

# Richard Webster

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/10225027/publications.pdf>

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13  
papers

338  
citations

933447

10  
h-index

1125743

13  
g-index

13  
all docs

13  
docs citations

13  
times ranked

386  
citing authors

#	ARTICLE	IF	CITATIONS
1	Passive and active immunization models of MuSK-Ab positive myasthenia: Electrophysiological evidence for pre and postsynaptic defects. <i>Experimental Neurology</i> , 2012, 234, 506-512.	4.1	112
2	Serological and experimental studies in different forms of myasthenia gravis. <i>Annals of the New York Academy of Sciences</i> , 2018, 1413, 143-153.	3.8	44
3	CHRND mutation causes a congenital myasthenic syndrome by impairing co-clustering of the acetylcholine receptor with rapsyn. <i>Brain</i> , 2006, 129, 2784-2793.	7.6	34
4	A mouse model of AChR deficiency syndrome with a phenotype reflecting the human condition. <i>Human Molecular Genetics</i> , 2004, 13, 2947-2957.	2.9	29
5	$\beta$ -2-Adrenergic receptor agonists ameliorate the adverse effect of long-term pyridostigmine on neuromuscular junction structure. <i>Brain</i> , 2019, 142, 3713-3727.	7.6	24
6	Fast-channel congenital myasthenic syndrome with a novel acetylcholine receptor mutation at the $\epsilon$ - $\mu$ subunit interface. <i>Neuromuscular Disorders</i> , 2014, 24, 143-147.	0.6	19
7	A novel congenital myasthenic syndrome due to decreased acetylcholine receptor ion-channel conductance. <i>Brain</i> , 2012, 135, 1070-1080.	7.6	18
8	Animal Models of the Neuromuscular Junction, Vitally Informative for Understanding Function and the Molecular Mechanisms of Congenital Myasthenic Syndromes. <i>International Journal of Molecular Sciences</i> , 2018, 19, 1326.	4.1	17
9	Rapsyn facilitates recovery from desensitization in fetal and adult acetylcholine receptors expressed in a muscle cell line. <i>Journal of Physiology</i> , 2019, 597, 3713-3725.	2.9	13
10	Interaction of Axonal Chondrolectin with Collagen XIXa1 Is Necessary for Precise Neuromuscular Junction Formation. <i>Cell Reports</i> , 2019, 29, 1082-1098.e10.	6.4	13
11	Congenital myasthenic syndrome due to a TOR1AIP1 mutation: a new disease pathway for impaired synaptic transmission. <i>Brain Communications</i> , 2020, 2, fcaa174.	3.3	11
12	Antagonistic postsynaptic and presynaptic actions of cyclohexanol on neuromuscular synaptic transmission and function. <i>Journal of Physiology</i> , 2021, 599, 5417-5449.	2.9	3
13	Slow-Channel Congenital Myasthenic Syndrome due to a Novel Mutation in the Acetylcholine Receptor Alpha Subunit in a South Asian: A Case Report. <i>Journal of Neuromuscular Diseases</i> , 2021, 8, 163-167.	2.6	1