

別紙4

研究成果の刊行に関する一覧表レイアウト
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
該当なし							

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
Uemura M, Nozaki H, Koyama A, Sakai N, Ando S, Kanazawa M, Onodera O	HTRA1 Mutations Identified in Symptomatic Carriers Have the Property of Interfering the Trimer-Dependent Activation Cascade	Frontiers in Neurology	10	1-6	2019
Sakai N, Uemura M, Kato T, Nozaki H, Koyama A, Ando S, Kamie H, Kato M and Onodera O	Hemorrhagic cerebral small vessel disease caused by a novel mutation in 3' UTR of collagen type IV alpha 1	Neurology Genetics	6	e383-4	2019
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富本秀和	遺伝性脳小血管病.	SRL宝函	40	21-27	2019
松山裕文、富本秀和	遺伝性血管性認知症	Medical Practice	36	603-607	2019
Senzaki S, Miura S, Ochi M, Kato T, Okada T, Matsumoto S, Shirakawa A, Ochi H, Igase M, Kitazawa R, Zhu B, Ikeuchi T, Ohyagi Y.	Sporadic Japanese case of adult-onset leukoencephalopathy with axonal spheroids and pigmented glia caused by a de novo p.Phe849del mutation in <i>CSF1R</i> .	Neurology Clinical Neuroscience	8	96-98	2020
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Funayama M, Sugihara M, Takata T, Mimura M, Ikeuchi T.	Remarkable behavioral signs and progressive non-fluent aphasia in a patient with adult-onset leukoencephalopathy with axonal spheroids and pigmented glia (ALSP)	Psychogeriatrics	19	282-285	2019
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