## Anna Nogalska

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Sporadic inclusion-body myositis: A degenerative muscle disease associated with aging, impaired muscle protein homeostasis and abnormal mitophagy. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2015, 1852, 633-643.	3.8	81
2	Sodium phenylbutyrate reverses lysosomal dysfunction and decreases amyloid-l²42 in an in vitro-model of inclusion-body myositis. Neurobiology of Disease, 2014, 65, 93-101.	4.4	10
3	Pathogenic Considerations in Sporadic Inclusion-Body Myositis, a Degenerative Muscle Disease Associated With Aging and Abnormalities of Myoproteostasis. Journal of Neuropathology and Experimental Neurology, 2012, 71, 680-693.	1.7	63
4	Guidelines for the use and interpretation of assays for monitoring autophagy. Autophagy, 2012, 8, 445-544.	9.1	3,122
5	Activation of the γ-secretase complex and presence of γ-secretase-activating protein may contribute to Aβ42 production in sporadic inclusion-body myositis muscle fibers. Neurobiology of Disease, 2012, 48, 141-149.	4.4	11
6	Novel demonstration of conformationally modified tau in sporadic inclusion-body myositis muscle fibers. Neuroscience Letters, 2011, 503, 229-233.	2.1	12
7	Abnormalities of NBR1, a novel autophagy-associated protein, in muscle fibers of sporadic inclusion-body myositis. Acta Neuropathologica, 2011, 122, 627-636.	7.7	49
8	Novel demonstration of amyloid-β oligomers in sporadic inclusion-body myositis muscle fibers. Acta Neuropathologica, 2010, 120, 661-666.	7.7	40
9	In AβPPâ€overexpressing cultured human muscle fibers proteasome inhibition enhances phosphorylation of AβPP751 and GSK3β activation: effects mitigated by lithium and apparently relevant to sporadic inclusionâ€body myositis. Journal of Neurochemistry, 2010, 112, 389-396.	3.9	35
10	Increased BACE1 mRNA and noncoding BACE1-antisense transcript in sporadic inclusion-body myositis muscle fibers—Possibly caused by endoplasmic reticulum stress. Neuroscience Letters, 2010, 474, 140-143.	2.1	28
11	Impaired Autophagy in Sporadic Inclusion-Body Myositis and in Endoplasmic Reticulum Stress-Provoked Cultured Human Muscle Fibers. American Journal of Pathology, 2010, 177, 1377-1387.	3.8	94
12	Amyloid-l²42 is preferentially accumulated in muscle fibers of patients with sporadic inclusion-body myositis. Acta Neuropathologica, 2009, 117, 569-574.	7.7	56
13	p62/SQSTM1 is overexpressed and prominently accumulated in inclusions of sporadic inclusion-body myositis muscle fibers, and can help differentiating it from polymyositis and dermatomyositis. Acta Neuropathologica, 2009, 118, 407-413.	7.7	133
14	In inclusion-body myositis muscle fibers Parkinson-associated DJ-1 is increased and oxidized. Free Radical Biology and Medicine, 2008, 45, 773-779.	2.9	24
15	Endoplasmic reticulum stress induces myostatin precursor protein and NF-κB in cultured human muscle fibers: Relevance to inclusion body myositis. Experimental Neurology, 2007, 204, 610-618.	4.1	50
16	Homocysteine-induced endoplasmic reticulum protein (Herp) is up-regulated in sporadic inclusion-body myositis and in endoplasmic reticulum stress-induced cultured human muscle fibers. Journal of Neurochemistry, 2006, 96, 1491-1499.	3.9	60