

BRAIN

A JOURNAL OF NEUROLOGY

Volume 143 Part 4 April 2020

Editorial

1045 Editorial. D. M. Kullmann

Scientific Commentaries

1046 Predicting incidences of neurodevelopmental disorders. J. R. Lemke

1048 All in the numbers? Computational prediction of optimal anaesthetic weaning in status epilepticus. S. Rüegg and R. Sutter

1050 Improving diagnosis and prognosis in disorders of consciousness. A. M. Owen

1053 Atrophy network mapping of transdiagnostic cognitive and neuropsychiatric symptoms. M. Veldsman

Review Articles

1057 Decoding the relationship between ageing and amyotrophic lateral sclerosis: a cellular perspective. V. A. Pandya and R. Patani

1073 Lipid metabolic pathways converge in motor neuron degenerative diseases. O. J. Rickman, E. L. Baple and A. H. Crosby

Update

1088 Revisiting 'brain modes' in a new computational era: approaches for the characterization of brain-behavioural associations. M. N. Toba, O. Godefroy, R. J. Rushmore, et al.

Reports

1099 A catalogue of new incidence estimates of monogenic neurodevelopmental disorders caused by *de novo* variants. J. A. López-Rivera, E. Pérez-Palma, J. Symonds, et al.

1106 Structural and functional footprint of visual snow syndrome. C. J. Shanks, F. H. Maniyan, D. E. Chou, et al.

Original Articles

1114 Novel congenital disorder of O-linked glycosylation caused by GALNT2 loss of function. M. Zilmer, A. C. Edmondson, S. A. Khetarpal, et al.

1127 Protective effects of 4-aminopyridine in experimental optic neuritis and multiple sclerosis. M. Dietrich, V. Koska, C. Hecker, et al.

1143 Electrophysiological predictors of successful weaning from anaesthetics in refractory status epilepticus. D. B. Rubin, B. Angelini, M. Shoukat, et al.

1158 Distinct patterns of structural damage underlie working memory and reasoning deficits after traumatic brain injury. A. E. Jolly, G. T. Scott, D. J. Sharp, et al.

1177 Prognosis for patients with cognitive motor dissociation identified by brain-computer interface. J. Pan, Q. Xie, P. Qin, et al.

1190 Variants in saposin D domain of prosaposin gene linked to Parkinson's disease. Y. Oji, T. Hatano, S.-I. Ueno, et al.

1206 White matter basis for the hub-and-spoke semantic representation: evidence from semantic dementia. Y. Chen, L. Huang, K. Chen, et al.

1220 Plasma tau, neurofilament light chain and amyloid- β levels and risk of dementia; a population-based cohort study. F. deWolf, M. Ghanbari, S. Licher, et al.

1233 Medial temporal lobe connectivity and its associations with cognition in early Alzheimer's disease. D. Berron, D. van Westen, R. Ossenkoppele, et al.

1249 Network localization of clinical, cognitive, and neuropsychiatric symptoms in Alzheimer's disease. A. M. Tetreault, T. Phan, D. Orlando, et al.

1261 Impaired theta phase coupling underlies frontotemporal dysconnectivity in schizophrenia. R. A. Adams, D. Bush, F. Zheng, et al.

Dorsal Column

Grey Matter

1278 The Arc de Siècle: functional neurological disorder during the 'forgotten' years of the 20th century. M. Fend, L. Williams, A. J. Carson, et al.

Letters to the Editor

e25 The repeat variant in *MSH3* is not a genetic modifier for spinocerebellar ataxia type 3 and Friedreich's ataxia. W. Y. Yau, M. Raposo, C. Bettencourt, et al.

e26 Reply: The repeat variant in *MSH3* is not a genetic modifier for spinocerebellar ataxia type 3 and Friedreich's ataxia. M. Flower, V. Lomeikaite, P. Holmans, et al.

e27 Interactions of interictal epileptic discharges with sleep slow waves and spindles. N. von Ellenrieder, A. Koupparis and J. Gotman

e28 Reply: Interactions of interictal epileptic discharges with sleep slow waves and spindles. P. Dahal, N. Ghani, A. Flinker, et al.

e29 Recurrent bi-allelic splicing variant c.454+3A>G in *TRAPPC4* is associated with progressive encephalopathy and muscle involvement. P. Kaur, R. Kadavigere, K. M. Girisha, et al.

e30 Reply: Recurrent bi-allelic splicing variant c.454+3A>G in *TRAPPC4* is associated with progressive encephalopathy and muscle involvement. N. J. Van Bergen and J. Christodoulou

e31 *RSRC1* loss-of-function variants cause mild to moderate autosomal recessive intellectual disability. M. Scala, M. Mojarad, S. Riazuddin, et al.

e32 Corrigenda

Evoked Response

e34 THC and CBD: is medical cannabis overhyped or under-prescribed? L. Lyon



OXFORD
UNIVERSITY PRESS