運動失調症の医療基盤に関する調査研究班

研究成果（雑誌）の刊行に関する一覧

発表者氏名	論文タイトル名	発表誌名	巻	ページ	出版年 ※H26年度	GRANTへの謝辞の有無
Kokubo K, Suzuki K, Hattori N, Miyai I, Mori E	Executive dysfunction in patients with putaminal hemorrhage	J Stroke Cerebrovasc Dis	24	1978-85	2015	**無**
Dorsch AK, Thomas S, Xu X, Kaiser W, Dobkin BH, on behalf of the Si, and on behalf of the SIRRACT investigators (Miyai I, Kawano T).	SIRRACT: An International Randomized Clinical Trial of Activity Feedback During Inpatient Stroke Rehabilitation Enabled by Wireless Sensing.	Neurorehabil Neural Rep	29	407-15	2015	**無**
宮井一郎	**脳卒中リハビリテーションの新潮流.**	**脳**21	18	192-197	2015	**無**
三原雅史,宮井一郎	Functional NIRS	Clinical Neurosci	33	787-800	2015	**無**
宮井一郎	**小脳性運動失調症のリハビリテーション**	**医学のあゆみ**	255	1068-73	2015	**無**
Sugiyama A, Ito S, Suichi T, Sakurai T, Mukai H, Yokota H, Yonezu T, Kuwabara S.	Putaminal hypointensity on T2*-weighted MR imaging is the most practically useful sign in diagnosing multiple system atrophy: A preliminary study.	J Neurol Sci.	349(1-2)	174-178	2015	**無**
Yamamoto T, Asahina M, Yamanaka Y, Uchiyama T, Hirano S, Sugiyama A, Sakakibara R, Kuwabara S.	Urinary dysfunctions are more severe in the parkinsonian phenotype of multiple system atrophy	Movement Disorders Clinical Practice	**印刷中**		2016	**無**
桑原　聡	**小脳の最新知見 「皮質性小脳萎縮症」**	**医学のあゆみ**	255(10)	1052-54	2015	**無**
荒木信之，山中義崇，Anupama Poudel, **藤沼好克，片桐明，桑原聡，朝比奈正**	**脊髄小脳失調症6型の皮膚交感神経機能(原著論文)**	**発汗学**	22(1)	10-12	2015	**無**

人.						
桑原　聡	多系統萎縮症の生命予後予測因子	SCD・MSA(脊髄小脳変性症・多系統萎縮症)情報誌『Update on SCD』	9	2-3	2015	無
Koh K, Kobayashi F, Miwa M, Shindo K, Isozaki E, Ishiura H, Tsuji S, and Takiyama Y.	Novel mutations in the PNPLA6 gene in Boucher-Neuhauser syndrome.	J Hum Genet	60	217-220	2015	有
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三井　純	多系統萎縮症	医学のあゆみ	255	1047-1051	2015年	無
三井　純	多系統萎縮症の遺伝学	Annual Review神経内科2015		35-41	2015年	無
Mitsui, J., Matsukawa, T., Sasaki, H., Yabe, I., Matsushima, M., Durr, A., Brice, A., Takashima, H., Kikuchi, A., Aoki, M., Ishiura, H., Yasuda, T., Date, H., Ahsan, B., Iwata, A., Goto, J., Ichikawa, Y., Nakahara, Y., Momose, Y., Takahashi, Y., Hara, K., Kakita, A., Yamada, M., Takahashi, H., Onodera, O., Nishizawa, M., Watanabe, H., Ito, M., Sobue, G., Ishikawa, K., Mizusawa, H., Kanai, K., Hattori, T., Kuwabara, S., Arai, K., Koyano, S., Kuroiwa, Y., Hasegawa, K., Yuasa, T., Yasui, K., Nakashima, K., Ito, H., Izumi, Y., Kaji, R., Kato, T., Kusunoki, S., Osaki, Y., Horiuchi, M., Kondo, T., Murayama, S., Hattori, N., Yamamoto, M., Murata, M., Satake, W.,	Variants associated with Gaucher disease in multiple system atrophy	Annals of clinical and translational neurology	2	417-426	2015	無

