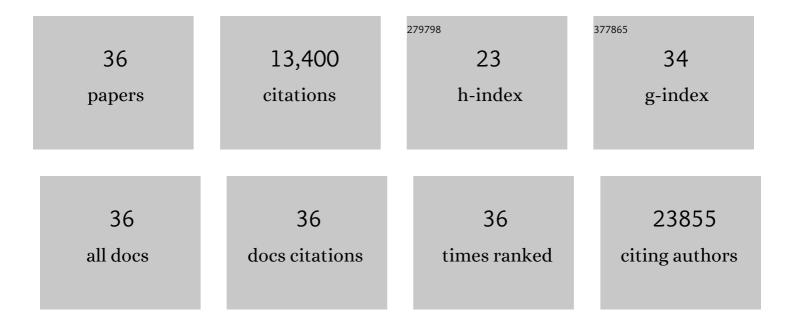
## Joan S Steffan

List of Publications by Year in descending order

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IOAN S STEEEAN

#	Article	IF	CITATIONS
1	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). Autophagy, 2016, 12, 1-222.	9.1	4,701
2	Guidelines for the use and interpretation of assays for monitoring autophagy. Autophagy, 2012, 8, 445-544.	9.1	3,122
3	Histone deacetylase inhibitors arrest polyglutamine-dependent neurodegeneration in Drosophila. Nature, 2001, 413, 739-743.	27.8	1,156
4	Suberoylanilide hydroxamic acid, a histone deacetylase inhibitor, ameliorates motor deficits in a mouse model of Huntington's disease. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 2041-2046.	7.1	805
5	SUMO Modification of Huntingtin and Huntington's Disease Pathology. Science, 2004, 304, 100-104.	12.6	627
6	IKK phosphorylates Huntingtin and targets it for degradation by the proteasome and lysosome. Journal of Cell Biology, 2009, 187, 1083-1099.	5.2	343
7	Nicotinamide Restores Cognition in Alzheimer's Disease Transgenic Mice via a Mechanism Involving Sirtuin Inhibition and Selective Reduction of Thr231-Phosphotau. Journal of Neuroscience, 2008, 28, 11500-11510.	3.6	339
8	Serines 13 and 16 Are Critical Determinants of Full-Length Human Mutant Huntingtin Induced Disease Pathogenesis in HD Mice. Neuron, 2009, 64, 828-840.	8.1	288
9	Inhibition of specific HDACs and sirtuins suppresses pathogenesis in a Drosophila model of Huntington's disease. Human Molecular Genetics, 2008, 17, 3767-3775.	2.9	248
10	The first 17 amino acids of Huntingtin modulate its sub-cellular localization, aggregation and effects on calcium homeostasis. Human Molecular Genetics, 2007, 16, 61-77.	2.9	247
11	Potential function for the Huntingtin protein as a scaffold for selective autophagy. Proceedings of the United States of America, 2014, 111, 16889-16894.	7.1	236
12	A bivalent Huntingtin binding peptide suppresses polyglutamine aggregation and pathogenesis in Drosophila. Nature Genetics, 2002, 30, 367-376.	21.4	167
13	Phosphorylation of Threonine 3. Journal of Biological Chemistry, 2009, 284, 29427-29436.	3.4	152
14	Ganglioside GM1 induces phosphorylation of mutant huntingtin and restores normal motor behavior in Huntington disease mice. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 3528-3533.	7.1	140
15	Selective histone deacetylase (HDAC) inhibition imparts beneficial effects in Huntington's disease mice: implications for the ubiquitin–proteasomal and autophagy systems. Human Molecular Genetics, 2012, 21, 5280-5293.	2.9	128
16	Treating the whole body in Huntington's disease. Lancet Neurology, The, 2015, 14, 1135-1142.	10.2	126
17	SUMO-2 and PIAS1 Modulate Insoluble Mutant Huntingtin Protein Accumulation. Cell Reports, 2013, 4, 362-375.	6.4	97
18	Human Neural Stem Cell Transplantation Rescues Functional Deficits in R6/2 and Q140 Huntington's Disease Mice. Stem Cell Reports, 2018, 10, 58-72.	4.8	76

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#	Article	IF	CITATIONS
19	PIAS1 Regulates Mutant Huntingtin Accumulation and Huntington's Disease-Associated Phenotypes InÂVivo. Neuron, 2016, 90, 507-520.	8.1	73
20	RRN11 Encodes the Third Subunit of the Complex Containing Rrn6p and Rrn7p That Is Essential for the Initiation of rDNA Transcription by Yeast RNA Polymerase I. Journal of Biological Chemistry, 1996, 271, 21062-21067.	3.4	70
21	Does Huntingtin play a role in selective macroautophagy?. Cell Cycle, 2010, 9, 3401-3413.	2.6	68
22	Serine 421 regulates mutant huntingtin toxicity and clearance in mice. Journal of Clinical Investigation, 2016, 126, 3585-3597.	8.2	44
23	PIAS1 modulates striatal transcription, DNA damage repair, and SUMOylation with relevance to Huntington's disease. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	28
24	IKKβ slows Huntington's disease progression in R6/1 mice. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 10952-10961.	7.1	23
25	Targeting aggregation in the development of therapeutics for the treatment of Huntington's disease and other polyglutamine repeat diseases. Expert Opinion on Therapeutic Targets, 2003, 7, 201-213.	3.4	18
26	Isoform-dependent lysosomal degradation and internalization of apolipoprotein E requires autophagy proteins. Journal of Cell Science, 2022, 135, .	2.0	16
27	Human Neural Progenitor Transplantation Rescues Behavior and Reduces α-Synuclein in a Transgenic Model of Dementia with Lewy Bodies. Stem Cells Translational Medicine, 2017, 6, 1477-1490.	3.3	14
28	Striatal Mutant Huntingtin Protein Levels Decline with Age in Homozygous Huntington's Disease Knock-In Mouse Models. Journal of Huntington's Disease, 2018, 7, 137-150.	1.9	14
29	Cooperation of cell adhesion and autophagy in the brain: Functional roles in development and neurodegenerative disease. Matrix Biology Plus, 2021, 12, 100089.	3.5	8
30	Serine residues 13 and 16 are key modulators of mutant huntingtin induced toxicity in Drosophila. Experimental Neurology, 2021, 338, 113463.	4.1	7
31	Diminished LC3-Associated Phagocytosis by Huntington's Disease Striatal Astrocytes. Journal of Huntington's Disease, 2022, 11, 25-33.	1.9	7
32	Transglutaminase 6 Is Colocalized and Interacts with Mutant Huntingtin in Huntington Disease Rodent Animal Models. International Journal of Molecular Sciences, 2021, 22, 8914.	4.1	6
33	Longitudinal Biochemical Assay Analysis of Mutant Huntingtin Exon 1 Protein in R6/2 Mice. Journal of Huntington's Disease, 2018, 7, 321-335.	1.9	5
34	A cause for childhood ataxia. ELife, 2016, 5, .	6.0	1
35	A20â€A role for transglutaminase 6 in hd pathology. , 2018, , .		0

APOE4 dysregulates autophagy in cultured cells. , 2022, 1, 29-33.

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