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Cover credit: Deep intronic splicing mutation in ACVRL1 as a cause of HHT, from Wooderchak-Donahue WL *et al* (see page 826).

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Contents

December 2018 Volume 55 Issue 12

Cancer genetics

785 Investigation of clinically relevant germline variants detected by next-generation sequencing in patients with childhood cancer: a review of the literature
D E Sylvester, Y Chen, R V Jamieson, L Dalla-Pozza, J A Byrne

794 MCQs Reclassification of *BRCA1* and *BRCA2* variants of uncertain significance: a multifactorial analysis of multicentre prospective cohort
J-S Lee, S Oh, S K Park, M-H Lee, J W Lee, S-W Kim, B H Son, D-Y Noh, J E Lee, H-L Park, M J Kim, S I Cho, Y K Lee, S S Park, M-W Seong

Neurogenetics

803 EDITOR'S CHOICE PEHO syndrome: the endpoint of different genetic epilepsies
M Chitre, M S Nahorski, K Stouffer, B Dunning-Davies, H Houston, E L Wakeling, A F Brady, S M Zuberi, M Suri, A P J Parker, C G Woods

814 Characterising the phenotype and mode of inheritance of patients with inherited peripheral neuropathies carrying MME mutations
V Lupo, M Frasquet, A Sánchez-Monteagudo, A L Pelayo-Negro, T García-Sobrino, M J Sedano, J Pardo, M Misiego, J García-García, M J Sobrido, M D Martínez-Rubio, M J Chumillas, J J Vílchez, J F Vázquez-Costa, C Espinós, T Sevilla

Diagnostics

824 Genome sequencing reveals a deep intronic splicing *ACVRL1* mutation hotspot in Hereditary Haemorrhagic Telangiectasia
W L Wooderchak-Donahue, J McDonald, A Farrell, G Akay, M Velinder, P Johnson, C VanSant-Webb, R Margraf, E Briggs, K J Whitehead, J Thomson, A E Lin, R E Pyeritz, G Marth, P Bayrak-Toydemir

Genome-wide studies

831 Common variants at 5q33.1 predispose to migraine in African-American children
X Chang, R Pellegrino, J Garifallou, M March, J Snyder, F Mentch, J Li, C Hou, Y Liu, P M A Sleiman, H Hakonarson

Developmental defects

837 Specific combinations of biallelic *POLR3A* variants cause Wiedemann-Rautenstrauch syndrome
S Paolacci, Y Li, E Agolini, E Bellacchio, C E Arboleda-Bustos, D Carrero, D Bertola, L Al-Gazali, M Alders, J Altmüller, G Arboleda, F Beleggia, A Bruselles, A Ciolfi, G Gillessen-Kaesbach, T Krieg, S Mohammed, C Müller, A Novelli, J Ortega, A Sandoval, G Velasco, G Yigit, H Arboleda, C Lopez-Otin, B Wollnik, M Tartaglia, R C Hennekam

Miscellaneous

846 Correction: *A common SLC26A4-linked haplotype underlying non-syndromic hearing loss with enlargement of the vestibular aqueduct*

Epigenetics

847 Paternal uniparental disomy of chromosome 19 in a pair of monochorionic diamniotic twins with dysmorphic features and developmental delay
K S Yeung, M S P Ho, S L Lee, A S Y Kan, K Y K Chan, M H Y Tang, C C Y Mak, G K C Leung, P L So, R Pfundt, C R Marshall, S W Scherer, S Choufani, R Weksberg, B Hon-Yin Chung