別添５

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
田中　章景ら	脊髄小脳変性症	技術情報協会	疾患原因遺伝子・タンパク質の解析技術と創薬／診断技術への応用	技術情報協会	東京	2022	第6章第5節

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
Hirayanagi K, Ozaki H, Tsukagoshi S, Furuta N, Ikeda Y.	Porphyrins ameliorate spinocerebellar ataxia type 36 GGCCTG repeat expansion-mediated cytotoxicity.	Neurosci Res.	Oct;171	92-102	2021
Aoki S, Nagashima K, Shibata M, Kasahara H, Fujita Y, Hashiguchi A, Takashima H, Ikeda Y.	Sibling Cases of Charcot-Marie-Tooth Disease Type 4H with a Homozygous FGD4 Mutation and Cauda Equina Thickening	Intern Med.	60(24)	3975-3981	2021
H. Aoki, M. Higashi, M. Okita, N. Ando, S. Murayama, K. Ishikawa, T. Yokota.	Thymidine kinase 2 and mitochondrial protein COX I in the cerebellum of patients with spinocerebellar ataxia type 31 caused by penta-nucleotide repeats (TTCCA)n	The Cerebellum	Jan 27. doi: 10.1007/s12311-021-01364-2.		2022
Kurumada K, Sugiyama A, Hirano S, Yamamoto T, Yamanaka Y, Araki N, Yakiyama M, Yoshitake M,	Pareidolia in Parkinson's disease and multipe system atrophy	Parkinson's Disease	2021	2704755.	2021
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Kimura M et al.	Takotsubo Cardiomyopathy in Bickerstaff Brainstem Encephalitis Triggered by COVID-19	Front Neurol	12	822247	2021
Lipponen J, et al.	Molecular epidemiology of hereditary ataxia in Finland.	BMC Neurol	21(1)	382	2021
Kubota S et al.	SGTA associates with intracellular aggregates in neurodegenerative diseases	Mol Brain	14(1)	59	2021
Ebina J, Hara K, Watanabe H, Kawabata K, Yamashita F, Kawaguchi A, Yoshida Y, Kato T, Ogura A, Masuda M, Ohdake R, Mori D, Maesawa S, Katsuno M, Kano O, Sobue G.	Individual voxel-based morphometry adjusting covariates in multiple system atrophy. Parkinsonism Relat Disord.	Parkinsonism Relat Disord	90	114-119	2021
Yoshiyuki Kishimoto	Quantitative evaluation of upper limb ataxia in spinocerebellar ataxias	Annals of Clinical and Translational Neurology	9 巻 4 号	529-539	2022
Matsushima A, Maruyama Y, Mizukami N, Tetsuya M, Hashimoto M, Yoshida K.	Gait training with a wearable curara® robot for cerebellar ataxia: a single-arm study.	Biomed Eng Online	20(1)	90	2021
Nishida K, Sakashita K, Yamasaki H, Futamura N	Impact of tracheostomy invasive ventilation on survival in Japanese patients with multiple system atrophy	Parkinsonism Related Disorders	In press		2022

コメントの追加 [祥之2]: 4 月出版になっておりますので、令和 3 年度の業績報告にはあげない方がよろしいでしょうか？

コメントの追加 [祥之1]: 発表者名は勝野先生のお名前の方が適切でしたでしょうか？

Takiyama Y, et al.	A clinical and genetic study of SPG31 in Japan.	J Hum Genet	Online aheaed of print		2022
Takiyama Y, et al.	A p.Glu420Gln mutation in SPAST is associated with infantile onset spastic paraplegia complicated by cerebellar ataxia, epilepsy, peripheral neuropathy, and hypoplasia of the corpus callosum.	Neurol Sci.	43(3)	2123-2126	2022
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Takiyama Y, et al.	Spastic paraplegia with Paget's disease of bone due to a VCP gene mutation.	Intern Med.	60(1)	141-144	2021
Takiyama Y, et al.	A Nepalese family with an REEP2 mutaion: clinical and genetic study. Jul66(7):749-752.	J Hum Genet.	66(7)	749-752	2021
Takiyama Y, et al.	Biallelic variants in HPDL cause pure and complicated hereditary spastic paraplegia.	Brain	144(5)	1422-1434	2021
髙尾昌樹	脳表ヘモジデリン沈着症のオーバビュー	神経治療学	38巻2号	80-82	2021
大平雅之	脳表ヘモジデリン沈着症の疫学	神経治療学	38巻2号	83-85	2021
N Miyazawa et al.	Case of cortical superficial siderosis presenting with corticobasal syndrome	Clin Neurosci	10(2)	95-97	2022
藤本宏明, 宮井一郎	脊髄小脳変性症のニューロリハビリテーション治療	Jpn J Rehabil Med	58(5)	536-543	2021