

Journal of
**MEDICAL
 GENETICS**

Original articles

- 665 Pancreatic function and gene deletion F508 in cystic fibrosis *G Borgo, G Mastella, P Gasparini, A Zorzanello, R Doro, P F Pignatti*
- 670 Adrenoleucodystrophy: a molecular genetic study in five families *R G Del Mastro, S Bunday, M W Kilpatrick*
- 676 In vitro gene amplification for prenatal diagnosis of congenital adrenal hyperplasia *G Rumsby, J W Honour*
- 679 Deletion analysis of DMD/BMD families from the German Democratic Republic and selected regions of Czechoslovakia and Hungary *A Speer, U Kräft, R Hanke, K Grade, C Couelle, K Wulff, M Wehnert, F H Herrmann, L Kadasi, E Kunert, U Müller, C Förster, C Wolf, R Szibor*
- 683 Theoretical considerations on germline mosaicism in Duchenne muscular dystrophy *T Grimm, B Müller, C R Müller, M Janka*
- 688 MASA syndrome: new clinical features and linkage analysis using DNA probes *C Schrander-Stumpel, E Legius, J P Fryns, J J Cassiman*
- 693 Recessive metaphyseal dysplasia without hypotrichosis. A syndrome clinically distinct from McKusick cartilage-hair hypoplasia *A Verloes, G E Pierard, M Le Merrer, P Maroteaux*
- 697 Exclusion of autosomal dominant polycystic kidney disease type II (ADPKD2) from 160 cM of chromosome 1 *S Kumar, W J Kimberling, P A Gabow, Y Y Shugart, S Pieke-Dahl*

**Syndrome of the month
 Annotation**

- 701 Craniodiaphyseal dysplasia *L A Brueton, R M Winter*
- 707 Teaching of clinical genetics in Britain: a report from the Royal College of Physicians of London *A W Johnston*

Editor's note

- 710 Delivery of genetic services *P S Harper*

Medical genetic services

- 711 Genetic services in Britain: a strategy for success after the National Health Service and Community Care Act 1990 *R Harris*

Case reports

- 715 Familial congenital laryngeal abductor paralysis: different expression in a family with one male and three females affected *A Schinzel, E Hof, P Dangel, W Robinson*
- 717 A child, homozygous for a stop codon in exon 11, shows milder cystic fibrosis symptoms than her heterozygous nephew *H Cuppens, P Marynen, C De Boeck, F De Baets, E Eggermont, H Van den Berghe, J J Cassiman*
- 720 Ankyloblepharon filiforme adnatum in trisomy 18 Edwards syndrome *D G R Evans, I D Evans, D Donnai, R H Lindenbaum*
- 722 Distal long arm deletions of the X chromosome and ovarian failure *A Bates, P J Howard*
- 724 Triploidy arising from a first meiotic non-disjunction in a mother carrying a reciprocal translocation *L Rochon, M J J Vekemans*

727-728 Letters to the Editor