

# **Contents**



SUBMISSION

GO TO

TO SUBMIT YOUR

MANUSCRIP

WEBSIT

## Review article

Hereditary paraganglioma targets diverse paraganglia B E Baysal

## **Original articles**

- 623 Fraser syndrome and cryptophthalmos: review of the diagnostic criteria and evidence for phenotypic modules in complex malformation syndromes A M Slavotinek, C J Tifft
- 634 Identification of a locus for a form of spondyloepiphyseal dysplasia on chromosome 15q26.1: exclusion of aggrecan as a candidate gene S Eyre, P Roby, K Wolstencroft, K Spreckley, R Aspinwall, R Bayoumi, L Al-Gazali, R Ramesar, P Beighton, G Wallis
- Apolipoprotein E4 is only a weak 639 predictor of dementia and cognitive decline in the general population A G Yip, C Brayne, D Easton, D C Rubinsztein, the Medical Research Council Cognitive Function and Ageing Study (MRC CFAS)
- 644 Identification of candidate lung cancer susceptibility genes in mouse using oligonucleotide arrays W J Lemon, H Bernert, H Sun, Y Wang, M You

## **Short reports**

- Mapping of a novel locus for achromatopsia (ACHM4) to 1p and identification of a germline mutation in the  $\alpha$  subunit of cone transducin (GNAT2) I A Aligianis, T Forshew, S Johnson, M Michaelides, C A Johnson, R C Trembath, D M Hunt, A T Moore, E R Maher
- Vitreoretinopathy with phalangeal epiphyseal dysplasia, a type II collagenopathy resulting from a novel mutation in the C-propeptide region of the molecule A.J. Richards, J. Morgan, PWP Bearcroft, EPickering, MJ Owen, P Holmans, N Williams, C Tysoe, F M Pope, M P Snead, H Hughes

#### Letters to JMG

September 2002

- 666 Spondyloepimetaphyseal dysplasia with multiple dislocations (Hall type): three further cases and evidence of autosomal dominant inheritance CM Hall, N H Elcioglu, K D MacDermot, A C Offiah, R M Winter
- 671 Familial chronic nail candidiasis with ICAM-1 deficiency: a new form of chronic mucocutaneous candidiasis D Zuccarello, D C Salpietro, S Gangemi, V Toscano, M V Merlino, S Briuglia, G Bisignano, M Mangino, R Mingarelli, B Dallapiccola
- Expression of wild type and mutant 676 TSC2, but not TSC1, causes an increase in the G1 fraction of the cell cycle in HEK293 cells L Khare, A Astrinidis, W Senapedis, P D Adams, E Petri Henske
- 681 Mutation in KCNQ1 that has both recessive and dominant characteristics A Murray, F Potet, C Bellocq, I Baró, W Reardon, H E Hughes, S Jeffery
- 686 Heterozygous truncating mutation in the human homeobox gene GSH2 has no discernable phenotypic effect J G Dauwerse, C E M de Die-Smulders, E Bakker, M H Breuning, D J M Peters
- 689 Coping style, psychological distress, risk perception, and satisfaction in subjects attending genetic counselling for hereditary cancer K Nordin, A Lidén, M Hansson, R Rosenquist, G Berglund
- 695 Genetic counselling for familial breast and ovarian cancer in Ontario A Andermann, S A Narod
- 697 The use of audiotapes in consultations with women from high risk breast cancer families: a randomised trial E Lobb, P Butow, B Meiser, A Barratt, J Kirk, M Gattas, E Haan, K Tucker
- 704 Instructions for authors

### **Electronic letters**

Parental attitude towards genetic testing for familial hypercholesterolaemia in children M A W Umans-Eckenhausen, F | Oort, K C M P Ferenschild, J C Defesche, J.J.P. Kastelein, J.C.J.M. de Haes

contd...