Kevin C Eggan

List of Publications by Citations

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| # | Paper | IF | Citations |
|-----|--|--------|-----------|
| 106 | Somatic coding mutations in human induced pluripotent stem cells. <i>Nature</i> , 2011 , 471, 63-7 | 50.4 | 998 |
| 105 | Reference Maps of human ES and iPS cell variation enable high-throughput characterization of pluripotent cell lines. <i>Cell</i> , 2011 , 144, 439-52 | 56.2 | 756 |
| 104 | A small-molecule inhibitor of tgf-Beta signaling replaces sox2 in reprogramming by inducing nanog. <i>Cell Stem Cell</i> , 2009 , 5, 491-503 | 18 | 650 |
| 103 | Non-cell autonomous effect of glia on motor neurons in an embryonic stem cell-based ALS model. <i>Nature Neuroscience</i> , 2007 , 10, 608-14 | 25.5 | 626 |
| 102 | Conversion of mouse and human fibroblasts into functional spinal motor neurons. <i>Cell Stem Cell</i> , 2011 , 9, 205-18 | 18 | 504 |
| 101 | Intrinsic membrane hyperexcitability of amyotrophic lateral sclerosis patient-derived motor neurons. <i>Cell Reports</i> , 2014 , 7, 1-11 | 10.6 | 444 |
| 100 | Axonal transport of TDP-43 mRNA granules is impaired by ALS-causing mutations. <i>Neuron</i> , 2014 , 81, 53 | 615,43 | 408 |
| 99 | DNA methylation dynamics of the human preimplantation embryo. <i>Nature</i> , 2014 , 511, 611-5 | 50.4 | 390 |
| 98 | A functionally characterized test set of human induced pluripotent stem cells. <i>Nature Biotechnology</i> , 2011 , 29, 279-86 | 44.5 | 379 |
| 97 | Human embryonic stem cell-derived motor neurons are sensitive to the toxic effect of glial cells carrying an ALS-causing mutation. <i>Cell Stem Cell</i> , 2008 , 3, 637-48 | 18 | 365 |
| 96 | Pathways disrupted in human ALS motor neurons identified through genetic correction of mutant SOD1. <i>Cell Stem Cell</i> , 2014 , 14, 781-95 | 18 | 300 |
| 95 | Human pluripotent stem cells recurrently acquire and expand dominant negative P53 mutations. <i>Nature</i> , 2017 , 545, 229-233 | 50.4 | 270 |
| 94 | Ovulated oocytes in adult mice derive from non-circulating germ cells. <i>Nature</i> , 2006 , 441, 1109-14 | 50.4 | 202 |
| 93 | Genetic Ablation of AXL Does Not Protect Human Neural Progenitor Cells and Cerebral Organoids from Zika Virus Infection. <i>Cell Stem Cell</i> , 2016 , 19, 703-708 | 18 | 185 |
| 92 | Modeling ALS with motor neurons derived from human induced pluripotent stem cells. <i>Nature Neuroscience</i> , 2016 , 19, 542-53 | 25.5 | 174 |
| 91 | iPSC-derived dopamine neurons reveal differences between monozygotic twins discordant for Parkinson\ddot\ddot\ddot\ddot\ddot\ddot\ddot\dd | 10.6 | 166 |
| 90 | ALS-causative mutations in FUS/TLS confer gain and loss of function by altered association with SMN and U1-snRNP. <i>Nature Communications</i> , 2015 , 6, 6171 | 17.4 | 162 |

