

JOURNAL OF MEDICAL GENETICS

- Genetic polymorphism and interethnic variability of plasma paroxonase activity J. R. PLAYFER, L. C. EZE, M. F. BULLEN, and D. A. P. EVANS 337
- Spinal dysraphism: genetic relation to neural tube malformations C. O. CARTER, K. A. EVANS, and K. TILL 343
- Annotation: Risks of miscarriage after amniocentesis C. O. CARTER 351
- An unusual family of benign 'X' linked muscular dystrophy with cardiac involvement R. S. WADIA, S. U. WADGAONKAR, R. B. AMIN, and H. V. SAR-DESAI 352
- Familial essential ('benign') chorea THOMAS D. BIRD, COLDEVIN B. CARLSON, and JUDITH G. HALL 357
- Haemoglobin Lepore_{Boston} in a Turkish family AYHAN O. ÇAVDAR and AYTEN ARCASOY 363
- Partial monosomy and partial trisomy 18 in two offspring of carrier of pericentric inversion of chromosome 18 ANGELA M. VIANNA-MORGANTE, MARIA JOSE NOZAKI, CLÁUDIO C. ORTEGA, VERÔNICA COATES, and YSAO YAMAMURA 366
- Clinical details, cytogenetic studies, and cellular physiology of a 69,XXX fetus, with comments on the biological effect of triploidy in man CHRISTINE M. GOSDEN, M. O. WRIGHT, W. G. PATERSON, and K. A. GRANT 371
- Structure and inheritance of some heterozygous Robertsonian translocations in man A. DANIEL and P. R. L. C. LAM-PO-TANG 381
- Association of D/D translocations with fetal wastage and aneuploidy. A report of four families PAUL M. FERNHOFF, D. N. SINGH, JAMES HANSON, SUZANNE TRUSLER, CHERYL R. DUMONT, and ANDREW T. L. CHEN 389
- Family study of inherited syndrome with multiple congenital deformities: symphalangism, carpal and tarsal fusion, brachydactyly, craniosynostosis, strabismus, hip osteochondritis V. VENTRUTO, R. DI GIROLAMO, B. FESTA, A. ROMANO, G. SEBASTIO, and L. SEBASTIO 394
- Short communication:**
- Linkage relations of locus for X-borne type of Charcot-Marie-Tooth muscular atrophy and that for Xg blood groups C. J. DE WEERDT, G. L. DANIELS, and PATRICIA TIPPETT 399
- Case reports:**
- Antenatal diagnosis of trisomy 13 with unexpected increase in alpha-feto protein J. S. FITZSIMMONS, G. M. FILSHIE, A. S. HILL, and R. KIME 400
- Discordance for Cornelia de Lange syndrome in twins ROGER E. STEVENSON and CHARLES I. SCOTT, Jr. 402
- The de Lange syndrome in one of twins GERSON CARAKUSHANSKY and C. BERTHIER 404
- Gardner's syndrome and steatocystoma multiplex. Two unusual genetically determined conditions occurring in same patient BARBARA LEPPARD and H. R. THOMPSON 407
- A giant short arm of no. 21 chromosome in mother of 21/21 translocation mongol E. TUNCBILEK, M. BOBROW, G. CLARKE, and K. TAYSI 411
- Book reviews 413