

## 研究成果の刊行に関する一覧表

発表者氏名：論文タイトル名．発表誌名 巻号：ページ，出版年
<p>Uruha A, Hayashi YK, Oya Y, Mori-Yoshimura M, Kanai M, Murata M, Kawamura M, Ogata K, Matsumura T, Suzuki S, Takahashi Y, Kondo T, Kawarabayashi T, Ishii Y, Kokubun N, Yokoi S, Yasuda R, Kira JI, Mitsuhashi S, <u>Noguchi S</u>, Nonaka I, <u>Nishino I</u>: Necklace cytoplasmic bodies in hereditary myopathy with early respiratory failure. <b><i>J Neurol Neurosurg Psychiatry</i></b>. [Epub Sep 2014] ahead of print</p>
<p>Endo Y, Furuta A, <u>Nishino I</u>: Danon disease: a phenotypic expression of LAMP-2 deficiency. <b><i>Acta Neuropathol</i></b>. 129(3): 391-398, Mar, 2015 [Epub Jan 2015]</p>
<p>Tanboon J, Hayashi YK, <u>Nishino I</u>, Sangruchi T: Kyphoscoliosis and easy fatigability in a 14-year-old boy. <b><i>Neuropathology</i></b>. 35(1): 91-93, Feb 2015 [Epub Aug 2014]</p>
<p>Dong M, <u>Noguchi S</u>, Endo Y, Hayashi YK, Yoshida S, Nonaka I, <u>Nishino I</u>: <i>DAG1</i> mutations associated with asymptomatic hyperCKemia and hypoglycosylation of <math>\alpha</math>-dystroglycan. <b><i>Neurology</i></b>. 84(3): 273-279, Jan, 2015 [Epub Dec 2014]</p>
<p>Yonekawa T, Malicdan MC, Cho A, Hayashi YK, Nonaka I, Mine T, Yamamoto T, <u>Nishino I</u>, <u>Noguchi S</u>: Sialyllactose ameliorates myopathic phenotypes in symptomatic GNE myopathy model mice. <b><i>Brain</i></b>. 137(10): 2670-2679, Oct, 2014 [Epub Jul 2014]</p>
<p>Anada RP, Wong KT, Malicdan MC, Goh KJ, Hayashi YK, <u>Nishino I</u>, <u>Noguchi S</u>: Absence of beta-amyloid deposition in the central nervous system of a transgenic mouse model of distal myopathy with rimmed vacuoles. <b><i>Amyloid</i></b>. 21(2): 138-139, Jun, 2014 [Epub Mar 2014]</p>

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