Kevin C Eggan

List of Publications by Year in Descending Order

106 papers	9,438 citations	40 h-index	97 g-index
122 ext. papers	11,313 ext. citations	16 avg, IF	5.78 L-index

#	Paper	IF	Citations
106	Considerations and practical implications of performing a phenotypic CRISPR/Cas survival screen.. *PLoS ONE*, **2022**, 17, e0263262	3.7	1
105	Loss of mouse Stmn2 function causes motor neuropathy.. *Neuron*, **2022**,	13.9	2
104	Whole-genome analysis of human embryonic stem cells enables rational line selection based on genetic variation.. *Cell Stem Cell*, **2022**,	18	1
103	Cx43 hemichannels contribute to astrocyte-mediated toxicity in sporadic and familial ALS.. *Proceedings of the National Academy of Sciences of the United States of America*, **2022**, 119, e2107391119	11.5	0
102	Spinal motor neuron transplantation to enhance nerve reconstruction strategies: Towards a cell therapy.. *Experimental Neurology*, **2022**, 353, 114054	5.7	0
101	The genetic architecture of DNA replication timing in human pluripotent stem cells. *Nature Communications*, **2021**, 12, 6746	17.4	4
100	Genoppi is an open-source software for robust and standardized integration of proteomic and genetic data. *Nature Communications*, **2021**, 12, 2580	17.4	3
99	Human amyotrophic lateral sclerosis excitability phenotype screen: Target discovery and validation. *Cell Reports*, **2021**, 35, 109224	10.6	11
98	Connecting TDP-43 Pathology with Neuropathy. *Trends in Neurosciences*, **2021**, 44, 424-440	13.3	10
97	Cancer-Related Mutations Identified in Primed Human Pluripotent Stem Cells. *Cell Stem Cell*, **2021**, 28, 10-11	18	7
96	De novo DNA methyltransferases DNMT3A and DNMT3B are essential for XIST silencing for erosion of dosage compensation in pluripotent stem cells. *Stem Cell Reports*, **2021**, 16, 2138-2148	8	1
95	C9orf72 suppresses systemic and neural inflammation induced by gut bacteria. *Nature*, **2020**, 582, 89-94	50.4	83
94	Absence of Survival and Motor Deficits in 500 Repeat C9ORF72 BAC Mice. *Neuron*, **2020**, 108, 775-783.e4	13.9	13
93	A High-Content Screen Identifies TPP1 and Aurora B as Regulators of Axonal Mitochondrial Transport. *Cell Reports*, **2019**, 28, 3224-3237.e5	10.6	17
92	RNA-seq as a tool for evaluating human embryo competence. *Genome Research*, **2019**, 29, 1705-1718	9.7	9
91	Cancer-Related Mutations Identified in Primed and Naive Human Pluripotent Stem Cells. *Cell Stem Cell*, **2019**, 25, 456-461	18	15
90	Dysregulated protocadherin-pathway activity as an intrinsic defect in induced pluripotent stem cell-derived cortical interneurons from subjects with schizophrenia. *Nature Neuroscience*, **2019**, 22, 229-242	25.5	50

