
Imaging the head: functional imaging G V Sawle


Cerebral blood flow and metabolism in children with severe head injuries. Part 2: cerebrovascular resistance and its determinants P M Sharplcs, D S F Matthews, J A Eyre

Clinicopathological study of 35 cases of multiple system atrophy G K Wenning, Y Ben-Shlomo, M Magalhaes, S E Daniel, N P Quin

Progressive supranuclear palsy: neuropathologically based diagnostic clinical criteria S J Collins, J E Abikog, J E Parisi, D M Maraganore

Neuropsychological pattern of striatonigral degeneration: comparison with Parkinson's disease and progressive supranuclear palsy B Pillon, N Goeuder-Khouja, B Desoer, M Vidaliher, C Malapane, B Dutoit, Y Agid


Advance information and movement in gilles de la Tourette's syndrome N Georgiou, J L Bradasht, J G Phillips, J A Bradasht, E Chu

Impaired prepulse inhibition of acoustic and tactile startle response in patients with Huntington's disease N R Steedlow, J Paulson, D L Braff, N Batters, M A Goyer, M R Swenson

Continuous response variable trial design in motor neuron disease: long term treatment with a TRH analogue (RX77368) A Goometiltes, R J Graifoff

Rhabdomyolysis and acute encephalopathy in late onset medium chain acyl-CoA dehydrogenase deficiency W Ruitenberg, P J E Ploos, D M Turnbull, B Garavaglia, R A Chalmers, R W Taylor, J M Gabriels

Phoedrine: a substitiute for hydroxymphetamine as a diagnostic eyedrop test in Horner's syndrome A T Bates, S Chamberlain, M Champion, I L Foley, E Hughes, B Jami, H Mehta, S B Smith

Loss of heterozygosity for DNA polymorphisms mapping to chromosomes 10 and 17 and prognosis in patients with giomas C E Jones, M B Davis, J L Darling, J F Geddes, D G T Thomas, A E Harding

Partial restoration of blink: reflex function after spinal accessory-facial nerve anastomosis N Danziger, B Chassande, G Lams, I Fligny, J Soudant, J C Willer

Thrombolytic treatment for acute occlusion of the basilar artery M Huemer, V Niederwisser, G Ladurner

A missense point mutation (Ser515Phe) in the adenomleukodystrophy gene in a family with adrenoleukodystrophy: a clinical, biochemical, and genetic study M Vorgerd, S Fuchs, M Tengenthoff, J-P Malin

Treatment of chronic limb spasticity with botulinum toxin A J W Dunne, N Heye, S L Dunne

"Pseudo" hypertrophic neuropathy of childhood M Baba, H Takada, H Miuara, T Okushima, M Matsuagree

Erdheim-Chester disease and slowly progressive cerebellar dysfunction T Fukazawa, E Tsukihima, H Sasaki, K Hamada, T Hamada, K Tashiro

Miller Fisher-Guillain-Barre overlap syndrome with enhancing lesions in the spinocerebellar tracts M Urashitani, T Usaka, M Kamayama