Contents

Review article
225 Genes other than BRCA1 and BRCA2 involved in breast cancer susceptibility
M M de Jong, I M Nolte, G J te Meerman, W T A van der Graaf, J C Oosterwijk, J H Kleibeuker, M Schaapveld, E G E de Vries

Commentaries
243 Assessing epidemiological evidence for the teratogenic effects of anticonvulsant medications
H Doll, P McElhatton
245 The teratogenicity of anticonvulsant drugs: a progress report
L B Holmes
248 Antiepileptic drug therapy during pregnancy: the neurologist’s perspective
S Shorvon

Original articles
251 Long term health and neurodevelopment in children exposed to antiepileptic drugs before birth
J C S Dean, H Hailey, S J Moore, D J Lloyd, P D Tumpey, J Little
260 Specific haplotypes of the RET proto-oncogene are over-represented in patients with sporadic papillary thyroid carcinoma
F Lesueur, M Corbex, J D McKay, J Lima, P Soares, P Grisenti, J Burgess, J Ceccherrini, S Landolfi, M Papotti, A Auronni, D E Goliga, G Romeo
266 Automated fluorescent genotyping detects 10% of cryptic subtelomeric rearrangements in idiopathic syndromic mental retardation
M Rio, F Molinari, S Heuerz, C Ozilou, P Gossot, O Roaul, V Commer-Daure, J Amiel, S Lyonnet, M le Menter, C Turleau, M-C de Blis, M Prieur, S Romana, M Vekemans, A Munnich, J Colleaux

Short report
271 Identification of mutations in the gene encoding sterol regulatory element binding protein (SREBP)-2 in hypercholesterolaemic subjects
P Y Muller, A R Miserez

Letters to JMG
276 A new MRXS locus maps to the X chromosome pericentromeric region: a new syndrome or narrow definition of Sutherland-Haan genetic locus?
M Fichera, E Borgione, E Avola, S Amata, M Sturino, C Romano, A Ragusa
281 Distinctive audiometric features between USH2A and USH2B subtypes of Usher syndrome
M Hmani-Aifa, S Ben Arab, K Kharrat, D J Orten, A Boulia-Eggied, M Dina, S Machida, W J Kimberling, H Ayadi
284 Dominant X linked retinitis pigmentosa is frequently accounted for by truncating mutations in exon ORF15 of the RPGR gene
286 Identification of cryptic splice site, exon skipping, and novel point mutations in type I CD36 deficiency
H Hanawa, K Watanabe, T Nakamura, Y Ogawa, K Toba, I Fuse, M Kodama, K Kato, K Fuse, Y Aizawa
292 Four novel mutations in the OFD1 (Coxorf1) gene in Finnish patients with oral-facial-digital syndrome 1
A Rakkolainen, S Ala-Mello, P Kristo, A Orpana, I Jarvela
297 Genetic characterisation of patients with multiple colonic polyps
C Albuquerque, M Cravo, C Cruz, P Lage, P Chaves, P Fidalgo, A Suspicio, C Nobre Leitao

Echoes
242 Another risk factor for MI?
259 Gene carriers face other risks
291 Limited role for HLA DR15 in MS
296 Genes predict outcome in multiple sclerosis
302 Cancers of the small and large bowel originate differently

303 Book reviews
303 Correction
304 Instructions for authors
contd…