

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
Yamada M, Sakai K, Hamaguchi T, Noguchi- Shinohara M.	Cerebral amyloid angiopathy: emerging evidence for novel pathophysiology and pathogenesis.	Lee SH	Stroke Revisited: Pathophysiology of Stroke	Springer	Singapore	2020	81-94
濱口 毅, 山田正仁	プリオン病, 遅発 性ウイルス感染症	園生雅弘, 北川一夫, 青木正志	脳神経疾患最新 の治療2021- 2023	南江堂	東京	2021	131- 135
金谷泰宏	健康支援と社会保 障②公衆衛生 難病対策	平野かよ子	ナーシング・グ ラフィカ	メディカ出 版	大阪	2021	196- 203

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
Hamaguchi T, Sakai K, Kobayashi A, Kitamoto T, Ae R, Nakamura Y, Sanjo N, Arai K, Koide M, Katada F, Harada M, Murai H, Murayama S, Tsukamoto T, Mizusawa H, Yamada M.	Characterization of sporadic Creutzfeldt-Jakob disease and history of neurosurgery to identify potentially iatrogenic cases.	Emerg Infect Dis	26	1140-1146	2020
Hamaguchi T, Sanjo N, Ae R, Nakamura Y, Sakai K, Takao M, Murayama S, Iwasaki Y, Satoh K, Murai H, Harada M, Tsukamoto T, Mizusawa H, Yamada M.	MM2 type sporadic Creutzfeldt-Jakob disease: new diagnostic criteria for MM2-cortical type.	J Neurol Neurosurg Psychiatry	91	1158-1165	2020
Matsubayashi T, Akaza M, Hayashi Y, Hamaguchi T, Yamada M, Shimohata T, Yokota T, Sanjo N.	Focal sharp waves are a specific early-stage marker of the MM2-cortical form of sporadic Creutzfeldt-Jakob disease.	Prion	14	207-213	2020
Sakai K, Hamaguchi T, Sanjo N, Murai H, Iwasaki Y, Hamano T, Honma M, Noguchi-Shinohara M, Nozaki I, Nakamura Y, Kitamoto T, Harada M, Mizusawa H, Yamada M.	Diffusion-weighted magnetic resonance imaging in dura mater graft-associated Creutzfeldt-Jakob disease.	J Neurol Sci	418	117094	2020
Matsuzono K, Kim Y, Honda H, Anan Y, Tsunoda M, Amano Y, Fukusima N, Iwaki T, Kitamoto T, Fujimoto S.	Prion gene PRNP Y162X truncation mutation can induce a refractory esophageal achalasia.	Am J Gastroenterol			In press
Hayashi Y, Iwasaki Y, Waza M, Kato S, Akagi A, Kimura A, Inuzuka T, Satoh K, Kitamoto T, Yoshida M, Shimohata T.	Clinicopathological findings of a long-term survivor of V180I genetic Creutzfeldt-Jakob disease.	Prion	14	109-117	2020
Nomura T, Iwata I, Naganuma R, Matsushima M, Satoh K, Kitamoto T, Yabe I.	A patient with spastic paralysis finally diagnosed as V180I genetic Creutzfeldt-Jakob disease 9 years after onset.	Prion	14	226-231	2020
Takahashi-Iwata I, Yabe I, Kudo A, Eguchi K, Wakita M, Shirai S, Matsushima M, Toyoshima T, Chiba S, Tanikawa S, Tanaka S, Satoh K, Kitamoto T, Sasaki H.	MM2 cortical form of sporadic Creutzfeldt-Jakob disease without progressive dementia and akinetic mutism: A case deviating from current diagnostic criteria.	J Neurol Sci	412	116759	2020

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版
Kobayashi A, Hirata T, Nishikaze T, Ninomiya A, Maki Y, Takada Y, Kitamoto T, Kinoshita T.	α 2,3 linkage of sialic acid to a GPI anchor and an unpredicted GPI attachment site in human prion protein.	J Biol Chem	295	7789-7798	2020
Akagi A, Iwasaki Y, Yamamoto A, Matsuura H, Ikeda T, Mimuro M, Riku Y, Miyahara H, Kitamoto T, Yoshida M.	Identification of intracerebral hemorrhage in the early-phase of MM1+2C-type sporadic Creutzfeldt-Jakob disease: A case report.	Neuropathology	40	399-406	2020
Honda H, Matsuzono K, Satoh K, Fujisawa M, Suzuki SO, Furuyama C, Kitamoto T, Fujimoto S, Abe K, Iwaki T.	Detection of cutaneous prion protein deposits could help diagnose GPI-anchorless prion disease with neuropathy.	Eur J Neurol			In press
Iwasaki Y, Mori K, Ito M, Kawai Y, Akagi A, Riku Y, Miyahara H, Kobayashi A, Kitamoto T, Yoshida M.	System degeneration in an MM1-type sporadic Creutzfeldt-Jakob disease case with an unusually prolonged akinetic mutism state.	Prion			In press
Shintaku M, Nakamura T, Kaneda D, Shinde A, Kusaka H, Takeuchi A, Kitamoto T.	Genetic Creutzfeldt-Jakob disease-M232R with the cooccurrence of multiple prion strains, M1 + M2C + M2T: Report of an autopsy case.	Neuropathology			In press
Hermann P, Appleby B, Brandel JP, Caughey B, Collins S, Geschwind MD, Green A, Haik S, Kovacs GG, Ladogana A, Llorens F, Mead S, Nishida N, Pal S, Parchi P, Pocchiari M, Satoh K, Zanusso G, Zerr I.	Biomarkers and diagnostic guidelines for sporadic Creutzfeldt-Jakob disease.	Lancet Neurol	20	235-246	2021
Dong TT, Satoh K.	The Latest research on RT-QuIC assays-A literature review.	Pathogens			In press
Honda H, Mori S, Watanabe A, Sasagasako N, Sadashima S, Dong T, Satoh K, Nishida N, Iwaki T.	Abnormal prion protein deposits with high seeding activities in the skeletal muscle, femoral nerve, and scalp of an autopsied case of sporadic Creutzfeldt-Jakob disease.	Neuropathology			In press
Fujita H, Ogaki K, Shiina T, Onuma H, Sakuramoto H, Satoh K, Suzuki K.	V180I genetic Creutzfeldt-Jakob disease with cardiac sympathetic nerve denervation masquerading as Parkinson's disease: A case report.	Medicine (Baltimore)	100	e24294	2021

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版
Nakagaki T, Ishibashi D, Mori T, Miyazaki Y, Takatsuki H, Tange H, Taguchi Y, Satoh K, Atarashi R, Nishida N.	Administration of FK506 from late stage of disease prolongs survival of human prion-inoculated mice.	Neurotherapeutics	17	1850-1860	2020
Matsubara T, Satoh K, Homma T, Nakagaki T, Yamaguchi N, Atarashi R, Sudo Y, Uezono Y, Ishibashi D, Nishida N.	Prion protein interacts with the metabotropic glutamate receptor 1 and regulates the organization of Ca ²⁺ signaling.	Biochem Biophys Res Commun	525	447-454	2020
濱口 毅, 山田正仁	プリオン病	日本医事新報	5010	44	2020
濱口 毅, 山田正仁	クロイツフェルト・ヤコブ病	薬局 (増刊号: 病気とくすり 2021)	72	199-203	2021
中村治雅, 水澤英洋	患者レジストリシステム	医学のあゆみ	273	123-127	2020
佐藤克也	プリオン病	Pharma Medica	39	63-68	2021