

Contents

**Editorial**

- The genetic testing of children *T M Marteau* 743

**Annotation**

- Syndrome of the month *I Young, A Wilkie* 744

**Original articles**

- Genetic linkage to the type VII collagen gene (COL7A1) in 26 families with generalised recessive dystrophic epidermolysis bullosa and anchoring fibril abnormalities *M G S Dunnill, A J Richards, G Milana, F Mollica, D Atherton, I Winship, M Farrall, L Al-Imara, R A J Eady, F M Pope* 745
- Mosaic uniparental disomy in Beckwith-Wiedemann syndrome *R E Slatter, M Elliott, K Welham, M Carrera, P N Schofield, E R Maher* 749
- X inactivation patterns in female monozygotic twins and their families *E Watkiss, T Webb, G Rysiecki, N Girdler, E Hewett, S Bunday* 754
- Cephalometric analysis of Rapp-Hodgkin syndrome *T C Hart, S Kyrkanides* 758
- Phenotypic variation of tuberous sclerosis in a single extended kindred *S L Smalley, F Burger, M Smith* 761
- Absence of linkage between idiopathic dilated cardiomyopathy and candidate genes involved in the immune function in a large Italian pedigree *M Krajcinovic, L Mestroni, G M Severini, B Pinamonti, F Camerini, A Falaschi, M Giacca* 766
- Osteocraniostenosis *A Verloes, F Narcy, B Grattagliano, A-L Delezoide, P Guibaud, J-P Schaaps, M Le Merrer, P Maroteaux* 772

**Syndrome of the month**

- Albright's hereditary osteodystrophy *L C Wilson, R C Trembath* 779

**Clinical practice in medical genetics**

- The genetic testing of children. Report of a Working Party of the Clinical Genetics Society (UK). Chairman: Dr Angus Clarke 785

**Brief papers**

- Intrachromosomal triplication of 15q11-q13 *A A Schinzel, L Brecevic, F Bernasconi, F Binkert, F Berthet, A Wuilloud, W P Robinson* 798
- A newborn with ring chromosome 10, aganglionic megacolon, and renal hypoplasia *G Calabrese, P G Franchi, L Stuppia, R Mingarelli, C Rossi, L Ramenghi, M Marino, E Morizio, R Peila, A Antonucci, G Palka* 804
- Down-Turner syndrome: case report and review *G J C M Van Buggenhout, B C J Hamel, J C M Trommelen, H Mieloo, D F C M Smeets* 807
- Linkage and mutation analysis in an extended family with Charcot-Marie-Tooth disease type 1B *E Nelis, V Timmerman, P De Jonghe, L Muylle, J-J Martin, C Van Broeckhoven* 811

**Short report**

- Chimaerism shown by cytogenetics and DNA polymorphism analysis *A J Green, D E Barton, P Jenks, J Pearson, J R W Yates* 816

**Abstracts**

- Medical genetics: advances in brief 818

**Letters to the Editor**

- Huntington's disease in two unrelated Arab kindreds and in an Afghani family resident in Saudi Arabia *E M Scrimgeour, S A Tahoona, T H Zawawi* 819
- The Prader-Willi-like phenotype in fragile X patients: a designation facilitating clinical (and molecular) differential diagnosis *B B A De Vries, M F Niermeijer* 820
- Arthrogryposis multiplex congenita, renal dysfunction, and cholestasis syndrome *J M Saraiva, H C Mota* 820

**Book reviews**

- 821

- Notices** 822