

*Journal of***MEDICAL
GENETICS**

- Original articles**
- 217 The value of investigation for genetic counselling in tuberous sclerosis *A E Fryer, A H Chalmers, J P Osborne*
- 224 Attitudes of general practitioners to presymptomatic testing for Huntington's disease *M E Mennie, S M Holloway, D J H Brock*
- 228 The clinical features of three babies with osteogenesis imperfecta resulting from the substitution of glycine by arginine in the pro $\alpha 1(I)$ chain of type I procollagen *W G Cole, C W Chow, J G Rogers, J F Bateman*
- 236 Becker muscular dystrophy: correlation of deletion type with clinical severity *A M Norman, N S T Thomas, H M Kingston, P S Harper*
- 240 Alphafetoprotein in midtrimester Down's syndrome fetal serum *M J Seller*
- 244 Prenatal diagnosis of junctional epidermolysis bullosa associated with pyloric atresia *V Nazzaro, U Nicolini, L De Luca, E Berti, R Caputo*
- 249 Reliability of prenatal diagnosis of genetic diseases by analysis of amplified trophoblast DNA *M C Rosatelli, R Sardu, T Tuveri, M T Scalas, A Di Tucci, M De Murtas, G Loudianos, G Monni, A Cao*
- 252 Ellis-van Creveld syndrome, Jeune syndrome, and renal-hepatic-pancreatic dysplasia: separate entities or disease spectrum? *L A Brueton, M J Dillon, R M Winter*
- Syndrome of the month**
- 256 Wiedemann-Rautenstrauch syndrome *H V Toriello*
- Case reports**
- 258 Rothmund-Thomson syndrome associated with trisomy 8 mosaicism *K L Ying, J Oizumi, C J R Curry*
- 261 Terminal deletion (14)(q32.3): a new case *N Telford, D A G Thomson, M J Griffiths, S Ilett, J L Watt*
- 264 Trisomy 10p syndrome owing to maternal pericentric inversion *K-I Ohba, S Ohdo, T Sonoda*
- 267 A constitutional 5q23 deletion *H Rivera, P Simi, S Rossi, L Pardelli, M C Di Paolo*
- 269 Kyphomelic dysplasia: the first 10 cases *P D Turnpenny, R A Dakwar, F N Boulos*
- Dysmorphology report**
- 273 Fetal brain disruption sequence: a milder variant *C G Bönnemann, P Meinecke*
- 275 Letters to the Editor
- 278 Book reviews
- 280 Notices