

# Jason A Mills

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6264399/publications.pdf>

Version: 2024-02-01

28  
papers

701  
citations

567281

15  
h-index

580821

25  
g-index

30  
all docs

30  
docs citations

30  
times ranked

1457  
citing authors

#	ARTICLE	IF	CITATIONS
1	Generation of human control iPSC line CHOPi004-A from juvenile foreskin fibroblast cells. <i>Stem Cell Research</i> , 2020, 49, 102084.	0.7	2
2	Functional Cortical Axon Tracts Generated from Human Stem Cell-Derived Neurons. <i>Tissue Engineering - Part A</i> , 2019, 25, 736-745.	3.1	10
3	CRISPR Activation Enhances In Vitro Potency of AAV Vectors Driven by Tissue-Specific Promoters. <i>Molecular Therapy - Methods and Clinical Development</i> , 2019, 13, 380-389.	4.1	35
4	Comparative AAV-eGFP Transgene Expression Using Vector Serotypes 1, 9, 7m8, and 8b in Human Pluripotent Stem Cells, RPEs, and Human and Rat Cortical Neurons. <i>Stem Cells International</i> , 2019, 2019, 1-11.	2.5	24
5	A Mini Review: Moving iPSC-Derived Retinal Subtypes Forward for Clinical Applications for Retinal Degenerative Diseases. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1185, 557-561.	1.6	2
6	Use of induced pluripotent stem cell models to probe the pathogenesis of Choroideremia and to develop a potential treatment. <i>Stem Cell Research</i> , 2018, 27, 140-150.	0.7	37
7	NIPBL+/haploinsufficiency reveals a constellation of transcriptome disruptions in the pluripotent and cardiac states. <i>Scientific Reports</i> , 2018, 8, 1056.	3.3	26
8	Amelioration of Neurosensory Structure and Function in Animal and Cellular Models of a Congenital Blindness. <i>Molecular Therapy</i> , 2018, 26, 1581-1593.	8.2	19
9	Strategies for retinal cell generation from human pluripotent stem cells. <i>Stem Cell Investigation</i> , 2017, 4, 65-65.	3.0	13
10	Personalized models reveal mechanistic and therapeutic insights into CEP290-associated Leber congenital amaurosis. <i>Stem Cell Investigation</i> , 2016, 3, 65-65.	3.0	2
11	Generation of Hermansky-Pudlak Syndrome Type 1 (HPS1) induced pluripotent stem cells (iPSCs). <i>Stem Cell Research</i> , 2016, 16, 233-235.	0.7	7
12	Generation of human control iPSC cell line CHOPWT9 from healthy adult peripheral blood mononuclear cells. <i>Stem Cell Research</i> , 2016, 16, 14-16.	0.7	3
13	Retinas in a Dish Peek into Inherited Retinal Degeneration. <i>Cell Stem Cell</i> , 2016, 18, 688-689.	11.1	4
14	Generation of Hermansky Pudlak syndrome type 2 (HPS2) induced pluripotent stem cells (iPSCs). <i>Stem Cell Research</i> , 2016, 16, 287-289.	0.7	2
15	Generation of poikiloderma with neutropenia (PN) induced pluripotent stem cells (iPSCs). <i>Stem Cell Research</i> , 2015, 15, 595-597.	0.7	2
16	Impaired Telomere Maintenance and Decreased Canonical WNT Signaling but Normal Ribosome Biogenesis in Induced Pluripotent Stem Cells from X-Linked Dyskeratosis Congenita Patients. <i>PLoS ONE</i> , 2015, 10, e0127414.	2.5	26
17	Dysregulation of the Transforming Growth Factor $\beta$ Pathway in Induced Pluripotent Stem Cells Generated from Patients with Diamond Blackfan Anemia. <i>PLoS ONE</i> , 2015, 10, e0134878.	2.5	27
18	Emergence of a Stage-Dependent Human Liver Disease Signature with Directed Differentiation of Alpha-1 Antitrypsin-Deficient iPSC Cells. <i>Stem Cell Reports</i> , 2015, 4, 873-885.	4.8	77

#	ARTICLE	IF	CITATIONS
19	OCT4 Coordinates with WNT Signaling to Pre-pattern Chromatin at the SOX17 Locus during Human ES Cell Differentiation into Definitive Endoderm. <i>Stem Cell Reports</i> , 2015, 5, 490-498.	4.8	29
20	The negative impact of Wnt signaling on megakaryocyte and primitive erythroid progenitors derived from human embryonic stem cells. <i>Stem Cell Research</i> , 2014, 12, 441-451.	0.7	49
21	Utilization of the AAVS1 safe harbor locus for hematopoietic specific transgene expression and gene knockdown in human ES cells. <i>Stem Cell Research</i> , 2014, 12, 630-637.	0.7	35
22	Efficient iPS Cell Generation from Blood Using Episomes and HDAC Inhibitors. <i>Journal of Visualized Experiments</i> , 2014, , e52009.	0.3	10
23	High-level transgene expression in induced pluripotent stem cell-derived megakaryocytes: correction of Glanzmann thrombasthenia. <i>Blood</i> , 2014, 123, 753-757.	1.4	54
24	Hematopoietic Differentiation of Pluripotent Stem Cells in Culture. <i>Methods in Molecular Biology</i> , 2014, 1185, 181-194.	0.9	42
25	Dysregulation of the TGF $\beta$ 2 Pathway in Induced Pluripotent Stem Cells (iPSCs) Generated from Patients with Diamond Blackfan Anemia (DBA). <i>Blood</i> , 2014, 124, 254-254.	1.4	0
26	Clonal genetic and hematopoietic heterogeneity among human-induced pluripotent stem cell lines. <i>Blood</i> , 2013, 122, 2047-2051.	1.4	75
27	AAV-Mediated Gene Therapy for Choroideremia: Preclinical Studies in Personalized Models. <i>PLoS ONE</i> , 2013, 8, e61396.	2.5	71
28	Tissue-Specific Transgene Expression in Induced Pluripotent Stem (iPS) Cell-Derived Megakaryocytes: Correction of Glanzmann Thrombasthenia (GT). <i>Blood</i> , 2012, 120, 387-387.	1.4	0