

III. 研究成果の刊行に関する一覧表

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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
伊藤雅之	てんかんの病理	辻 貞俊	最新医学別冊. 新しい診断と治療のABC 74	最新医学社	大阪	2012	72-82

雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Arai A, Saito T, Hanai S, Otsuki T, Nakagawa E, Takahashi A, Kaneko Y, Kaido T, Saito Y, Sugai K, Sasaki M, Goto Y, Itoh M.	Abnormal maturation and differentiation of neocortical neurons in epileptogenic cortical malformation: unique distribution of layer-specific marker cells of focal cortical dysplasia and hemimegalencephaly.	Brain Res	1470	89-97	2012
Sakakibara T, Sukigara S, Otsuki T, Takahashi A, Kaneko Y, Kaido T, Yuko Saito Y, Sato N, Nakagawa E, Sugai K, Sasaki M, Goto Y, Itoh M.	Imbalance of interneuron distribution between neocortex and basal ganglia: Consideration of epileptogenesis of focal cortical dysplasia.	J Neurol Sci	323	128 - 133	2012
Sakakibara T, Saito T, Otsuki T, Takahashi A, Kaneko Y, Kaido T, Saito Y, Sato N, Nakagawa E, Sugai K, Sasaki M, Goto Y, Itoh M.	Delayed maturation of neurons of focal cortical dysplasia IIA and IIB: consideration from specific neocortical-layer marker expression.	J Neuropathol Exp Neurol	71	741 - 749	2012
Itoh M, Tahimic CGT, Ide S, Otsuki A, Saoka T, Noguchi S, Oshimura M, Goto Y, Kurimasa A.	Methyl CpG-binding protein isoform MeCP2_e2 is dispensable for phenotypes but essential for embryo viability and placenta development.	J Biol Chem	287	13859 - 13867	2012
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Kaido T, Otsuki T, Saito Y, Sugai K, Takahashi A, Kaneko Y, Sakakibara T, Saito Y, Takahashi H, Honda R, Nakagawa E, Sasaki M, Kakita A, Itoh M.	Novel pathological abnormalities of deep brain structures including dysplastic neurons in anterior striatum associated with focal cortical dysplasia in epilepsy.	J Neurosurg Pediatr	10	217 - 225	2012

Mizuuchi T, Kimura A, Tanaka A, Muto A, Nittono H, Seki Y, Takahashi T, Kurosawa T, Kage M, Takikawa H, Matsuishi T.	Characterization of urinary bile acids in a pediatric BRIC-1 patient: Effect of rifampicin treatment.	Clin Chim Acta	413	1301 - 1304	2012
Okabe Y, Takahashi T, Mitsumasu C, Koshi K, Tanaka E, Matsuishi T.	Alterations of Gene Expression and Glutamate Clearance in Astrocytes Derived from an MeCP2-null Mouse Model of Rett Syndrome.	PLoS ONE	7	E35354	2012
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Takahashi S, Matsumoto N, Okayama A, Suzuki N, Araki A, Okajima K, Tanaka H, Miyamoto A.	FOXP1 mutations in Japanese patients with the congenital variant of Rett syndrome.	Clin Genet	82	569 - 573	2012
Nagai M, Meguro-Horiike M, Horike S.	Epigenetic defects related to assisted reproductive technologies: Large offspring syndrome (LOS).	DNA Methylation-Genomic Technologies and Impact		167 - 182	2012