

Letters to JMG

- 244** Analysis of mtDNA variant segregation during early human embryonic development: a tool for successful NARP preimplantation diagnosis *J Steffann, N Frydman, N Gigarel, P Burlet, P F Ray, R Fanchin, E Feyereisen, V Kerbrat, G Tachdjian, J-P Bonnefont, R Frydman, A Munnich*
- 248** Congenital hyperinsulinism and mosaic abnormalities of the ploidy *I Giurgea, D Sanlaville, J-C Fournet, C Sempoux, C Bellanné-Chantelot, G Touati, L Hubert, M-S Groos, F Brunelle, J Rahier, J-C Henquin, M J Dunne, F Jaubert, J-J Robert, C Nihoul-Fékété, M Vekemans, C Junien, P de Lonlay*
- 255** Aortic aneurysmal disease and cutis laxa caused by defects in the elastin gene *Z Szabo, M W Crepeau, A L Mitchell, M J Stephan, R A Puntel, K Yin Loke, R C Kirk, Z Urban*
- 259** Spastin mutations are frequent in sporadic spastic paraparesis and their spectrum is different from that observed in familial cases *C Depienne, C Tallaksen, J Y Lephay, B Bricka, S Poëa-Guyon, B Fontaine, P Labauge, A Brice, A Durr*
- 266** Two high throughput technologies to detect segmental aneuploidies identify new Williams-Beuren syndrome patients with atypical deletions *C Howald, G Merla, M C Digilio, S Amenta, R Lyle, S Deutsch, U Choudhury, A Bottani, S E Antonarakis, H Fryssira, B Dallapiccola, A Reymond*
- 274** A mutation in the hair matrix and cuticle keratin KRTHB5 gene causes ectodermal dysplasia of hair and nail type *M Naeem, M Wajid, K Lee, S M Leal, W Ahmad*
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- 285** Age associated increase in the prevalence of chromosome 22q loss of heterozygosity in histological subsets of benign meningioma *M E Baser, T Y Poussaint*

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- 288** Genetic Disorders of the Indian Subcontinent *Y Yamada*

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linked to a 1.7 Mb region on chromosome 2p13.3-14 J Warner, D R Nyholt, F Busfield, M Epstein, J Burgess, S Stranks, P Hill, D Perry-Keene, D Learoyd, B Robinson, B T Teh, J B Prins, J W Cardinal

- e13** Mapping of hereditary mixed polyposis syndrome (HMPS) to chromosome 10q23 by genomewide high-density single nucleotide polymorphism (SNP) scan and identification of *BMPRIA* loss of function X Cao, K W Eu, M P Kumarasinghe, H H Li, C Loi, P Y Cheah