(2019-2016)

| 89 | Loss-of-function mutations in the C9ORF72 mouse ortholog cause fatal autoimmune disease. <i>Science Translational Medicine</i> , 2016 , 8, 347ra93 | 17.5 | 157 |
|----|---|---------------------------------|-----|
| 88 | Opportunities and challenges of pluripotent stem cell neurodegenerative disease models. <i>Nature Neuroscience</i> , 2013 , 16, 780-9 | 25.5 | 156 |
| 87 | ALS-implicated protein TDP-43 sustains levels of STMN2, a mediator of motor neuron growth and repair. <i>Nature Neuroscience</i> , 2019 , 22, 167-179 | 25.5 | 154 |
| 86 | Modeling pain in vitro using nociceptor neurons reprogrammed from fibroblasts. <i>Nature Neuroscience</i> , 2015 , 18, 17-24 | 25.5 | 135 |
| 85 | Constructing and deconstructing stem cell models of neurological disease. <i>Neuron</i> , 2011 , 70, 626-44 | 13.9 | 124 |
| 84 | Efficient CRISPR-Cas9-mediated generation of knockin human pluripotent stem cells lacking undesired mutations at the targeted locus. <i>Cell Reports</i> , 2015 , 11, 875-883 | 10.6 | 111 |
| 83 | Genetic variation in human DNA replication timing. <i>Cell</i> , 2014 , 159, 1015-1026 | 56.2 | 102 |
| 82 | Generation of neuropeptidergic hypothalamic neurons from human pluripotent stem cells. <i>Development (Cambridge)</i> , 2015 , 142, 633-43 | 6.6 | 93 |
| 81 | How to make spinal motor neurons. <i>Development (Cambridge)</i> , 2014 , 141, 491-501 | 6.6 | 92 |
| 80 | Combining NGN2 Programming with Developmental Patterning Generates Human Excitatory Neurons with NMDAR-Mediated Synaptic Transmission. <i>Cell Reports</i> , 2018 , 23, 2509-2523 | 10.6 | 90 |
| 79 | C9orf72 suppresses systemic and neural inflammation induced by gut bacteria. <i>Nature</i> , 2020 , 582, 89-94 | 50.4 | 83 |
| 78 | 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 |
| 77 | Reactive Astrocytes Promote ALS-like Degeneration and Intracellular Protein Aggregation in Human Motor Neurons by Disrupting Autophagy through TGF-II. Stem Cell Reports, 2017, 9, 667-680 | 8 | 61 |
| 76 | Generation of a Motor Nerve Organoid with Human Stem Cell-Derived Neurons. <i>Stem Cell Reports</i> , 2017 , 9, 1441-1449 | 8 | 60 |
| 75 | From Dish to Bedside: Lessons Learned While Translating Findings from a Stem Cell Model of Disease to a Clinical Trial. <i>Cell Stem Cell</i> , 2015 , 17, 8-10 | 18 | 59 |
| 74 | Monitoring peripheral nerve degeneration in ALS by label-free stimulated Raman scattering imaging. <i>Nature Communications</i> , 2016 , 7, 13283 | 17.4 | 56 |
| 73 | Exome sequencing in amyotrophic lateral sclerosis implicates a novel gene, DNAJC7, encoding a heat-shock protein. <i>Nature Neuroscience</i> , 2019 , 22, 1966-1974 | 25.5 | 56 |
| 72 | Dysregulated protocadherin-pathway activity as an intrinsic defect in induced pluripotent stem cell-derived cortical interneurons from subjects with schizophrenia. <i>Nature Neuroscience</i> , 2019 , 22, 229- | 2 ² 452 ⁵ | 50 |