89	Comparison of three congruent patient-specific cell types for the modelling of a human genetic Schwann-cell disorder. *Nature Biomedical Engineering*, **2019**, 3, 571-582	19	9
88	Herpesviral lytic gene functions render the viral genome susceptible to novel editing by CRISPR/Cas9. *ELife*, **2019**, 8,	8.9	19
87	Exome sequencing in amyotrophic lateral sclerosis implicates a novel gene, DNAJC7, encoding a heat-shock protein. *Nature Neuroscience*, **2019**, 22, 1966-1974	25.5	56
86	A Stem Cell-Based Screening Platform Identifies Compounds that Desensitize Motor Neurons to Endoplasmic Reticulum Stress. *Molecular Therapy*, **2019**, 27, 87-101	11.7	12
85	ALS-implicated protein TDP-43 sustains levels of STMN2, a mediator of motor neuron growth and repair. *Nature Neuroscience*, **2019**, 22, 167-179	25.5	154
84	Dipeptide repeat proteins activate a heat shock response found in C9ORF72-ALS/FTLD patients. *Acta Neuropathologica Communications*, **2018**, 6, 55	7.3	15
83	Convergence of independent DISC1 mutations on impaired neurite growth via decreased UNC5D expression. *Translational Psychiatry*, **2018**, 8, 245	8.6	12
82	All-optical synaptic electrophysiology probes mechanism of ketamine-induced disinhibition. *Nature Methods*, **2018**, 15, 823-831	21.6	22
81	Oligodendrocyte differentiation of induced pluripotent stem cells derived from subjects with schizophrenias implicate abnormalities in development. *Translational Psychiatry*, **2018**, 8, 230	8.6	25
80	Comparative genomic analysis of embryonic, lineage-converted and stem cell-derived motor neurons. *Development (Cambridge)*, **2018**, 145,	6.6	8
79	TDP-43 induces p53-mediated cell death of cortical progenitors and immature neurons. *Scientific Reports*, **2018**, 8, 8097	4.9	22
78	All-Optical Electrophysiology for High-Throughput Functional Characterization of a Human iPSC-Derived Motor Neuron Model of ALS. *Stem Cell Reports*, **2018**, 10, 1991-2004	8	34
77	Combining NGN2 Programming with Developmental Patterning Generates Human Excitatory Neurons with NMDAR-Mediated Synaptic Transmission. *Cell Reports*, **2018**, 23, 2509-2523	10.6	90
76	The C9orf72-interacting protein Smcr8 is a negative regulator of autoimmunity and lysosomal exocytosis. *Genes and Development*, **2018**, 32, 929-943	12.6	41
75	Human pluripotent stem cells recurrently acquire and expand dominant negative P53 mutations. *Nature*, **2017**, 545, 229-233	50.4	270
74	Modelling Zika Virus Infection of the Developing Human Brain In Vitro Using Stem Cell Derived Cerebral Organoids. *Journal of Visualized Experiments*, **2017**,	1.6	12
73	A Scaled Framework for CRISPR Editing of Human Pluripotent Stem Cells to Study Psychiatric Disease. *Stem Cell Reports*, **2017**, 9, 1315-1327	8	7
72	Reactive Astrocytes Promote ALS-like Degeneration and Intracellular Protein Aggregation in Human Motor Neurons by Disrupting Autophagy through TGF-β1. *Stem Cell Reports*, **2017**, 9, 667-680	8	61

71	Generation of a Motor Nerve Organoid with Human Stem Cell-Derived Neurons. *Stem Cell Reports*, **2017**, 9, 1441-1449	8	60
70	Loss-of-function mutations in the C9ORF72 mouse ortholog cause fatal autoimmune disease. *Science Translational Medicine*, **2016**, 8, 347ra93	17.5	157
69	Comprehensive Protocols for CRISPR/Cas9-based Gene Editing in Human Pluripotent Stem Cells. *Current Protocols in Stem Cell Biology*, **2016**, 38, 5B.6.1-5B.6.60	2.8	23
68	Generation of a TLE1 homozygous knockout human embryonic stem cell line using CRISPR-Cas9. *Stem Cell Research*, **2016**, 17, 430-432	1.6	3
67	CAT7 and cat7l Long Non-coding RNAs Tune Polycomb Repressive Complex 1 Function during Human and Zebrafish Development. *Journal of Biological Chemistry*, **2016**, 291, 19558-72	5.4	26
66	Two familial ALS proteins function in prevention/repair of transcription-associated DNA damage. *Proceedings of the National Academy of Sciences of the United States of America*, **2016**, 113, E7701-E7709	11.5	71
65	Monitoring peripheral nerve degeneration in ALS by label-free stimulated Raman scattering imaging. *Nature Communications*, **2016**, 7, 13283	17.4	56
64	SLC52A3, A Brown-Vialetto-van Laere syndrome candidate gene is essential for mouse development, but dispensable for motor neuron differentiation. *Human Molecular Genetics*, **2016**, 25, 1814-23	5.6	8
63	Genetic Ablation of AXL Does Not Protect Human Neural Progenitor Cells and Cerebral Organoids from Zika Virus Infection. *Cell Stem Cell*, **2016**, 19, 703-708	18	185
62	Modeling ALS with motor neurons derived from human induced pluripotent stem cells. *Nature Neuroscience*, **2016**, 19, 542-53	25.5	174
61	Generation of a TLE3 heterozygous knockout human embryonic stem cell line using CRISPR-Cas9. *Stem Cell Research*, **2016**, 17, 441-443	1.6	4
60	From Dish to Bedside: Lessons Learned While Translating Findings from a Stem Cell Model of Disease to a Clinical Trial. *Cell Stem Cell*, **2015**, 17, 8-10	18	59
59	Efficient CRISPR-Cas9-mediated generation of knockin human pluripotent stem cells lacking undesired mutations at the targeted locus. *Cell Reports*, **2015**, 11, 875-883	10.6	111
58	A perspective on stem cell modeling of amyotrophic lateral sclerosis. *Cell Cycle*, **2015**, 14, 3679-88	4.7	15
57	Modeling pain in vitro using nociceptor neurons reprogrammed from fibroblasts. *Nature Neuroscience*, **2015**, 18, 17-24	25.5	135
56	Generation of neuropeptidergic hypothalamic neurons from human pluripotent stem cells. *Development (Cambridge)*, **2015**, 142, 633-43	6.6	93
55	ALS-causative mutations in FUS/TLS confer gain and loss of function by altered association with SMN and U1-snRNP. *Nature Communications*, **2015**, 6, 6171	17.4	162
54	Motoneurons derived from induced pluripotent stem cells develop mature phenotypes typical of endogenous spinal motoneurons. *Journal of Neuroscience*, **2015**, 35, 1291-306	6.6	38

