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M. Mizuguchi (Japan)

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### New guidelines for management of febrile seizures in Japan

J. Natsume, S.-i. Hamano, K. Iyoda, H. Kanemura, M. Kubota, M. Mimaki, S. Nijijima, T. Tanabe, H. Yoshinaga, N. Kojimahara, H. Komaki, K. Sugai, T. Fukuda, Y. Maegaki, H. Sugie (Japan)

### Prenatal irradiation-induced brain neuropathology and cognitive impairment

B. Yang, B.X. Ren, F.R. Tang (People's Republic of China, Singapore)

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### The pathology of incipient polymicrogyria

P. Diamandis, D. Chitayat, A. Toi, S. Blaser, P. Shannon (Canada)

### Longitudinal change in white matter in preterm infants without magnetic resonance imaging abnormalities: Assessment of serial diffusion tensor imaging and their relationship to neurodevelopmental outcomes

S. Kidowaki, M. Morimoto, K. Yamada, K. Sakai, M. Zuiki, H. Maeda, S. Yamashita, T. Morita, T. Hasegawa, T. Chiyonobu, S. Tokuda, H. Hosoi (Japan)

### Efficacy of bezafibrate on fibroblasts of glutaric acidemia type II patients evaluated using an *in vitro* probe acylcarnitine assay

K. Yamada, H. Kobayashi, R. Bo, J. Purevsuren, Y. Mushimoto, T. Takahashi, Y. Hasegawa, T. Taketani, S. Fukuda, S. Yamaguchi (Japan)

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### A Japanese case of $\beta$ -ureidopropionase deficiency with dysmorphic features

T. Akiyama, T. Shibata, H. Yoshinaga, T. Kuhara, Y. Nakajima, T. Kato, Y. Maeda, M. Ohse, M. Oka, M. Kageyama, K. Kobayashi (Japan)

### Cerebral arteriopathy associated with heterozygous Arg179Cys mutation in the *ACTA2* gene: Report in 2 newborn siblings

J. de Grazia, I. Delgado, A. Sanchez-Montanez, S. Boronat, M. del Campo, E. Vazquez (Spain)

### Epileptic phenotype of *FGFR3*-related bilateral medial temporal lobe dysgenesis

T. Okazaki, Y. Saito, R. Ueda, T. Awashima, Y. Nishimura, I. Yuasa, Y. Shinohara, K. Adachi, M. Sasaki, E. Nanba, Y. Maegaki (Japan)

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- Infantile neuroaxonal dystrophy and *PLA2G6*-associated neurodegeneration: An update for the diagnosis  
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