| 71 | Notch inhibition allows oncogene-independent generation of iPS cells. <i>Nature Chemical Biology</i> , 2014 , 10, 632-639 | 11.7 | 48 |
|----|--|------|----|
| 70 | The role of maternal-specific H3K9me3 modification in establishing imprinted X-chromosome inactivation and embryogenesis in mice. <i>Nature Communications</i> , 2014 , 5, 5464 | 17.4 | 43 |
| 69 | The C9orf72-interacting protein Smcr8 is a negative regulator of autoimmunity and lysosomal exocytosis. <i>Genes and Development</i> , 2018 , 32, 929-943 | 12.6 | 41 |
| 68 | FUS is sequestered in nuclear aggregates in ALS patient fibroblasts. <i>Molecular Biology of the Cell</i> , 2014 , 25, 2571-8 | 3.5 | 40 |
| 67 | Motoneurons derived from induced pluripotent stem cells develop mature phenotypes typical of endogenous spinal motoneurons. <i>Journal of Neuroscience</i> , 2015 , 35, 1291-306 | 6.6 | 38 |
| 66 | Nanog-independent reprogramming to iPSCs with canonical factors. <i>Stem Cell Reports</i> , 2014 , 2, 119-26 | 8 | 34 |
| 65 | All-Optical Electrophysiology for High-Throughput Functional Characterization of a Human iPSC-Derived Motor Neuron Model of ALS. <i>Stem Cell Reports</i> , 2018 , 10, 1991-2004 | 8 | 34 |
| 64 | CAT7 and cat7l Long Non-coding RNAs Tune Polycomb Repressive Complex 1 Function during Human and Zebrafish Development. <i>Journal of Biological Chemistry</i> , 2016 , 291, 19558-72 | 5.4 | 26 |
| 63 | Oligodendrocyte differentiation of induced pluripotent stem cells derived from subjects with schizophrenias implicate abnormalities in development. <i>Translational Psychiatry</i> , 2018 , 8, 230 | 8.6 | 25 |
| 62 | Comprehensive Protocols for CRISPR/Cas9-based Gene Editing in Human Pluripotent Stem Cells. <i>Current Protocols in Stem Cell Biology</i> , 2016 , 38, 5B.6.1-5B.6.60 | 2.8 | 23 |
| 61 | All-optical synaptic electrophysiology probes mechanism of ketamine-induced disinhibition. <i>Nature Methods</i> , 2018 , 15, 823-831 | 21.6 | 22 |
| 60 | TDP-43 induces p53-mediated cell death of cortical progenitors and immature neurons. <i>Scientific Reports</i> , 2018 , 8, 8097 | 4.9 | 22 |
| 59 | Genetic validation of a therapeutic target in a mouse model of ALS. <i>Science Translational Medicine</i> , 2014 , 6, 248ra104 | 17.5 | 21 |
| 58 | Herpesviral lytic gene functions render the viral genome susceptible to novel editing by CRISPR/Cas9. <i>ELife</i> , 2019 , 8, | 8.9 | 19 |
| 57 | A High-Content Screen Identifies TPP1 and Aurora B as Regulators of Axonal Mitochondrial Transport. <i>Cell Reports</i> , 2019 , 28, 3224-3237.e5 | 10.6 | 17 |
| 56 | Cancer-Related Mutations Identified in Primed and Naive Human Pluripotent Stem Cells. <i>Cell Stem Cell</i> , 2019 , 25, 456-461 | 18 | 15 |
| 55 | A perspective on stem cell modeling of amyotrophic lateral sclerosis. <i>Cell Cycle</i> , 2015 , 14, 3679-88 | 4.7 | 15 |
| 54 | Dipeptide repeat proteins activate a heat shock response found in C9ORF72-ALS/FTLD patients. <i>Acta Neuropathologica Communications</i> , 2018 , 6, 55 | 7.3 | 15 |

