

Author Index to Volume 37

- Abe, S., see Okumura, A. 978
- Abe, S., see Nakazawa, M. 864
- Abe, S., see Okumura, A. 339
- Abe, Y., Terashima, H., Hoshino, H., Sassa, K., Sakai, T., Ohtake, A., Kubota, M., and Yamanouchi, H. Characteristic MRI features of chronic inflammatory demyelinating polyradiculoneuropathy 894
- Aburano, H., see Ueda, F. 439
- Achour, N.B., see Mrabet, S. 904
- Adachi, M., see Shimada, S. 515
- Agan, K., see Kilinc, O. 352
- Agematsu, K., see Motobayashi, M. 140
- Aiba, K., see Yoshitomi, S. 874
- Aida, N., see Takano, K. 638
- Aida, N., see Niwa, T. 967
- Akasaka, M., see Nishimura, T. 808
- Akasaka, M., see Nakazawa, M. 418
- Akasaka, M., see Kamei, A. 137
- Akasaka, M., see Saitoh, M. 463
- Akasaka, N., see Magara, S. 725
- Akasaka, N., see Kobayashi, Y. 356
- Akashi, N., see Saito, Y. 175
- Akiyama, T., see Yoshinaga, H. 13
- Akiyama, T., see Kobayashi, K. 487
- Aksu, B., see Kurtcan, S. 546
- Alkan, A., see Kurtcan, S. 546
- Altunbasak, S., see Inceci, F. 311
- Altunbasak, S., see Inceci, F. 803
- Altunbasak, S., see Inceci, F. 66
- Ameilia, A., Shatriah, I., Wan-Hitam, W.H., and Yunus, R. A unilateral optic perineuritis in a teenager – A case report 635
- Amemiya, K., see Saitoh, M. 463
- Anai, C., see Egami, C. 840
- Ando, N., see Okanishi, T. 423
- Ando, S., see Kuwabara, H. 130
- Anzai, R., see Takano, K. 638
- Anzai, Y., see Saitoh, M. 463
- Aoyama, Y., see Vatanavicharn, N. 698
- Arai, E., see Okumura, A. 978
- Arai, H., see Kitai, Y. 758
- Araki, A., Ikegami, M., Okayama, A., Matsumoto, N., Takahashi, S., Azuma, H., and Takahashi, M. Improved prefrontal activity in AD/HD children treated with atomoxetine: A fNIRS study 76
- Araki, A., see Yamamoto, S. 829
- Araki, T., see Kuwabara, H. 130
- Araya, N., see Kamei, A. 137
- Arrigoni, F., see Romaniello, R. 273
- Arsenault, L., see Rebour, R. 643
- Asami, M., see Kamei, A. 137
- Ashida, A., see Kashiwagi, M. 618
- Awan, I., see Puppala, B. 206
- Awaya, T., see Ide, M. 825
- Azuma, H., see Yamamoto, S. 829
- Azuma, H., see Araki, A. 76
- Azuma, J., Nabatame, S., Nakano, S., Iwatani, Y., Kitai, Y., Tominaga, K., Kagitani-Shimono, K., Okinaga, T., Yamamoto, T., Nagai, T., and Ozono, K. Prognostic factors for acute encephalopathy with bright tree appearance 191
- Baba, S., see Mohammadi, M. 59
- Baba, S., see Saito, Y. 175
- Badilla-Porras, R., see Munoz, T. 168
- Bakar, E.E., see Değerliyurt, A. 250
- Baldassarri, M., see Mari, F. 527
- Balestri, P., see Mari, F. 527
- Bassi, M.T., see Romaniello, R. 273
- Batnini, S., and Uno, A. Investigation of basic cognitive predictors of reading and spelling abilities in Tunisian third-grade primary school children 579
- Baykan, N., see Ozcan, U.A. 495
- Ben Rhouma, H., see Rebai, I. 153
- Ben Rhouma, H., see Mrabet, S. 904
- Ben Youssef-Turki, I., see Rebai, I. 153
- Berg, A.T., see Lee, Y.-J. 37
- Bernasconi, P., see Fusco, C. 891
- Borgatti, R., see Romaniello, R. 273
- Boulay, C., see Cabasson, S. 943
- Boyd, S.G., see Spagnoli, C. 704
- Bozlu, G., Cobanogullari Direk, M., and Okuyaz, C. Subacute sclerosing panencephalitis with parkinsonian features in a child: A case report 901
- Brittain, H., see Spagnoli, C. 704
- Briyal, S., see Puppala, B. 206
- Broussolle, E., see Rebour, R. 643
- Byun, C.K., see Lee, J.S. 402
- Cabasson, S., Tardieu, M., Meunier, A., Rouanet-Larriviere, M.-F., Boulay, C., and Pedespan, J.-M. Childhood CIDP: Study of 31 patients and comparison between slow and rapid-onset groups 943
- Caffarelli, C., see Mari, F. 527
- Canali, E., see Fusco, C. 891
- Cao, A., see Ma, L. 387
- Cao, Z., see Han, B. 592
- Carignani, G., see Mari, F. 527
- Cerutti, A., see Sartori, S. 2
- Ceylaner, S., see Inceci, F. 803
- Chae, J.-H., see Lee, J.S. 223
- Chae, J.-H., see Lee, J.S. 402
- Chen, C.-H., see Lee, H.-F. 599
- Chen, L., see Zhang, P. 747
- Chen, Y., see Yi, Z. 23
- Cheng, G., see Zhang, P. 747
- Cheon, J.-E., see Lee, J.S. 223

