

Review article

Familial gastric cancer: overview and guidelines for management C Caldas, F Carneiro, H T Lynch, J Yokota, G L Wiesner, S M Powell, F R Lewis, D G Huntsman, P D P Pharoah, J A Jankowski, P MacLeod, H Vogelsang, G Keller, K G M Park, F M Richards, E R Maher, S A Gayther, C Oliveira, N Grehan, D Wight, R Seruca, F Raviello, B A J Ponder, C E Jackson 873

Original articles

- Mutations of the cathepsin C gene are responsible for Papillon-Lefèvre syndrome T C Hart, P S Hart, D W Bowden, M D Michalec, S A Collison, S J Walker, Y Zhang, E Firatli 881
 Popliteal pterygium syndrome: a clinical study of three families and report of linkage to the Van der Woude syndrome locus on 1q32 M M Lees, R M Winter, S Malcolm, H M Saal, L Chitty 888
 Unusual clustering of brain tumours in a family with NF1 and variable expression of cutaneous features F Faravelli, M Upadhyaya, M Osborn, S M Huson, R Hayward, R Winter 893
 Psychological functioning before predictive testing for Huntington's disease: the role of the parental disease, risk perception, and subjective proximity of the disease M Decruyenaere, G Evers-Kiebooms, A Boogaerts, J J Cassiman, T Cloostermans, K Demeyttereire, R Dom, J P Fryns 897
 Presymptomatic testing for BRCA1 and BRCA2: how distressing are the pre-test weeks? L N Lodder, P G Frets, R W Trisburg, E J Meijers-Heijboer, J G M Klijn, H J Duivenvoorden, A Tibben, A Wagner, C A van der Meer, P Devilee, C J Cornelisse, M F Niermeijer, and other members of the Rotterdam/Leiden Genetics Working Group 906
 Neocentromere formation in a stable ring 1p32-p36.1 chromosome H R Slater, S Nouri, E Earle, A W I Lo, L G Hale, K H A Choo 914

Short reports

- A proven de novo germline mutation in HNPCC C Kraus, S Kastl, K Günther, S Klessinger, W Hohenberger, W G Ballhausen 919
 Mutational analysis of the HGO gene in Finnish alkaptonuria patients D Beltrán-Valero de Bernabé, P Peterson, K Luopajarvi, P Matintalo, A Alho, Y Konttinen, K Krohn, S Rodriguez de Córdoba, A Ranki 922
 Skewed sex ratios in familial holoprosencephaly and in people with isolated single maxillary central incisor G Suthers, S Smith, S Springbett 924

Letters to the Editor

- Leigh syndrome transmitted by uniparental disomy of chromosome 9 V Tiranti, E Lamantea, G Uziel, M Zeviani, P Gasparini, R Marzella, M Rocchi, M Fried 927
 A case of Williams syndrome with a large, visible cytogenetic deletion Y-Q Wu, E Nickerson, L G Shaffer, K Keppler-Noreuil, A Mullenberg 928
 First molecular evidence for a de novo mutation in RS1 (XLRS1) associated with X linked juvenile retinoschisis A Gehrig, B H F Weber, B Lorenz, M Andressi 932
 Pathogenicity of homoplasmic mitochondrial DNA mutation and nuclear gene involvement M Odawara, H Maki, N Yamada 934
 Identification and clinical presentation of β thalassaemia mutations in the eastern region of Saudi Arabia E-H A El-Harith, W Kühnau, J Schmidtke, M Stuhmann, Z Nassarallah, A Al-Shahri 935
 Rapid screening for the most common β thalassaemia mutations in south east Asia by PCR based restriction fragment length polymorphism analysis (PCR-RFLP) P Pramoonjago, A Harahap, R A Taufani, I Setianingsih, S Marzuki 937
 Clinical and molecular findings in a patient with a deletion on the long arm of chromosome 12 A F Brady, M M Elsawi, C R Jamieson, K Marks, S Jeffery, M A Patton, L Murtaza, M O Savage 939
 Amelogenesis imperfecta, sensorineural hearing loss, and Beau's lines: a second case report of Heimler's syndrome M Tischkowitz, C Clenaghan, S Davies, L Hunter, J Potts, S Verhoeven 941
 PTEN and LKB1 genes in laryngeal tumours R W Chen, E Avizienyte, S Roth, I Leivo, A A Mäkitie, L-M Aaltonen, L A Aaltonen 943
 Index 945

