

## 研究成果の刊行に関する一覧表

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
Inaguma Y, Ito H, Hara A, Iwamoto I, Matsumoto A, Yamagata T, Tabata H, Nagata KI	Morphological characterization of mammalian Timeless in the mouse brain development.	Neurosci Res	92	21-28	2015
Mizuno M, Matsumoto A, Hamada N, Ito H, Miyauchi A, Jimbo EF, Momoi MY, Tabata H, Yamagata T, Nagata KI	Role of an adaptor protein Lin-7B in brain development: possible involvement in autism spectrum disorders.	J Neurochem	123	61-69	2014
Uehara N, Mori M, Tokuzawa Y, Mizuno Y, Tamaru S, Kohda M, Moriyama Y, Nakachi Y, Matoba N, Sakai T, Yamazaki T, Harashima H, Murayama K, Hattori K, Hayashi J, Yamagata T, Fujita Y, Ito M, Tanaka M, Nibu K, Ohtake A, Okazaki Y	New MT-ND6 and NDUFA1 mutations in mitochondrial respiratory chain disorders.	Ann Clin Transl Neurol	1	361-369	2014
Miyamoto Y, Ishikawa Y, Iegaki N, Sumi K, Fu K, Sato K, Furukawa-Hibi Y, Muramatsu S, Nabeshima T, Uno K, Nitta A	Overexpression of Shati/Nat8l, an <i>N</i> -acetyltransferase, in the nucleus accumbens attenuates the response to methamphetamine via activation of group II mGluRs in mice.	Int J Neuropsychopharmacol	17	1283-1294	2014
Ito H, Fujita K, Tagawa K, Chen X, Homma H, Sasabe T, Shimizu J, Shimizu S, Tamura T, Muramatsu S, Okazawa H	HMGB1 facilitates repair of mitochondrial DNA damage and extends the lifespan of mutant ataxin-1 knock-in mice.	EMBO Mol Med	7(1)	78-101	2014
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Shimbo H, Takagi M, Okuda M, Tsuyusaki Y, Takano K, Iai M, Yamashita S, Murayama K, Ohtake A, Goto Y, Aida N, Osaka H.	A rapid screening with direct sequencing from blood samples for the diagnosis of Leigh syndrome.	Mol Genet Metab Report	1	133-138	2014
Ohshiro-Sasaki A, Shimbo H, Takano K, Wada T, Osaka H.	A Three-Year-Old Boy With Glucose Transporter Type 1 Deficiency Syndrome Presenting With Episodic Ataxia.	Pediatr Neurol	50	99-100	2014
Miyatake S, Osaka H, Shiina M, Sasaki M, Takanashi J, Haginoya K, Wada T, Morimoto M, Ando N, Ikuta Y, Nakashima M, Tsurusaki Y, Miyake N, Ogata K, Matsumoto N, Saitsu H.	Expanding the phenotypic spectrum of TUBB4A-associated hypomyelinating leukoencephalopathies.	Neurology	82	2230-2237	2014
Nakamura K, Osaka H, Murakami Y, Anzai R, Nishiyama K, Kodera H, Nakashima M, Tsurusaki Y, Miyake N, Kinoshita T, Matsumoto N, Saitsu H.	PIGO mutations in intractable epilepsy and severe developmental delay with mild elevation of alkaline phosphatase levels.	Epilepsia	55	e13-17	2014