

### **III. 研究成果の刊行に関する一覧表**

書籍

著者氏名	論文タイトル名	書籍全体の編集者名	書籍名	出版社名	出版地	出版年	ページ
水口雅, 他		小児急性脳症診療ガイドライン策定ワーキンググループ	小児急性脳症診療ガイドライン2016	診断と治療社	東京	2016	
前垣義弘, 他		小児けいれん重積治療ガイドライン策定ワーキンググループ	小児けいれん重積治療ガイドライン2017	診断と治療社	東京	2017	
山内秀雄, 他		Yamanouchi H, Moshé LS, Okumura A	Acute Encephalopathy and Encephalitis in Infancy and Its Related Disorders.1st Ed.	Elsevier	St. Louis	2017	

雑誌

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Yamamoto H, Okumura A, Natsume J, Kojima S, Mizuguchi M.	A severity score for acute necrotizing encephalopathy.	Brain and Development	37(3)	322-327	2015
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<u>Saitoh M</u> , Kobayashi K, Ohmori I, Tanaka Y, Tanaka K, Inoue T, Horino A, Ohmura K, Kumakura A, Takei Y, Hirabayashi S, Kajimoto M, Uchida T, Yamazaki S, Shiihara T, Kumagai T, Kasai M, Terashima H, <u>Kubota M</u> , <u>Mizuguchi M</u> .	Cytokine-related and sodium channel polymorphism as candidate predisposing factors for childhood encephalopathy FIRES/AE RRPS.	Journal of the Neurological Sciences	368	272-276	2016
Kurahashi H, Azuma Y, Masuda A, Okuno T, Nakahara E, Imamura T, <u>Saitoh M</u> , <u>Mizuguchi M</u> , Shimizu T, Ohno K, <u>Okumura A</u> .	<i>MYRF</i> is associated with encephalopathy with reversible myelin vacuolization.	Annals of Neurology	83(1)	98-106	2018
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Omata T, Fujii K, <u>Takanashi J</u> , Murayama K, Takayanagi M, Muta K, Kodama K, Iida Y, Watanabe Y, Shimomoto N.	Drugs indicated for mitochondrial dysfunction as treatments for acute encephalopathy.	Journal of the Neurological Sciences	360	57-60	2016
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Ishida S, Yasukawa K, Koizumi M, Abe K, Hirai N, Honda T, Sakuma S, Tada H, <u>Takanashi J</u> .	Excitotoxicity in encephalopathy associated with STEC O-157 infection.	Brain and Development	40(4)	357-360	2018
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<u>Sakuma H</u> , Tanuma N, Kuki I, Takahashi Y, Shiomi M, Hayashi M.	Intrathecal overproduction of proinflammatory cytokines and chemokines in febrile infection-related refractory status epilepticus.	Journal of Neurology, Neurosurgery and Psychiatry	86(7)	820-822	2015

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Negishi Y, Miya F, Hattori A, Johmura Y, Nakagawa M, Ando N, Hori I, Togawa T, Aoyama K, Ohashi K, Fukumura S, Mizuno S, Umemura A, Kishimoto Y, Okamoto N, Kato M, Tsunoda T, Yamasaki M, Kanemura Y, Kosaki K, Nakanishi M, <u>Saitoh S</u> .	A combination of genetic and biochemical analyses for the diagnosis of PI3K-AKT-mTOR pathway-associated megalencephaly.	BMC Medical Genetics	18	4	2017
Ishii A, Watkins JC, Chen D, <u>Hirose S</u> , Hammer MF.	Clinical implications of SCN1A missense and truncation variants in a large Japanese cohort with Dravet syndrome.	Epilepsia	58(2)	282-290	2017