- Cheon, K.R., see Park, S.H. 625
 Chi, C.-S., see Lee, H.-F. 599
 Chida, S., see Kamei, A. 137
 Chiyonobu, T., see Shimada, S. 515
 Choi, J.E., see Lee, J.S. 402
 Choi, J.K., see Kim, J.-I. 359
 Chong, P.F., Ogata, R., Kobayashi, H., Koizumi, A., and Kira, R.
 Early onset of moyamoya syndrome in a Down syndrome patient with the genetic variant *RNF213* p.R4810K 822
 Chong, P.F., Haraguchi, K., Torio, M., Kirino, M., Ogata, R., Matsukura, M., Sakai, Y., Ishizaki, Y., Yamamoto, T., and Kira, R.
 A case of pontine tegmental cap dysplasia with comorbidity of oculoauriculovertebral spectrum 171
 Chugani, H.T., see Kamson, D.O. 370
 Chung, J.K., see Park, S.H. 625
 Cianci, P., see Mari, F. 527
 Cobanogullari Direk, M., see Bozlu, G. 901
 Cristofoli, F., see Mari, F. 527
 de Palma, L., see Sartori, S. 2
 De Simone, R., see Ranieri, A. 179
 Değerliyurt, A., Yalnizoglu, D., Bakar, E.E., Topcu, M., and Turanli, G.
 Electrical status epilepticus during sleep: A study of 22 patients 250
 Deguchi, K., see Yamamoto, T. 988
 Deiters, L., see Nögel, S.C. 394
 Delporte, L., see Rebour, R. 643
 Demirkol, E., see Kilinc, O. 352
 Densupsoontorn, N., see Vatanavicharn, N. 698
 Di Marco, C., see Mari, F. 527
 Dincer, A., see Ozcan, U.A. 495
 Ding, D., see Sun, H. 563
 Ding, Y., see Liu, Y. 286
 Ding, Y., see Wang, Q. 163
 Ding, Y., see Li, X. 952
 Doi, H., see Kobayashi, K. 572
 Dosa, L., see Mari, F. 527
 Dotti, M.T., see Mari, F. 527
 Egami, C., Yamashita, Y., Tada, Y., Anai, C., Mukasa, A., Yuge, K., Nagamitsu, S., and Matsuishi, T.
 Developmental trajectories for attention and working memory in healthy Japanese school-aged children 840
 Ekinci, G., see Kilinc, O. 352
 Endo, F., see Toda, Y. 230
 Endo, W., see Inui, T. 449
 Endzinienė, M., see Gelžinienė, G. 409
 Facini, C., see Orivoli, S. 833
 Facini, C., Pavlidis, E., Turco, E.C., and Pisani, F.
 Hereditary Hemorrhagic Telangiectasia presenting as migraine: A case report 974
 Fallerini, C., see Mari, F. 527
 Feng, H., see Xie, H. 797
 Fimiani, M., see Mari, F. 527
 Flores-Sarnat, L., see Sarnat, H.B. 553
 Frattini, D., see Fusco, C. 891
 Frattini, D., Nardocci, N., Pasarella, R., Panteghini, C., Garavaglia, B., and Fusco, C.
 Downbeat nystagmus as the presenting symptom of infantile neuroaxonal dystrophy: A case report 270
 Fu, X.J., see Kato, T. 911
 Fujihara, K., see Hino-Fukuyo, N. 849
 Fujii, T., see Ito, Y. 780
 Fujii, Y., see Niwa, T. 967
 Fujimaki, T., see Okumura, A. 978
 Fujita, K., see Nishiyama, M. 328
 Fukao, T., see Vatanavicharn, N. 698
 Fukasawa, T., Kubota, T., Negoro, T., Saitoh, M., Mizuguchi, M., Ihara, Y., Ishii, A., and Hirose, S.
 A case of recurrent encephalopathy with *SCN2A* missense mutation 631
 Fukuda, M., Kawabe, M., Takehara, M., Iwano, S., Kuwabara, K., Kikuchi, C., Wakamoto, H., Morimoto, T., Suzuki, Y., and Ishii, E.
