

Journal of

MEDICAL GENETICS

Original articles

Case reports

Abstracts

Letters to the Editor

- 505 Immunohistochemical studies show truncated dystrophins in the myotubes of three fetuses at risk for Duchenne muscular dystrophy I B Ginjaar, E Bakker, MBM van Paassen, J T den Dunnen, A Wessels, E E Zubrzycka-Gaarn, A F M Moorman, G-J B van Ommen
- 511 Maternal uniparental disomy for chromosome 14 I K Temple, A Cockwell, T Hassold, D Pettay, P Jacobs
- 515 Age at onset in Huntington's disease: effect of line of inheritance and patient's sex RACRoos, M Vegter-van der Vlis, J Hermans, HMElshove, ACMoll, JJP van de Kamp, G W Bruyn
- 520 Linkage disequilibrium and recombination make a telomeric site for the Huntington's disease gene unlikely L Barron, A Curtis, A E Shrimpton, S Holloway, H May, R G Snell, DJ H Brock
- 523 Reproductive behaviour of families segregating for Cooley's anaemia before and after the availability of prenatal diagnosis M R Gamberini, M Lucci, C Vullo, B Anderson, R Canella, I Barrai
- 530 Single maxillary central incisor in a girl with del(18p) syndrome $D\mathcal{J}$ Aughton, A A AlSaadi; $D\mathcal{J}$ Transue
- 533 Interstitial deletion of chromosome 13: prognosis and adult phenotype J C S Dean, S Simpson, D A Couzin, G S Stephen
- 536 De novo ring chromosome 3: a new case with a mild phenotype M McKinley, A Colley, P Sinclair, D Donnai, T Andrews
- 539 De novo interstitial deletion of 1p (pter \rightarrow p34.1::p32.3 \rightarrow qter) M Yoshino, Y Watanabe, N Harada, K Abe
- 541 Pallister-Killian syndrome: additional manifestations of cleft palate and sacral appendage D R McLeod, L R Wesselman, D I Hoar
- 544 Facial dysmorphism: a marker of autosomal dominant cranial diabetes insipidus R B S Laing, J C S Dean, D W M Pearson, A W Johnston
- 547 Neurosonography and pathology in the Schinzel-Giedion syndrome A C Maclennan, D Doyle, R M Simpson
- 550 Abstracts of the meeting of the Clinical Genetics Society held on 28 November 1990 at the Institute of Education, London
- 562 Abstracts of the meeting of the Clinical Genetics Society held on 20 to 22 March 1991 at Belfast City Hospital
- 575 Cystic fibrosis screening and community genetics G Turner, B Wilcken, H Griffiths
 - 575 False positive results with immunoreactive trypsinogen screening for cystic fibrosis owing to trisomy 13 F J Priest, N C Nevin
 - 576 Notices

BRITISH MEDICAL ASSOCIATION TAVISTOCK SQUARE LONDON WC1H 9JR

J Med Genet: first published as on 1 August 1991. Downloaded from http://jmg.bmj.com/ on April 19, 2024 by guest. Protected by copyright