

# James A Mastrianni

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4649844/publications.pdf>

Version: 2024-02-01

15  
papers

5,972  
citations

1163117

8  
h-index

1125743

13  
g-index

17  
all docs

17  
docs citations

17  
times ranked

14360  
citing authors

#	ARTICLE	IF	CITATIONS
1	Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). <i>Autophagy</i> , 2016, 12, 1-222.	9.1	4,701
2	A Transmembrane Form of the Prion Protein in Neurodegenerative Disease. <i>Science</i> , 1998, 279, 827-834.	12.6	687
3	Prion Protein Conformation in a Patient with Sporadic Fatal Insomnia. <i>New England Journal of Medicine</i> , 1999, 340, 1630-1638.	27.0	186
4	The genetics of prion diseases. <i>Genetics in Medicine</i> , 2010, 12, 187-195.	2.4	111
5	The Prion Diseases. <i>Journal of Geriatric Psychiatry and Neurology</i> , 2010, 23, 277-298.	2.3	109
6	Rapamycin Delays Disease Onset and Prevents PrP Plaque Deposition in a Mouse Model of Gerstmann-Str�ussler-Scheinker Disease. <i>Journal of Neuroscience</i> , 2012, 32, 12396-12405.	3.6	92
7	A New Transgenic Mouse Model of Gerstmann-Str�ussler-Scheinker Syndrome Caused by the A117V Mutation of <i>PRNP</i> . <i>Journal of Neuroscience</i> , 2009, 29, 10072-10080.	3.6	51
8	The Prion Diseases: Creutzfeldt-Jakob, Gerstmann-Str�ussler-Scheinker, and Related Disorders. <i>Journal of Geriatric Psychiatry and Neurology</i> , 1998, 11, 78-97.	2.3	15
9	Impaired transmissibility of atypical prions from genetic CJDG114V. <i>Neurology: Genetics</i> , 2018, 4, e253.	1.9	7
10	IVIG Delays Onset in a Mouse Model of Gerstmann-Str�ussler-Scheinker Disease. <i>Molecular Neurobiology</i> , 2019, 56, 2353-2361.	4.0	4
11	Anle138b prevents PrP plaque accumulation in Tg(PrP-A116V) mice but does not mitigate clinical disease. <i>Journal of General Virology</i> , 2019, 100, 1027-1037.	2.9	4
12	A Novel TBK1 Variant (Lys694del) Presenting With Corticobasal Syndrome in a Family With FTD-ALS Spectrum Diseases: Case Report. <i>Frontiers in Neurology</i> , 2022, 13, 826676.	2.4	3
13	A novel <i>PRNP</i> -G131R variant associated with familial prion disease. <i>Neurology: Genetics</i> , 2020, 6, e454.	1.9	2
14	An atypical presentation of primary central nervous system lymphoma. <i>Medicine (United States)</i> , 2020, 99, e22062.	1.0	0
15	A Novel Human Disease with Abnormal Prion Protein Sensitive to Protease. <i>FASEB Journal</i> , 2008, 22, 173.3.	0.5	0