Contents

Review article
Molecular genetics of neurological tumours  R Y Chung, B R Seizinger  361

Original articles
Detection and characterisation of an overmodified type III collagen by analysis of non-cutaneous connective tissues in a patient with Ehlers-Danlos syndrome IV  L Nuytinck, P Narcisi, A Nicholls, J P Renard, F M Pope, A De Paepe  375
Linkage of autosomal dominant dystrophic epidermolysis bullosa in three British families to the marker D3S2 close to the COL7A1 locus  L Al-Imara, A J Richards, R A J Eady, I M Leigh, M Farrall, F M Pope  381
Limb-girdle type muscular dystrophy in a large family with distal myopathy: homozygous manifestation of a dominant gene?  B Uld  383
Confirmation of an association between RFLPs at the transforming growth factor-alpha locus and non-syndromic cleft lip and palate  S E Holder, G M Vintiner, B Farren, S Malcolm, R M Winter  390
No evidence of linkage between the transforming growth factor-alpha gene in families with apparently autosomal dominant inheritance of cleft lip and palate  G M Vintiner, S E Holder, R M Winter, S Malcolm  393
A complex rearrangement associated with sex reversal and the Wolf-Hirschhorn syndrome: a cytogenetic and molecular study  K Coles, M Mackenzie, J Crolla, J Harvey, J Starr, F Howard, P Jacobs  400
Distal 8p deletion (8p23.1→8pter): a common deletion?  R Hutchinson, M Wilson, L Voullaire  407

Syndrome of the month
Angelman syndrome  J Clayton-Smith, M E Pembrey  412

Case reports
Ovarian cancer family and prophylactic choices  D G R Evans, G Ribiero, D Warrell, D Donnai  416
Acromegalo id facial appearance (AFA) syndrome: report of a second family  B Dallapiccola, L Zelante, L Accadina, R Mingarelli  419
An infant with multiple congenital abnormalities and biochemical findings suggesting a variant of galactosialidosis  B Soy, F A Hommes, S A Malik, N J Carpenter  423
Cardiofaciocutaneous syndrome with new ectodermal manifestations  P D Tumpenny, J C S Dean, I A Auchterlonie, A W Johnston  428
Recurrence of orbital cysts in the branchio-oculo-facial syndrome  D W Fielding, A E Fryer  430
Terminal 22q deletion associated with a partial deficiency of arylsulphatase A  K Narahara, Y Takahashi, M Murakami, K Tsuji, Y Yokoyama, R Murakami, S Ninomiya, Y Seino  432
Two sibs with Wiedemann-Rautenstrauch syndrome: possibilities of prenatal diagnosis by ultrasound  G Castiheyr, M Panol, H L Presas, E Goldschmidt, J M Sanchez  434

Short report
The demonstration of monozygosity in twins discordant for sacral agenesis  M d'A Crawford, J Cheshire, T M Wilson, C R J Woodhouse  437

Abstracts
Medical genetics: advances in brief  439

Letter to the Editor
Palmoplantar hyperkeratosis and deafness  J Verbov  440

Book reviews
440

Notice
440

BMJ PUBLISHING GROUP  TAVISTOCK SQUARE  LONDON WC1H 9JR