

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
Inoue K	Genetic risk factors for neurodegenerative diseases.	Wada K	Neurodegenerative Disorders as Systemic Diseases.	Springer Japan	Tokyo	2015	117-134
高梨潤一	小児神経疾患におけるMR spectroscopy (MRS) の臨床応用.	日本小児神経学会	続・イメージからせまる小児神経疾患	診断と治療社	東京	2015	19-22

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
Osaka H, Inoue K.	Pathophysiology and emerging therapeutic strategies in Pelizaeus-Merzbacher disease.	Expert Opinion on Orphan Drugs	3(12)	1447-1459	2015
Omata T, Nagai J, Shimbo H, Koizume S, Miyagi Y, Kurosawa K, Yamashita S, Osaka H, Inoue K.	A splicing mutation of proteolipid protein 1 in Pelizaeus-Merzbacher disease.	Brain Dev.	38(6)	581-584	2016
Ito Y, Inoue N, Inoue YU, Nakamura S, Matsuda Y, Inagaki M, Ohkubo T, Asami J, Terakawa YW, Kohsaka S, Goto Y, Akazawa C, Inoue T, Inoue K.	Additive dominant effect of a <i>SOX10</i> mutation underlies a complex phenotype of PCWH.	Neurobiol Dis.	80	1-14	2015
Sumida K, Inoue K, Takanashi J, Sasaki M, Watanabe K, Suzuki M, Kurahashi H, Omata T, Tanaka M, Yokochi K, Iio J, Iyoda K, Kurokawa T, Matsuo M, Sato T, Iwaki A, Osaka H, Kurosawa K, Yamamoto T, Matsumoto N, Maikusa N, Mastuda H, Sato N.	The magnetic resonance imaging spectrum of Pelizaeus-Merzbacher disease: A multicenter study of 19 patients.	Brain Dev.	38(6)	571-580	2016

Hoshino H, Kubota M	Clinical features and recent advances in research.	Pediatrics International	56	477-483	2014
久保田 雅也	Canavan病	日本臨床別冊神経症候群IV	神経症候群(第2版)	pp159-164	2014
Miyatake C, Koizumi S, Narazaki H, Asano T, Osaka H, Kurosawa K, Takanashi J, Fujino O.	Clinical pictures in Pelizaeus-Merzbacher disease: a report of a case.	J Nippon Med Sch.	82	74-75	2015
Kim Y, Asano Y, Koide R, Kimura H, Saitsu H, Matsumoto N, Bandoh M.	Callosal disconnection syndrome in symptomatic female carrier of Pelizaeus-Merzbacher disease.	J Neurol Sci.	358(1-2)	461-462	2015
Tsurusaki Y, Tanaka R, Shimada S, Shimojima K, Shiina M, Nakashima M, Saitsu H, Miyake N, Ogata K, Yamamoto T, Matsumoto N.	Novel compound heterozygous <i>L1AS</i> mutations cause glycine encephalopathy.	J Hum Genet.	60(10)	631-635	2015
Takanashi J	Neurochemistry of hypomyelination investigated with MR spectroscopy.	Magn Reson Med Sci.	14	85-91	2015
Yamamoto T, Takanashi J, Kurosawa K, Deguchi K, Osaka H, Inoue K	Comment on "Delayed myelination is not a constant feature of Allan-Herndon-Dudley syndrome: Report of a new case and review of the literature" by Azzolini S et al.	Brain Dev.	37	988-989	2015
Okanishi T, Yamamoto H, Hosokawa T, Ando N, Nagayama Y, Hashimoto Y, Maihara T, Goto T, Kutota M, Kawaguchi C, Yoshida H, Sugiura K, Itomi S, Ohno K, Takanashi J, Hayakawa M, Otsubo H, Okumura A.	Diffusion-weighted MRI for early diagnosis of neonatal herpes simplex encephalitis.	Brain Dev.	37	423-431	2015
Takeuchi A, Okamoto N, Fujinaga N, Morita H, Shimizu J, Akiyama T, Ninomiya S, Takanashi J, Kubo T.	Progressive brain atrophy in Schinzel-Giedion syndrome with a SETBP1 mutation.	Eur J Med Genet.	58	369-371	2015
Miyatake S, Tada H, Moriya S, Takanashi J, Hirano Y, Hayashi M, Oya Y, Nakashima M, Tsurusaki Y, Miyake N, Matsumoto N, Saitsu H.	Atypical giant axonal neuropathy arising from a homozygous mutation by uniparental isodisomy.	Clin Genet.	87	395-397	2015

Yamamoto T, Yoshioka S, Tsurusaki Y, Shino S, Shimojima K, Shigematsu Y, Takeuchi Y, Matsumoto N.	White matter abnormalities in an adult patient with L-2-hydroxyglutaric aciduria.	Brain Dev.	38	142-144	2016
Shimada S, Shimojima K, Sangu N, Hoshino A, Hachiya Y, Ohto T, Hashi Y, Nishida K, Mitani M, Kinjo S, Tsurusaki Y, Matsumoto N, Morimoto M, Yamamoto T.	Mutations in the genes encoding eukaryotic translation initiation factor 2B in Japanese patients with vanishing white matter disease.	Brain Dev.	37	960-966	2015
Masuda T, Ueda M, Ueyama H, Shimada S, Ishizaki M, Imamura S, Yamamoto T, Ando Y.	Megalencephalic leukoencephalopathy with subcortical cysts caused by compound heterozygous mutations in MLC1, in patients with and without subcortical cysts in the brain.	J Neurol Sci	351	211-213	2015
Yoshida T, Mizuta I, Saito K, Kimura Y, Park K, Ito Y, Haji S, Nakagawa M, Mizuno T.	Characteristic abnormal signals in medulla oblongata- 'eye spot' sign: four cases of elderly-onset Alexander disease.	Neurology clinical practice.	5	259-262	2015
Sugiyama A, Sawai S, Ito S, Mukai H, Beppu M, Yoshida T, Kuwabara S.	Incidental diagnosis of an asymptomatic adult-onset Alexander disease by brain magnetic resonance imaging for preoperative evaluation.	J Neurol Sci.	354	131-132	2015
Iwasaki Y, Saito Y, Mori K, Ito M, Mimuro M, Aiba I, Saito K, Mizuta I, Yoshida T, Nakagawa M, Yoshida M.	An autopsied case of adult-onset bulbospinal form Alexander disease with a novel S393R mutation in the GFAP gene.	Clin Neuropathol.	34	207-214	2015