| 53 | Absence of Survival and Motor Deficits in 500 Repeat C9ORF72 BAC Mice. Neuron, 2020, 108, 775-783. | 24 13.9 | 13 |
|----|---|----------------|----|
| 52 | Modelling Zika Virus Infection of the Developing Human Brain In Vitro Using Stem Cell Derived Cerebral Organoids. <i>Journal of Visualized Experiments</i> , 2017 , | 1.6 | 12 |
| 51 | A Stem Cell-Based Screening Platform Identifies Compounds that Desensitize Motor Neurons to Endoplasmic Reticulum Stress. <i>Molecular Therapy</i> , 2019 , 27, 87-101 | 11.7 | 12 |
| 50 | Convergence of independent DISC1 mutations on impaired neurite growth via decreased UNC5D expression. <i>Translational Psychiatry</i> , 2018 , 8, 245 | 8.6 | 12 |
| 49 | Human amyotrophic lateral sclerosis excitability phenotype screen: Target discovery and validation. <i>Cell Reports</i> , 2021 , 35, 109224 | 10.6 | 11 |
| 48 | Connecting TDP-43 Pathology with Neuropathy. <i>Trends in Neurosciences</i> , 2021 , 44, 424-440 | 13.3 | 10 |
| 47 | RNA-seq as a tool for evaluating human embryo competence. <i>Genome Research</i> , 2019 , 29, 1705-1718 | 9.7 | 9 |
| 46 | Comparison of three congruent patient-specific cell types for the modelling of a human genetic Schwann-cell disorder. <i>Nature Biomedical Engineering</i> , 2019 , 3, 571-582 | 19 | 9 |
| 45 | SLC52A3, A Brown-Vialetto-van Laere syndrome candidate gene is essential for mouse development, but dispensable for motor neuron differentiation. <i>Human Molecular Genetics</i> , 2016 , 25, 1814-23 | 5.6 | 8 |
| 44 | Ketamine exposure in early development impairs specification of the primary germ cell layers. <i>Neurotoxicology and Teratology</i> , 2014 , 43, 59-68 | 3.9 | 8 |
| 43 | Comparative genomic analysis of embryonic, lineage-converted and stem cell-derived motor neurons. <i>Development (Cambridge)</i> , 2018 , 145, | 6.6 | 8 |
| 42 | A Scaled Framework for CRISPR Editing of Human Pluripotent Stem Cells to Study Psychiatric Disease. <i>Stem Cell Reports</i> , 2017 , 9, 1315-1327 | 8 | 7 |
| 41 | Cancer-Related Mutations Identified in Primed Human Pluripotent Stem Cells. <i>Cell Stem Cell</i> , 2021 , 28, 10-11 | 18 | 7 |
| 40 | The genetic architecture of DNA replication timing in human pluripotent stem cells. <i>Nature Communications</i> , 2021 , 12, 6746 | 17.4 | 4 |
| 39 | Generation of a TLE3 heterozygous knockout human embryonic stem cell line using CRISPR-Cas9. <i>Stem Cell Research</i> , 2016 , 17, 441-443 | 1.6 | 4 |
| 38 | Generation of a TLE1 homozygous knockout human embryonic stem cell line using CRISPR-Cas9. <i>Stem Cell Research</i> , 2016 , 17, 430-432 | 1.6 | 3 |
| 37 | Dolly\legacy: human nuclear transplantation and better medicines for our children. <i>Cloning and Stem Cells</i> , 2007 , 9, 21-5 | | 3 |
| 36 | In vitro Differentiation of Human ES Cells149-167 | | 3 |

| 35 | Publicly available hiPSC lines with extreme polygenic risk scores for modeling schizophrenia | | 3 |
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| 34 | Genoppi is an open-source software for robust and standardized integration of proteomic and genetic data. <i>Nature Communications</i> , 2021 , 12, 2580 | 17.4 | 3 |
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| 30 | Short-circuiting epiblast development. <i>Cell Stem Cell</i> , 2007 , 1, 131-2 | 18 | 2 |
| 29 | Biological insights from the whole genome analysis of human embryonic stem cells | | 2 |
| 28 | Loss of mouse Stmn2 function causes motor neuropathy <i>Neuron</i> , 2022 , | 13.9 | 2 |
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| 22 | De novo DNA methyltransferases DNMT3A and DNMT3B are essential for XIST silencing for erosion of dosage compensation in pluripotent stem cells. <i>Stem Cell Reports</i> , 2021 , 16, 2138-2148 | 8 | 1 |
| 21 | Considerations and practical implications of performing a phenotypic CRISPR/Cas survival screen <i>PLoS ONE</i> , 2022 , 17, e0263262 | 3.7 | 1 |
| 20 | Whole-genome analysis of human embryonic stem cells enables rational line selection based on genetic variation <i>Cell Stem Cell</i> , 2022 , | 18 | 1 |
| 19 | Cx43 hemichannels contribute to astrocyte-mediated toxicity in sporadic and familial ALS <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022 , 119, e210739111 | 9 ^{11.5} | 0 |
| 18 | Spinal motor neuron transplantation to enhance nerve reconstruction strategies: Towards a cell therapy <i>Experimental Neurology</i> , 2022 , 353, 114054 | 5.7 | O |

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