Toda, T., Filla, A., Klockgether, T., Wullner, U., Nicholson, G., Gilman, S., Tanner, C.M., Kukull, W.A., Stern, M.B., Lee, V.M., Trojanowski, J.Q., Masliah, E., Low, P.A., Sandroni, P., Ozelius, L.J., Foroud, T. & Tsuji, S.						
Fujisawa, T., Yamaguchi, N., Kadowaki, H., Tsukamoto, Y., Tsuburaya, N., Tsubota, A., Takahashi, H., Naguro, I., Takahashi, Y., Goto, J., Tsuji, S., Nishitoh, H., Homma, K. & Ichijo, H.	systematic immunoprecipitation approach reinforces the concept of common conformational alterations in amyotrophic lateral sclerosis-linked SOD1 mutants.	Neurobiology of Diseases	82	478-486	2015	**無**
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Adachi T, Kitayama M, Nakano T, Adachi Y, Kato S, Nakashima K.	Autopsy case of spinocerebellar ataxia type 31 with severe dementia at the terminal stage.	Neuropathology.	35(3)	273-279	2015	**有**

Ozaki K, Doi H, Mitsui J, Sato N, Iikuni Y, Majima T, Yamane K, Irioka T, Ishiura H, Doi K, Morishita S, Higashi M, Sekiguchi T, Koyama K, Ueda N, Miura Y, Miyatake S, Matsumoto N, Yokota T, Tanaka F, Tsuji S, Mizusawa H, Ishikawa K.	A novel mutation in ELOVL4 leading to spinocerebellar ataxia (SCA) with the hot cross bun sign but lacking erythrokeratodermia: a broadened spectrum of SCA34.	JAMA Neurology	72	797-805	H27年	有
Pedroso JL, Abrahao A, Ishikawa K, Raskin S, de Souza PV, de Rezende Pinto WB, Braga-Neto P, de Albuquerque MV, Mizusawa H, Barsottini OG	When should we test patients with familial ataxias for SCA31? A misdiagnosed condition outside Japan?	J Neurol Sci.	355(1-2)	206-208	H27年	無
Aikawa T, Mogushi K, Iijima-Tsutsui K, Ishikawa K, Sakurai M, Tanaka H, Mizusawa H, Watase K.	Loss of MyD88 alters neuroinflammatory response and attenuates early Purkinje cell loss in a spinocerebellar ataxia type 6 mouse model.	Hum Mol Genet.	24(17)	4780-4791	H27年	無
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Mitsui J, Matsukawa T, Sasaki H, Yabe I, Matsushima M, Dürr A, Brice A, Takashima H, Kikuchi A, Aoki M, Ishiura H, Yasuda T, Date H, Ahsan B, Iwata A, Goto J, Ichikawa Y, Nakahara Y, Momose Y, Takahashi Y, Hara K, Kakita A, Yamada M, Takahashi H, Onodera O, Nishizawa M, Watanabe H, Ito M, Sobue G, Ishikawa K, Mizusawa H, Kanai K, Hattori T, Kuwabara S, Arai K, Koyano S, Kuroiwa Y, Hasegawa K, Yuasa T, Yasui K, Nakashima K, Ito H, Izumi Y, Kaji R, Kato T, Kusunoki S, Osaki Y, Horiuchi M, Kondo T, Murayama S, Hattori N, Yamamoto M, Murata M, Satake W, Toda T, Filla A, Klockgether T, Wüllner U, Nicholson G, Gilman S, Tanner CM, Kukull WA, Stern	Variants associated with Gaucher disease in multiple system atrophy.	Ann Clin Transl Neurol.	2(4)	417-426	H27年	無

