

Contents

Editorial page 145

Original articles

A reappraisal of the CHARGE association C A OLEY, M BARAITSER, AND D B GRANT *page 147*

Further delineation of the G syndrome: a manageable genetic cause of infantile dysphagia G N WILSON AND W J OLIVER *page 157*

Intellectual development in Apert's syndrome: a long term follow up of 29 patients M A PATTON, J GOODSHIP, R HAYWARD, AND R LANSDOWN *page 164*

Familial spastic paraplegia, bilateral sensorineural deafness, and intellectual retardation associated with a progressive nephropathy J S FITZSIMMONS, A R WATSON, D MELLOR, AND P R GUILBERT *page 168*

Immunodeficiency, centromeric heterochromatin instability of chromosomes 1, 9, and 16, and facial anomalies: the ICF syndrome P MARASCHIO, O ZUFFARDI, T DALLA FIOR, AND L TIEPOLO *page 173*

Family size limitation: a method for demonstrating recessive inheritance J F Y BROOKFIELD, R J POLLITT, AND I D YOUNG *page 181*

The frequency of consanguineous marriage among British Pakistanis A DARR AND B MODELL *page 186*

Absence of close linkage between benign hereditary chorea and the locus *D4S10* (probe G8) O W J QUARRELL, S YOUNGMAN, M SARFARAZI, AND P S HARPER *page 191*

Clinical, haematological, and genetic studies of type 2 normal Hb A₂ β thalassaemia A METAXOTOU-MAVROMATI, C KATTAMIS, L MATATHIA, M TZETIS, AND E KANAVAKIS *page 195*

Syndrome of the month Sclerosteosis P BEIGHTON *page 200*

Case reports

Hirschsprung disease associated with polydactyly, unilateral renal agenesis, hypertelorism, and congenital deafness: a new autosomal recessive syndrome H SANTOS, J MATEUS, AND M J LEAL *page 204*

Unusual dual genital duct remnants in true hermaphroditism C WILLIAMS AND I A HUGHES *page 206*

Short reports

De novo 10q23 interstitial deletion M A MORI, F GOMEZ-SABRIDO, A DIAZ DE BUSTAMANTE, I PINEL, AND M L MARTINEZ-FRIAS *page 209*

Interstitial deletion (6)(q11→q15) in an infant with congenital abnormalities H R SLATER, A ROBB, L A FORSYTH, D A HAMILTON, M C CLARK, AND C A S GALLOWAY *page 210*

Terminal deletion of chromosome 1(q43) in a female infant G P GARANI, L TAMISARI, S VOLPATO, AND V VIGI *page 211*

Correspondence page 213

Announcements page 194

ASTM CODEN: JMDGAE (25) 145-216 (1988) ISSN 0022-2593

British Medical Association Tavistock Square London WC1