## Laura Pirisinu

List of Publications by Year in descending order

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516215 580395 25 25 837 16 h-index citations g-index papers 25 25 25 720 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Novel Type of Chronic Wasting Disease Detected in Moose ( <i>Alces alces</i> ), Norway. Emerging Infectious Diseases, 2018, 24, 2210-2218.	2.0	106
2	Chronic Wasting Disease in Bank Voles: Characterisation of the Shortest Incubation Time Model for Prion Diseases. PLoS Pathogens, 2013, 9, e1003219.	2.1	88
3	Prion Disease in Dromedary Camels, Algeria. Emerging Infectious Diseases, 2018, 24, 1029-1036.	2.0	88
4	Studies in bank voles reveal strain differences between chronic wasting disease prions from Norway and North America. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 31417-31426.	3.3	57
5	Cofactors influence the biological properties of infectious recombinant prions. Acta Neuropathologica, 2018, 135, 179-199.	3.9	56
6	Gerstmann-StrÃ <b>¤</b> ssler-Scheinker disease subtypes efficiently transmit in bank voles as genuine prion diseases. Scientific Reports, 2016, 6, 20443.	1.6	54
7	A New Method for the Characterization of Strain-Specific Conformational Stability of Protease-Sensitive and Protease-Resistant PrPSc. PLoS ONE, 2010, 5, e12723.	1.1	42
8	Small Ruminant Nor98 Prions Share Biochemical Features with Human Gerstmann-StrÃøssler-Scheinker Disease and Variably Protease-Sensitive Prionopathy. PLoS ONE, 2013, 8, e66405.	1.1	37
9	PrP C Governs Susceptibility to Prion Strains in Bank Vole, While Other Host Factors Modulate Strain Features. Journal of Virology, 2016, 90, 10660-10669.	1.5	37
10	Chronic wasting disease in Europe: new strains on the horizon. Acta Veterinaria Scandinavica, 2021, 63, 48.	0.5	37
11	Atypical Creutzfeldt-Jakob disease with PrP-amyloid plaques in white matter: molecular characterization and transmission to bank voles show the M1 strain signature. Acta Neuropathologica Communications, 2017, 5, 87.	2.4	25
12	Variable Protease-Sensitive Prionopathy Transmission to Bank Voles. Emerging Infectious Diseases, 2019, 25, 73-81.	2.0	25
13	Characterization of goat prions demonstrates geographical variation of scrapie strains in Europe and reveals the composite nature of prion strains. Scientific Reports, 2020, 10, 19.	1.6	22
14	Isolation of infectious, non-fibrillar and oligomeric prions from a genetic prion disease. Brain, 2020, 143, 1512-1524.	3.7	21
15	Biochemical Characterization of Prion Strains in Bank Voles. Pathogens, 2013, 2, 446-456.	1.2	20
16	Molecular Discrimination of Sheep Bovine Spongiform Encephalopathy from Scrapie. Emerging Infectious Diseases, 2011, 17, 695-698.	2.0	19
17	Prions in Variably Protease-Sensitive Prionopathy: An Update. Pathogens, 2013, 2, 457-471.	1.2	19
18	Oral pravastatin prolongs survival time of scrapie-infected mice. Journal of General Virology, 2009, 90, 1775-1780.	1.3	16

#	Article	IF	CITATIONS
19	Transmissibility of Gerstmann–Strässler–Scheinker syndrome in rodent models: New insights into the molecular underpinnings of prion infectivity. Prion, 2016, 10, 421-433.	0.9	14
20	Isolation of a Defective Prion Mutant from Natural Scrapie. PLoS Pathogens, 2016, 12, e1006016.	2.1	14
21	Effect of PrP genotype and route of inoculation on the ability of discriminatory Western blot to distinguish scrapie from sheep bovine spongiform encephalopathy. Journal of General Virology, 2012, 93, 450-455.	1.3	11
22	Four types of scrapie in goats differentiated from each other and bovine spongiform encephalopathy by biochemical methods. Veterinary Research, 2019, 50, 97.	1.1	11
23	In vitro replication highlights the mutability of prions. Prion, 2014, 8, 154-160.	0.9	9
24	A single amino acid residue in bank vole prion protein drives permissiveness to Nor98/atypical scrapie and the emergence of multiple strain variants. PLoS Pathogens, 2022, 18, e1010646.	2.1	7
25	Stability of BSE infectivity towards heat treatment even after proteolytic removal of prion protein. Veterinary Research, 2021, 52, 59.	1.1	2