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Mitsui J, Matsukawa T, Sasaki H, Yabe I, Matsushima M, Dürr A, Brice A, Takashima H, Kikuchi A, Aoki M, Ishiura H, Yasuda T, Date H, Ahsan B, Iwata A, Goto J, Ichikawa Y, Nakahara Y, Momose Y, Takahashi Y, Hara K, Kakita A, Yamada M, Takahashi H, Onodera O, Nishizawa M, Watanabe H, Ito M, Sobue G, Ishikawa K, Mizusawa H, Kanai K, Hattori T, Kuwabara S, Arai K, Koyano S, Kuroiwa Y, Hasegawa K, Yuasa T, Yasui K, Nakashima K, Ito H, Izumi Y, Kaji R, Kato T, Kusunoki S, Osaki Y, Horiuchi M, Kondo T, Murayama S, Hattori N, Yamamoto M, Murata M, Satake W, Toda T, Filla A, Klockgether T, Wüllner U, Nicholson G, Gilman S, Tanner CM, Kukull WA, Stern MB, Lee VM, Trojanowski JQ, Masliah E, Low PA, Sandroni P, Ozelius LJ, Foroud T, Tsuji S.	Variants associated with Gaucher disease in multiple system atrophy.	Ann Clin Transl Neurol.	2(4)	417-26	2015	**有**

Nakazato Y, Mochizuki H, Ishii N, Ohkubo R, Hirano R, Takashima H, Shiomi K, Nakazato M.	Spinocerebellar ataxia 36 accompanied by cervical dystonia.	J Neurol Sci.	357(1-2)	304-6	2015	**無**
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Koichihara R, Saito T, Ishiyama A, Komaki H, Yuasa S, Saito Y, Nakagawa E, Sugai K, Shiihara T, Shioya A, Saito Y, Higuchi Y, Hashiguchi A, Takashima H, Sasaki M.	A mild case of giant axonal neuropathy without central nervous system manifestation	Brain Dev.	Sep 14. pii:S0387-7604(15)	00180-1	2015	**有**
Tanji K, Odagiri S, Miki Y, Maruyama A, Nikaido Y, Mimura J, Mori F, Warabi E, Yanagawa T, Ueno S, Itoh K, Wakabayashi K	p62 deficiency enhances α-synuclein pathology in mice	Brain Pathol	25	552-564	2015	**有**
Nakamura K, Mori F, Tanji K, Miki Y, Yamada M, Kakita A, Takahashi H, Utsumi J, Sasaki H, Wakabayashi K	Isopentenyl diphosphate isomerase, a cholesterol synthesizing enzyme, is localized in Lewy bodies	Neuropathology	35	432-440	2015	**有**
Kon T, Miki Y, Tanji K, Mori F, Tomiyama M, Toyoshima Y, Kakita A, Takahashi H, Utsumi J, Sasaki H, Wakabayashi K	Localization of nuclear receptor subfamily 4, group A, member 3 (NR4A3) in Lewy body disease and multiple system atrophy	Neuropathology	35	503-509	2015	**有**
Nakamura K, Mori F, Kon T, Tanji K, Miki Y, Tomiyama M, Kurotaki H, Toyoshima Y, Kakita A, Takahashi H, Yamada M, Wakabayashi K	Filamentous aggregations of phosphorylated α-synuclein in Schwann cells (Schwann cell cytoplasmic inclusions) in multiple system atrophy	Acta Neuropathol Comm	3	29	2015	**有**

