J Zunich, T E Kelly, J S Fryburg* 465
Diagnostic issues in a family with late onset type 2 neurofibromatosis *D G R Evans, D Bourn, A Wallace, R T Ramsden, J D Mitchell, T Strachan* 470

Brief papers

- Identification of RB1 germline mutations in Argentinian families with sporadic bilateral retinoblastoma *I Szijan, D R Lohmann, D L Parma, B Brandt, B Horsthemke* 475
H714Q mutation in Wilson disease is associated with late, neurological presentation *R H J Houwen, J Juyn, T U Hoogenraad, J K Ploos van Amstel, R Berger* 480
Inheritance of CMT1A duplication from a mosaic father *E Sorour, P Thompson, J MacMillan, M Upadhyaya* 483
Good growth response to growth hormone treatment in the ring chromosome 15 syndrome *M Nuutinen, K Kouvalainen, M Knip* 486

Short report

- A new PAX6 mutation in familial aniridia *I Hanson, A Brown, V van Heyningen* 488

Letters to the Editor

- GIG response to the UK Clinical Genetics Society report "The genetic testing of children" *S Dalby* 490
The genetic testing of children *A Clarke* 492
Possible genetic heterogeneity in hypochondroplasia *I Stoilov, M W Kilpatrick, P Tsipouras, T Costa* 492
UPD 13: no indication of maternal or paternal imprinting of genes on chromosome 13 *H Slater, J H Shaw, A Bankier, S M Forrest, G Dawson* 493
A family with autosomal dominant polycystic kidney disease linked to 4q21-23 *S Jeffery, S Morgan, V J Warmington, G A MacGregor, A K Saggarr-Malik* 493

Book reviews

495