 Carnitine deficiency: Risk factors and incidence in children with epilepsy 790
 Fukui, M., see Shimakawa, S. 733
 Fukui, M., see Nomura, S. 983
 Fukumoto, R., see Shibata, Y. 501
 Fukuyama, T., see Kubota, K. 158
 Fukuyama, T., see Motobayashi, M. 140
 Funato, M., see Morioka, I. 753
 Fung, C.W., see Wong, V.C.N. 729
 Furushima, W., Kaga, M., Nakamura, M., Gunji, A., and Inagaki, M.
 Auditory agnosia as a clinical symptom of childhood adrenoleukodystrophy 690
 Fusco, C., Frattini, D., Salerno, G.G., Canali, E., Bernasconi, P., and Maggi, L.
 New phenotype and neonatal onset of sodium channel myotonia in a child with a novel mutation of *SCN4A* gene 891
 Fusco, C., see Frattini, D. 270
 Fushiki, S., see Ogi, H. 739
 Gabata, T., see Ueda, F. 439
 Gandelman-Marton, R., see Heyman, E. 307
 Garavaglia, B., see Frattini, D. 270
 Garavelli, L., see Mari, F. 527
 Gelžinienė, G., Endzinienė, M., and Jurkevičienė, G.
 EEG activation by neuropsychological tasks in idiopathic generalized epilepsy of adolescence 409
 Gearing, G., see Kemmuir, C. 916
 Girschick, H.J., see Glaser, K. 347
 Glaser, K., Girschick, H.J., Schropp, C., and Speer, C.P.
 Psychomotor development following early treatment of severe infantile vitamin B12 deficiency and West syndrome – Is everything fine? A case report and review of literature 347
 Go, C.Y., see Mohammadi, M. 59
 Goto, M., Saito, Y., Honda, R., Saito, T., Sugai, K., Matsuda, Y., Miyatake, C., Takeshita, E., Ishiyama, A., Komaki, H., Nakagawa, E., Sasaki, M., Uto, C., Kikuchi, K., Motoki, T., and Saitoh, S.
 Episodic tremors representing cortical myoclonus are characteristic in Angelman syndrome due to *UBE3A* mutations 216
 Goto, T., see Okanishi, T. 423
 Gouider-Khouja, N., see Rebai, I. 153
 Gradišnik, P., Zagradisnik, B., Palfy, M., Kokalj-Vokac, N., and Marcun-Varda, N.
 Predictive value of paroxysmal EEG abnormalities for future epilepsy in focal febrile seizures 868
 Grossi, S., see Mari, F. 527
 Gulati, A., see Puppala, B. 206
 Guler, S., Kucukkoc, M., and Iscan, A.
 Prognosis and demographic characteristics of SSPE patients in Istanbul, Turkey 612
 Guler, S., see Kurtcan, S. 546
 Gunji, A., see Furushima, W. 690
 Hachiya, Y., see Shimada, S. 960
 Haginoya, K., see Inui, T. 449
 Haginoya, K., see Kobayashi, S. 858
 Haginoya, K., see Hino-Fukuyo, N. 849
 Haginoya, K., see Shimada, S. 515
 Haginoya, K., see Nakayama, T. 362
 Hallett, M.
 Tourette Syndrome: Update 651
 Hamano, S., see Nakamura, S. 925
 Han, B., Zou, H., Han, B., Zhu, W., Cao, Z., and Liu, Y.
 Diagnosis, treatment and follow-up of patients with tetrahydrobiopterin deficiency in Shandong province, China 592
 Han, B., see Han, B. 592

Download English Version:

<https://daneshyari.com/en/article/3036898>

Download Persian Version:

<https://daneshyari.com/article/3036898>

[Daneshyari.com](https://daneshyari.com)