

# Esther Becker

## List of Publications by Year in descending order

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46  
papers

4,210  
citations

186265

28  
h-index

243625

44  
g-index

47  
all docs

47  
docs citations

47  
times ranked

6635  
citing authors

#	ARTICLE	IF	CITATIONS
1	Moonwalker Mouse. , 2022, , 1773-1788.		0
2	Cerebellar Modelling Using Human Induced Pluripotent Stem Cells. <i>Neuromethods</i> , 2022, , 1-21.	0.3	2
3	High-resolution transcriptional landscape of xeno-free human induced pluripotent stem cell-derived cerebellar organoids. <i>Scientific Reports</i> , 2021, 11, 12959.	3.3	32
4	Caspr2 interacts with type 1 inositol 1,4,5-trisphosphate receptor in the developing cerebellum and regulates Purkinje cell morphology. <i>Journal of Biological Chemistry</i> , 2020, 295, 12716-12726.	3.4	3
5	Deconstructing cerebellar development cell by cell. <i>PLoS Genetics</i> , 2020, 16, e1008630.	3.5	32
6	Moonwalker Mouse. , 2020, , 1-16.		1
7	A gene expression signature in developing Purkinje cells predicts autism and intellectual disability co-morbidity status. <i>Scientific Reports</i> , 2019, 9, 485.	3.3	14
8	TRPC3 is a major contributor to functional heterogeneity of cerebellar Purkinje cells. <i>ELife</i> , 2019, 8, .	6.0	45
9	Genotypeâ€phenotype correlations, dystonia and disease progression in spinocerebellar ataxia type 14. <i>Movement Disorders</i> , 2018, 33, 1119-1129.	3.9	26
10	A Simplified Method for Generating Purkinje Cells from Human-Induced Pluripotent Stem Cells. <i>Cerebellum</i> , 2018, 17, 419-427.	2.5	48
11	Neurodegeneration in SCA14 is associated with increased PKC $\delta$ kinase activity, mislocalization and aggregation. <i>Acta Neuropathologica Communications</i> , 2018, 6, 99.	5.2	37
12	The Use of Stem Cell-Derived Neurons for Understanding Development and Disease of the Cerebellum. <i>Frontiers in Neuroscience</i> , 2018, 12, 646.	2.8	5
13	Cerebellar involvement in autism and ADHD. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2018, 155, 61-72.	1.8	56
14	From Mice to Men: TRPC3 in Cerebellar Ataxia. <i>Cerebellum</i> , 2017, 16, 877-879.	2.5	10
15	Functional expression of calciumâ€permeable canonical transient receptor potential 4â€containing channels promotes migration of medulloblastoma cells. <i>Journal of Physiology</i> , 2017, 595, 5525-5544.	2.9	30
16	Dominant Mutations in GRM1 Cause Spinocerebellar Ataxia Type 44. <i>American Journal of Human Genetics</i> , 2017, 101, 451-458.	6.2	62
17	Recent advances in modelling of cerebellar ataxia using induced pluripotent stem cells. <i>Journal of Neurology and Neuromedicine</i> , 2017, 2, 11-15.	0.9	11
18	A Transient Translaminar GABAergic Interneuron Circuit Connects Thalamocortical Recipient Layers in Neonatal Somatosensory Cortex. <i>Neuron</i> , 2016, 89, 536-549.	8.1	124