Tanji K, Miki Y, Maruyama A, Mimura J, Matsumiya T, Mori F, Imaizumi T, Itoh K, Wakabayashi K	Trehalose intake induces chaperone molecules along with autophagy in a mouse model of Lewy body disease	Biochem Biophys Res Com	465	746-752	2015	有
Mori F, Tanji K, Miki Y, Toyoshima Y, Yoshida M, Kakita A, Takahashi H, Utsumi J, Sasaki H, Wakabayashi K	G protein-coupled receptor 26 immunoreactivity in intranuclear inclusions associated with polyglutamine and intranuclear inclusion body diseases	Neuropathology			in press	有
Nakamura K, Mori F, Kon T, Tanji K, Miki Y, Tomiyama M, Kurotaki H, Toyoshima Y, Kakita A, Takahashi H, Yamada M, Wakabayashi K	Accumulation of phosphorylated α-synuclein in subpial and periventricular astrocytes in multiple system atrophy of long duration	Neuropathology			in press	有
Miki Y, Tanji K, Mori F, Utsumi J, Sasaki H, Kakita A, Takahashi H, Wakabayashi K	Alteration of upstream autophagy-related proteins (ULK1, ULK2, Beclin1, VPS34, and AMBRA1) in Lewy body disease	Brain Pathol			in press	有
Nakamura K, Mori F, Tanji K, Miki Y, Toyoshima Y, Kakita A, Takahashi H, Yamada M, Wakabayashi K	α-Synuclein pathology in the cranial and spinal nerves in Lewy body disease	Neuropathology			in press	有
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Sato Y, Nakatani E, Watanabe Y, Fukushima M, Nakashima K, Kannagi M, Kanatani Y, Mizushima H.	Prediction of prognosis of ALS: Importance of active denervation findings of the cervical-upper limb area and trunk area.	Intractable & Rare Diseases Research.	4	181-189	2015	有
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金谷泰宏	**わが国における難病とは.**	**日医雑誌**	144	1137-1139	2015	**無**
田中　洋康、武田　篤	**内の神経伝達物質と関連症状　パーキンソン病に於けるドパミン系・アセチルコリン系の低下とそれに伴う神経徴候**	**神経心理学**	31(2)	99-107	2015	**無**

谷口さやか、武田篤	【Parkinson病の治療　内科医に必要な新しい治験】Parkinson病の新しい理解　非運動症状を含めて	日本内科学会雑誌	104(8)	1546-1551	2015	無
大泉　英樹、武田　篤	【嗅覚障害臨床の最近の進歩】他科疾患と嗅覚障害　パーキンソン病と嗅覚障害	Progress in Medicine	35(4)	687-691	2015	無
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Tsuboi T, Watanabe H, Tanaka Y, Ohdake R, Yoneyama N, Hara K, Ito M, Hirayama M, Yamamoto M, Fujimoto Y, Kajita Y, Wakabayashi T, Sobue G	Characteristic laryngoscopic findings in Parkinson's disease patients after subthalamic nucleus deep brain stimulation and its correlation with voice disorder.	J Neural Transm (Vienna)	122(12)	1663-72.	2015	無
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原一洋, 渡辺宏久, 伊藤瑞規, 祖父江元	**拡散強調画像を用いた多系統萎縮症の上小脳脚病変の評価**	**神経内科**	82(6)	562-6	2015	**無**
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Yabe, I., Matsushima, M., Yoshida, K., Ishikawa, K., Shirai, S., Takahashi, I., Sasaki, H	Rare frequency of downbeat positioning nystagmus in spinocerebellar ataxia type 31	J Neurol Sci	350	90-92	2015	**有**
Sakushima, K., Nishimoto, N., Nojima, M., Matsushima, M., Yabe, I., Sato, N., Mori, M., Sasaki, H	Epidemiology of Multiple System Atrophy in Hokkaido, the Northernmost Island of Japan	Cerebellum	14	682-687	2015	**有**
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Shirai, S., Yabe, I., Matsushima, M., Ito, Y., M., Yoneyama, M., Sasaki, H	Quantitative evaluation of gait ataxia by accelerometers	J Neurol Sci	358	253-258	2015	**有**
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Mitsui Y, Kusunoki S, Arimura K, Kaji R, Kanda T, Kuwabara S, Sonoo M, Takada K; and the Japanese GBS Study Group.	A multicentre prospective study of Guillain-Barre Syndrome in Japan: a focus on the incidence of subtypes.	J Neurol Neurosurg Psychiatry.	86	110-4	2015	
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