

# Peter J Brophy

## List of Publications by Year in descending order

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34  
papers

2,892  
citations

201674

27  
h-index

377865

34  
g-index

36  
all docs

36  
docs citations

36  
times ranked

2529  
citing authors

#	ARTICLE	IF	CITATIONS
1	Neurofascins Are Required to Establish Axonal Domains for Saltatory Conduction. <i>Neuron</i> , 2005, 48, 737-742.	8.1	306
2	An Oligodendrocyte Cell Adhesion Molecule at the Site of Assembly of the Paranodal Axo-Glial Junction. <i>Journal of Cell Biology</i> , 2000, 150, 657-666.	5.2	280
3	Peripheral Demyelination and Neuropathic Pain Behavior in Periaxin-Deficient Mice. <i>Neuron</i> , 2000, 26, 523-531.	8.1	194
4	Specific Disruption of a Schwann Cell Dystrophin-Related Protein Complex in a Demyelinating Neuropathy. <i>Neuron</i> , 2001, 30, 677-687.	8.1	189
5	Restricted growth of Schwann cells lacking Cajal bands slows conduction in myelinated nerves. <i>Nature</i> , 2004, 431, 191-195.	27.8	187
6	A Glial Signal Consisting of Gliomedin and NrCAM Clusters Axonal Na <sup>+</sup> Channels during the Formation of Nodes of Ranvier. <i>Neuron</i> , 2010, 65, 490-502.	8.1	179
7	Glial and neuronal isoforms of Neurofascin have distinct roles in the assembly of nodes of Ranvier in the central nervous system. <i>Journal of Cell Biology</i> , 2008, 181, 1169-1177.	5.2	171
8	Periaxin, a novel protein of myelinating schwann cells with a possible role in axonal ensheathment. <i>Neuron</i> , 1994, 12, 497-508.	8.1	157
9	A Critical Role for Neurofascin in Regulating Action Potential Initiation through Maintenance of the Axon Initial Segment. <i>Neuron</i> , 2011, 69, 945-956.	8.1	139
10	Periaxin mutations cause a broad spectrum of demyelinating neuropathies. <i>Annals of Neurology</i> , 2002, 51, 709-715.	5.3	106
11	Two PDZ Domain Proteins Encoded by the Murine Periaxin Gene Are the Result of Alternative Intron Retention and Are Differentially Targeted in Schwann Cells. <i>Journal of Biological Chemistry</i> , 1998, 273, 5794-5800.	3.4	79
12	Increasing Internodal Distance in Myelinated Nerves Accelerates Nerve Conduction to a Flat Maximum. <i>Current Biology</i> , 2012, 22, 1957-1961.	3.9	79
13	Clinicopathological and genetic study of early-onset demyelinating neuropathy. <i>Brain</i> , 2004, 127, 2540-2550.	7.6	76
14	Acceleration of conduction velocity linked to clustering of nodal components precedes myelination. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, E321-8.	7.1	65
15	Proteome profile of peripheral myelin in healthy mice and in a neuropathy model. <i>ELife</i> , 2020, 9, .	6.0	63
16	A Tripartite Nuclear Localization Signal in the PDZ-domain Protein L-periaxin. <i>Journal of Biological Chemistry</i> , 2000, 275, 4537-4540.	3.4	58
17	The paranodal cytoskeleton clusters Na <sup>+</sup> channels at nodes of Ranvier. <i>ELife</i> , 2017, 6, .	6.0	57
18	Drp2 and Periaxin Form Cajal Bands with Dystroglycan But Have Distinct Roles in Schwann Cell Growth. <i>Journal of Neuroscience</i> , 2012, 32, 9419-9428.	3.6	53

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19	Differential Stability of PNS and CNS Nodal Complexes When Neuronal Neurofascin Is Lost. <i>Journal of Neuroscience</i> , 2014, 34, 5083-5088.	3.6	49
20	Schwann Cell O-GlcNAc Glycosylation Is Required for Myelin Maintenance and Axon Integrity. <i>Journal of Neuroscience</i> , 2016, 36, 9633-9646.	3.6	48
21	Periaxin is required for hexagonal geometry and membrane organization of mature lens fibers. <i>Developmental Biology</i> , 2011, 357, 179-190.	2.0	47
22	Absence of Dystrophin Related Protein-2 disrupts Cajal bands in a patient with Charcot-Marie-Tooth disease. <i>Neuromuscular Disorders</i> , 2015, 25, 786-793.	0.6	40
23	Loss of Glial Neurofascin155 Delays Developmental Synapse Elimination at the Neuromuscular Junction. <i>Journal of Neuroscience</i> , 2014, 34, 12904-12918.	3.6	39
24	Neurofascin 140 Is an Embryonic Neuronal Neurofascin Isoform That Promotes the Assembly of the Node of Ranvier. <i>Journal of Neuroscience</i> , 2015, 35, 2246-2254.	3.6	37
25	Input-Output Relationship of CA1 Pyramidal Neurons Reveals Intact Homeostatic Mechanisms in a Mouse Model of Fragile X Syndrome. <i>Cell Reports</i> , 2020, 32, 107988.	6.4	37
26	FAK Is Required for Schwann Cell Spreading on Immature Basal Lamina to Coordinate the Radial Sorting of Peripheral Axons with Myelination. <i>Journal of Neuroscience</i> , 2014, 34, 13422-13434.	3.6	36
27	Homozygous mutation in the Neurofascin gene affecting the glial isoform of Neurofascin causes severe neurodevelopment disorder with hypotonia, amimia and areflexia. <i>Human Molecular Genetics</i> , 2018, 27, 3669-3674.	2.9	34
28	Assembly of CNS Nodes of Ranvier in Myelinated Nerves Is Promoted by the Axon Cytoskeleton. <i>Current Biology</i> , 2017, 27, 1068-1073.	3.9	32
29	A murine model of Charcot-Marie-Tooth disease 4F reveals a role for the C-terminus of periaxin in the formation and stabilization of Cajal bands. <i>Wellcome Open Research</i> , 2018, 3, 20.	1.8	12
30	Direct Binding of the Flexible C-Terminal Segment of Periaxin to $\alpha 24$ Integrin Suggests a Molecular Basis for CMT4F. <i>Frontiers in Molecular Neuroscience</i> , 2019, 12, 84.	2.9	12
31	Neurofascin and Kv7.3 are delivered to somatic and axon terminal surface membranes en route to the axon initial segment. <i>ELife</i> , 2020, 9, .	6.0	12
32	Completion of neuronal remodeling prompts myelination along developing motor axon branches. <i>Journal of Cell Biology</i> , 2021, 220, .	5.2	7
33	Loss of protohaem IX farnesyltransferase in mature dentate granule cells impairs short-term facilitation at mossy fibre to CA3 pyramidal cell synapses. <i>Journal of Physiology</i> , 2017, 595, 2147-2160.	2.9	6
34	Dynamic early clusters of nodal proteins contribute to node of Ranvier assembly during myelination of peripheral neurons. <i>ELife</i> , 2021, 10, .	6.0	6