#	ARTICLE	IF	CITATIONS
19	Consensus Paper: Cerebellar Development. <i>Cerebellum</i> , 2016, 15, 789-828.	2.5	337
20	Do mutations in the murine ataxia gene <i>TRPC3</i> cause cerebellar ataxia in humans?. <i>Movement Disorders</i> , 2015, 30, 284-286.	3.9	78
21	The mutant Moonwalker <i>TRPC3</i> channel links calcium signaling to lipid metabolism in the developing cerebellum. <i>Human Molecular Genetics</i> , 2015, 24, 4114-4125.	2.9	24
22	Modeling Suggests <i>TRPC3</i> Hydrogen Bonding and Not Phosphorylation Contributes to the Ataxia Phenotype of the Moonwalker Mouse. <i>Biochemistry</i> , 2015, 54, 4033-4041.	2.5	10
23	Induced pluripotent stem cell technology for modelling and therapy of cerebellar ataxia. <i>Open Biology</i> , 2015, 5, 150056.	3.6	38
24	Reciprocal regulation of two G protein-coupled receptors sensing extracellular concentrations of Ca <sup>2+</sup> and H <sup>+</sup> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 10738-10743.	7.1	27
25	The Moonwalker Mouse: New Insights into <i>TRPC3</i> Function, Cerebellar Development, and Ataxia. <i>Cerebellum</i> , 2014, 13, 628-636.	2.5	41
26	Next generation sequencing for molecular diagnosis of neurological disorders using ataxias as a model. <i>Brain</i> , 2013, 136, 3106-3118.	7.6	146
27	Early Onset of Ataxia in Moonwalker Mice Is Accompanied by Complete Ablation of Type II Unipolar Brush Cells and Purkinje Cell Dysfunction. <i>Journal of Neuroscience</i> , 2013, 33, 19689-19694.	3.6	41
28	Autism Spectrum Disorder and the Cerebellum. <i>International Review of Neurobiology</i> , 2013, 113, 1-34.	2.0	197
29	Contactin-associated protein-2 antibodies in non-paraneoplastic cerebellar ataxia. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, 437-440.	1.9	105
30	Candidate Screening of the <i>TRPC3</i> Gene in Cerebellar Ataxia. <i>Cerebellum</i> , 2011, 10, 296-299.	2.5	27
31	<i>Oxr1</i> Is Essential for Protection against Oxidative Stress-Induced Neurodegeneration. <i>PLoS Genetics</i> , 2011, 7, e1002338.	3.5	130
32	A JIP3-Regulated GSK3 <sup>β</sup> /DCX Signaling Pathway Restricts Axon Branching. <i>Journal of Neuroscience</i> , 2010, 30, 16766-16776.	3.6	51
33	A point mutation in <i>TRPC3</i> causes abnormal Purkinje cell development and cerebellar ataxia in moonwalker mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 6706-6711.	7.1	187
34	Activation of FOXO1 by Cdk1 in Cycling Cells and Postmitotic Neurons. <i>Science</i> , 2008, 319, 1665-1668.	12.6	167
35	Pin1 in Neuronal Apoptosis. <i>Cell Cycle</i> , 2007, 6, 1332-1335.	2.6	23
36	A Conserved MST-FOXO Signaling Pathway Mediates Oxidative-Stress Responses and Extends Life Span. <i>Cell</i> , 2006, 125, 987-1001.	28.9	758

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37	Pin1 Mediates Neural-Specific Activation of the Mitochondrial Apoptotic Machinery. <i>Neuron</i> , 2006, 49, 655-662.	8.1	73
38	p38 MAP Kinase Mediates Apoptosis through Phosphorylation of BimEL at Ser-65. <i>Journal of Biological Chemistry</i> , 2006, 281, 25215-25222.	3.4	195
39	Bim Regulation of Lumen Formation in Cultured Mammary Epithelial Acini Is Targeted by Oncogenes. <i>Molecular and Cellular Biology</i> , 2005, 25, 4591-4601.	2.3	130
40	Beyond proliferation—cell cycle control of neuronal survival and differentiation in the developing mammalian brain. <i>Seminars in Cell and Developmental Biology</i> , 2005, 16, 439-448.	5.0	33
41	Characterization of the c-Jun N-Terminal Kinase-BimEL Signaling Pathway in Neuronal Apoptosis. <i>Journal of Neuroscience</i> , 2004, 24, 8762-8770.	3.6	108
42	Cell cycle regulation of neuronal apoptosis in development and disease. <i>Progress in Neurobiology</i> , 2004, 72, 1-25.	5.7	274
43	Apoptosis Induced by p75NTR Overexpression Requires Jun Kinase-Dependent Phosphorylation of Bad. <i>Journal of Neuroscience</i> , 2003, 23, 11373-11381.	3.6	156
44	JNK Phosphorylation and Activation of BAD Couples the Stress-activated Signaling Pathway to the Cell Death Machinery. <i>Journal of Biological Chemistry</i> , 2002, 277, 40944-40949.	3.4	212
45	Specific role for cathepsin S in the generation of antigenic peptides <i>in vivo</i> . <i>European Journal of Immunology</i> , 2002, 32, 467-476.	2.9	98
46	Carbon source-dependent transcriptional regulation of the QCR8 gene in <i>Kluyveromyces lactis</i> .. <i>Current Genetics</i> , 2001, 39, 311-318.	1.7	6