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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Tankyrase-mediated ADP-ribosylation is a regulator of TNF-induced death. Science Advances, 2022, 8, eabh2332.	10.3	9
2	Mitochondrial NAD+ Controls Nuclear ARTD1-Induced ADP-Ribosylation. Molecular Cell, 2021, 81, 340-354.e5.	9.7	31
3	Establishment of a Mass-Spectrometry-Based Method for the Identification of the <i>In Vivo</i> Whole Blood and Plasma ADP-Ribosylomes. Journal of Proteome Research, 2021, 20, 3090-3101.	3.7	7
4	Engineering Af1521 improves ADP-ribose binding and identification of ADP-ribosylated proteins. Nature Communications, 2020, 11, 5199.	12.8	49
5	Development of a Corneal Bioluminescence Mouse for Real-Time In Vivo Evaluation of Gene Therapies. Translational Vision Science and Technology, 2020, 9, 44.	2.2	2
6	Lysosomal protease deficiency or substrate overload induces an oxidative-stress mediated STAT3-dependent pathway of lysosomal homeostasis. Nature Communications, 2018, 9, 5343.	12.8	52
7	Comprehensive ADPâ€ribosylome analysis identifies tyrosine as an ADPâ€ribose acceptor site. EMBO Reports, 2018, 19, .	4.5	75
8	Identification of PARP-Specific ADP-Ribosylation Targets Reveals a Regulatory Function for ADP-Ribosylation in Transcription Elongation. Molecular Cell, 2016, 63, 181-183.	9.7	10
9	Keratin 12 missense mutation induces the unfolded protein response and apoptosis in Meesmann epithelial corneal dystrophy. Human Molecular Genetics, 2016, 25, 1176-1191.	2.9	22
10	Keratin 9 Is Required for the Structural Integrity and Terminal Differentiation of the Palmoplantar Epidermis. Journal of Investigative Dermatology, 2014, 134, 754-763.	0.7	87
11	In vivo gene silencing following non-invasive siRNA delivery into the skin using a novel topical formulation. Journal of Controlled Release, 2014, 196, 355-362.	9.9	34
12	siRNA Silencing of the Mutant Keratin 12 Allele in Corneal Limbal Epithelial Cells Grown From Patients With Meesmann's Epithelial Corneal Dystrophy., 2014, 55, 3352.		28
13	Allele-Specific siRNA Silencing for the Common Keratin 12 Founder Mutation in Meesmann Epithelial Corneal Dystrophy., 2013, 54, 494.		34
14	Generic and Personalized RNAi-Based Therapeutics for a Dominant-Negative Epidermal Fragility Disorder. Journal of Investigative Dermatology, 2012, 132, 1627-1635.	0.7	38