53	Intrinsic membrane hyperexcitability of amyotrophic lateral sclerosis patient-derived motor neurons. *Cell Reports*, **2014**, 7, 1-11	10.6	444
52	Pathways disrupted in human ALS motor neurons identified through genetic correction of mutant SOD1. *Cell Stem Cell*, **2014**, 14, 781-95	18	300
51	How to make spinal motor neurons. *Development (Cambridge)*, **2014**, 141, 491-501	6.6	92
50	iPSC-derived dopamine neurons reveal differences between monozygotic twins discordant for Parkinson's disease. *Cell Reports*, **2014**, 9, 1173-82	10.6	166
49	Genetic variation in human DNA replication timing. *Cell*, **2014**, 159, 1015-1026	56.2	102
48	DNA methylation dynamics of the human preimplantation embryo. *Nature*, **2014**, 511, 611-5	50.4	390
47	Axonal transport of TDP-43 mRNA granules is impaired by ALS-causing mutations. *Neuron*, **2014**, 81, 536-543	13.9	408
46	Notch inhibition allows oncogene-independent generation of iPS cells. *Nature Chemical Biology*, **2014**, 10, 632-639	11.7	48
45	FUS is sequestered in nuclear aggregates in ALS patient fibroblasts. *Molecular Biology of the Cell*, **2014**, 25, 2571-8	3.5	40
44	The role of maternal-specific H3K9me3 modification in establishing imprinted X-chromosome inactivation and embryogenesis in mice. *Nature Communications*, **2014**, 5, 5464	17.4	43
43	Genetic validation of a therapeutic target in a mouse model of ALS. *Science Translational Medicine*, **2014**, 6, 248ra104	17.5	21
42	Ketamine exposure in early development impairs specification of the primary germ cell layers. *Neurotoxicology and Teratology*, **2014**, 43, 59-68	3.9	8
41	Nanog-independent reprogramming to iPSCs with canonical factors. *Stem Cell Reports*, **2014**, 2, 119-26	8	34
40	Opportunities and challenges of pluripotent stem cell neurodegenerative disease models. *Nature Neuroscience*, **2013**, 16, 780-9	25.5	156
39	Reference Maps of human ES and iPS cell variation enable high-throughput characterization of pluripotent cell lines. *Cell*, **2011**, 144, 439-52	56.2	756
38	Conversion of mouse and human fibroblasts into functional spinal motor neurons. *Cell Stem Cell*, **2011**, 9, 205-18	18	504
37	Constructing and deconstructing stem cell models of neurological disease. *Neuron*, **2011**, 70, 626-44	13.9	124
36	Somatic coding mutations in human induced pluripotent stem cells. *Nature*, **2011**, 471, 63-7	50.4	998

35	A functionally characterized test set of human induced pluripotent stem cells. *Nature Biotechnology* , **2011**, 29, 279-86	44.5	379
34	A small-molecule inhibitor of tgf-Beta signaling replaces sox2 in reprogramming by inducing nanog. *Cell Stem Cell*, **2009**, 5, 491-503	18	650
33	Human embryonic stem cell-derived motor neurons are sensitive to the toxic effect of glial cells carrying an ALS-causing mutation. *Cell Stem Cell*, **2008**, 3, 637-48	18	365
32	Dolly's legacy: human nuclear transplantation and better medicines for our children. *Cloning and Stem Cells*, **2007**, 9, 21-5		3
31	Non-cell autonomous effect of glia on motor neurons in an embryonic stem cell-based ALS model. *Nature Neuroscience*, **2007**, 10, 608-14	25.5	626
30	Short-circuiting epiblast development. *Cell Stem Cell*, **2007**, 1, 131-2	18	2
29	Ovulated oocytes in adult mice derive from non-circulating germ cells. *Nature*, **2006**, 441, 1109-14	50.4	202
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