

# Ariadna Bargiela

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9257658/publications.pdf>

Version: 2024-02-01

14  
papers

1,687  
citations

1307594

7  
h-index

1199594

12  
g-index

14  
all docs

14  
docs citations

14  
times ranked

2344  
citing authors

#	ARTICLE	IF	CITATIONS
1	Neuroprotective properties of queen bee acid by autophagy induction. <i>Cell Biology and Toxicology</i> , 2023, 39, 751-770.	5.3	7
2	The hallmarks of myotonic dystrophy type 1 muscle dysfunction. <i>Biological Reviews</i> , 2021, 96, 716-730.	10.4	40
3	Practicing logical reasoning through <i>Drosophila</i> segmentation gene mutants. <i>Biochemistry and Molecular Biology Education</i> , 2021, 49, 729-736.	1.2	0
4	Defined d-hexapeptides bind CUG repeats and rescue phenotypes of myotonic dystrophy myotubes in a <i>Drosophila</i> model of the disease. <i>Scientific Reports</i> , 2021, 11, 19417.	3.3	0
5	Musashi-2 contributes to myotonic dystrophy muscle dysfunction by promoting excessive autophagy through miR-7 biogenesis repression. <i>Molecular Therapy - Nucleic Acids</i> , 2021, 25, 652-667.	5.1	12
6	Inhibition of autophagy rescues muscle atrophy in a LGMDD2 <i>Drosophila</i> model. <i>FASEB Journal</i> , 2021, 35, e21914.	0.5	6
7	Guidelines for the use and interpretation of assays for monitoring autophagy (4th) <i>Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50 502</i>	9.1	1,430
8	miR-7 Restores Phenotypes in Myotonic Dystrophy Muscle Cells by Repressing Hyperactivated Autophagy. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 19, 278-292.	5.1	29
9	Increased Muscleblind levels by chloroquine treatment improve myotonic dystrophy type 1 phenotypes in in vitro and in vivo models. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 25203-25213.	7.1	32
10	Six Serum miRNAs Fail to Validate as Myotonic Dystrophy Type 1 Biomarkers. <i>PLoS ONE</i> , 2016, 11, e0150501.	2.5	7
11	Increased autophagy and apoptosis contribute to muscle atrophy in a myotonic dystrophy type 1 <i>Drosophila</i> model. <i>DMM Disease Models and Mechanisms</i> , 2015, 8, 679-690.	2.4	74
12	Two Enhancers Control Transcription of <i>Drosophila</i> muscleblind in the Embryonic Somatic Musculature and in the Central Nervous System. <i>PLoS ONE</i> , 2014, 9, e93125.	2.5	13
13	Muscleblind, BSF and TBPH are mislocalized in the muscle sarcomere of a <i>Drosophila</i> myotonic dystrophy model. <i>DMM Disease Models and Mechanisms</i> , 2013, 6, 184-96.	2.4	36
14	A GFP-tagged Muscleblind C protein isoform reporter construct. <i>Fly</i> , 2010, 4, 333-337.	1.7	1