

Contents

Editorial page 385

Original articles

Osteogenesis imperfecta type IIA: evidence for dominant inheritance I D YOUNG, E M THOMPSON, C M HALL, AND M E PEMBREY *page 386*

Recurrence risks and prognosis in severe sporadic osteogenesis imperfecta E M THOMPSON, I D YOUNG, C M HALL, AND M E PEMBREY *page 390*

Prenatal prediction of osteogenesis imperfecta (OI type IV): exclusion of inheritance using a collagen gene probe P TSIPOURAS, R C SCHWARTZ, J D GOLDBERG, R L BERKOWITZ, AND F RAMIREZ *page 406*

Wolf-Hirschhorn locus is distal to *D4S10* on short arm of chromosome 4 C McKEOWN, A P READ, A DODGE, O STECKO, A MERCER, AND R HARRIS *page 410*

The fragile X syndrome in a large family. III Investigations on linkage of flanking DNA markers with the fragile site Xq27 H VEENEMA, N J CARPENTER, E BAKKER, M H HOFKER, A MILLINGTON WARD, AND P L PEARSON *page 413*

Syndrome of the month Postaxial acrofacial dysostosis (Miller) syndrome D DONNAL, H E HUGHES, AND R M WINTER *page 422*

Case reports

18q- syndrome and extraskelatal Ewing's sarcoma M M VALTUEÑA, J M GARCIA-SAGREDO, A M VILLA, C L GIMÉNEZ, AND J M A MEIX *page 426*

An isodicentric X chromosome with short arm fusion in a woman without somatic features of Turner's syndrome I C S BARNES, D J CURTIS, AND S L B DUNCAN *page 428*

Translocation X:13 in a patient with retinoblastoma G PONZIO, E SAVIN, G CATTANEO, M P GHIOTTI, A MARRA, O ZUFFARDI, AND C DANESINO *page 431*

Partial monosomy 12p13-1→13:3 D R ROMAIN, J GOLDSMITH, L M COLUMBANO-GREEN, C J CHAPMAN, R H SMYTHIE, AND R G PARFITT *page 434*

De novo 2q+ masquerading as Smith-Lemli-Opitz syndrome A E DONNENFELD, E H ZACKAI, D M McDONALD, R AQUINO, AND B S EMANUEL *page 436*

Incontinentia pigmenti in a boy with Klinefelter's syndrome A D ORMEROD, M I WHITE, E MCKAY, AND A W JOHNSTON *page 439*

Obituary page 442

Book reviews page 445

ASTM CODEN: JMDGAE (24) 385-448 (1987) ISSN 0022-2593

British Medical Association Tavistock Square London WC1