Early prognosis

Familial

Persistence

Some reflections

Cu/Zn superoxide dismutase gene mutations in amyotrophic lateral sclerosis: correlation between genotype and clinical features A Radunovic, P N Leigh on behalf of the European Familial ALS Group

Massive

Spinal

Acute

Disordered

Changes

Neuropsychological stability

Association between interictal mood and acute epilepsy J H Rees, H J Willison

Bilateral total deafness due to pontine haematoma C A Egan, L Davies, G M Halmagyi

Spinal cord MRI in multiple sclerosis with multicoil arrays: a comparison between fast spin echo and fast FLAIR M Filippi, T A Youory, H Akhadia, M Stehling, M A Horsfield, R Volz

Massive nerve root enlargement in chronic inflammatory demyelinating polyneuropathy W Schudy, P J Goulding, B R F Lecky, R H M King, G M L Smith

Changes in spinal cord excitability in a patient with rhythmic segmental myoclonus V Di Lazzaro, D Rostuccia, R Nardona, A Oliviero, P Profice, A Imola, P Tomasi, J C Rothwell

Disordered axial movement in Parkinson’s disease M J Steiger, P D Thompson, C D Marsden

Acute oropharyngeal palsy is associated with antibodies to Q01b and GT1a gangliosides C P O’Leary, J Veitch, W F Durward, A M Thomas, J H Rees, H J Willison

Persistence of MRI changes in central pontine myelinolysis P Maddison, P K Newman, N Bradby

Brain MRI, lumbar CSF monoamine concentrations, and clinical descriptors of patients with spinocerebellar ataxia mutations J J Higgins, J D Harvey-White, L E Neo, M J Colli, T A Grossi, I J Kogan

Double cortical stimulation in amyotrophic lateral sclerosis T Yokota, A Yoshino, A Inaba, Y Saito

Intercital mood and personality disorders in temporal lobe epilepsy and juvenile myoclonic epilepsy G I Perini, C Tonin, G Carraro, G Bernasconi, M P Canevini, R Canger, A Pellegrini, G Testa

Association between HIV distal symmetric polyneuropathy and mycobacterium avium complex infection G R Norton, J Sweeney, D Marriott, M G Law, B J Brew

Early prognosis in coma after cardiac arrest: a prospective clinical, electrophysiological, and biochemical study of 60 patients C Bassetti, F Bonini, J Mathis, C W Hess

Familial hemiplegic migraine in the west of Scotland: a clinical and genetic study of seven families M A S Ahmed, E Reid, A Cooke, R Arragrinsson, J L Tomie, J B P Stephenson

Neuropsychological stability over two years in asymptomatic carriers of the Huntington’s disease mutation J R Campodinico, A M Codori, J Brandt

Huntington’s disease: confirmation of diagnosis and presymptomatic testing in Spanish families by genetic analysis A Sanchez, S Castelleti-Bel, M Mila, D Genis, M Calopa, D Jimenez, X Estivill

Bilateral total deafness due to pontine haematoma C A Egan, L Davies, G M Halmagyi