

Jim Selfridge

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

27
papers

3,584
citations

18
h-index

30
g-index

30
ext. papers

4,065
ext. citations

15.3
avg, IF

4.71
L-index

#	Paper	IF	Citations
27	Reversal of neurological defects in a mouse model of Rett syndrome. <i>Science</i> , 2007 , 315, 1143-7	33.3	898
26	CpG islands influence chromatin structure via the CpG-binding protein Cfp1. <i>Nature</i> , 2010 , 464, 1082-6	50.4	507
25	The role of MeCP2 in the brain. <i>Annual Review of Cell and Developmental Biology</i> , 2011 , 27, 631-52	12.6	342
24	Embryonic lethal phenotype reveals a function of TDG in maintaining epigenetic stability. <i>Nature</i> , 2011 , 470, 419-23	50.4	282
23	Enhanced CpG mutability and tumorigenesis in MBD4-deficient mice. <i>Science</i> , 2002 , 297, 403-5	33.3	266
22	Rett syndrome mutations abolish the interaction of MeCP2 with the NCoR/SMRT co-repressor. <i>Nature Neuroscience</i> , 2013 , 16, 898-902	25.5	252
21	Up-regulation of glucocorticoid-regulated genes in a mouse model of Rett syndrome. <i>Human Molecular Genetics</i> , 2005 , 14, 2247-56	5.6	152
20	Kaiso-deficient mice show resistance to intestinal cancer. <i>Molecular and Cellular Biology</i> , 2006 , 26, 199-208	14.8	136
19	Morphological and functional reversal of phenotypes in a mouse model of Rett syndrome. <i>Brain</i> , 2012 , 135, 2699-710	11.2	109
18	Base excision by thymine DNA glycosylase mediates DNA-directed cytotoxicity of 5-fluorouracil. <i>PLoS Biology</i> , 2009 , 7, e91	9.7	90
17	Radically truncated MeCP2 rescues Rett syndrome-like neurological defects. <i>Nature</i> , 2017 , 550, 398-401	50.4	84
16	MeCP2 recognizes cytosine methylated tri-nucleotide and di-nucleotide sequences to tune transcription in the mammalian brain. <i>PLoS Genetics</i> , 2017 , 13, e1006793	6	76
15	Postnatal inactivation reveals enhanced requirement for MeCP2 at distinct age windows. <i>Human Molecular Genetics</i> , 2012 , 21, 3806-14	5.6	75
14	The molecular basis of variable phenotypic severity among common missense mutations causing Rett syndrome. <i>Human Molecular Genetics</i> , 2016 , 25, 558-70	5.6	54
13	A dominant role for the methyl-CpG-binding protein Mbd2 in controlling Th2 induction by dendritic cells. <i>Nature Communications</i> , 2015 , 6, 6920	17.4	53
12	A single allele of Hdac2 but not Hdac1 is sufficient for normal mouse brain development in the absence of its paralog. <i>Development (Cambridge)</i> , 2014 , 141, 604-616	6.6	52
11	Reduced seizure threshold and altered network oscillatory properties in a mouse model of Rett syndrome. <i>Neuroscience</i> , 2013 , 231, 195-205	3.9	42

10	Exclusive expression of MeCP2 in the nervous system distinguishes between brain and peripheral Rett syndrome-like phenotypes. <i>Human Molecular Genetics</i> , 2016 , 25, 4389-4404	5.6	38
9	Mice with DNA repair gene Ercc1 deficiency in a neural crest lineage are a model for late-onset Hirschsprung disease. <i>DNA Repair</i> , 2010 , 9, 653-60	4.3	17
8	Toxicity of overexpressed MeCP2 is independent of HDAC3 activity. <i>Genes and Development</i> , 2018 , 32, 1514-1524	12.6	16
7	A mutation-led search for novel functional domains in MeCP2. <i>Human Molecular Genetics</i> , 2018 , 27, 2531-2545	5.6	14
6	Affinity for DNA Contributes to NLS Independent Nuclear Localization of MeCP2. <i>Cell Reports</i> , 2018 , 24, 2213-2220	10.6	14
5	Neuronal non-CG methylation is an essential target for MeCP2 function. <i>Molecular Cell</i> , 2021 , 81, 1260-1275.e12	17.6	5
4	SALL4 controls cell fate in response to DNA base composition. <i>Molecular Cell</i> , 2021 , 81, 845-858.e8	17.6	5
3	An Orphan CpG Island Drives Expression of a miRNA Precursor with an Important Role in Mouse Development. <i>Epigenomes</i> , 2019 , 3, 7	2.3	1
2	Domains of methylated CAC and CG target MeCP2 to tune transcription in the brain		1
1	Neuronal non-CG methylation is an essential target for MeCP2 function		1