

# Christina A Pacak

## List of Publications by Year in descending order

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38  
papers

1,069  
citations

567281

15  
h-index

414414

32  
g-index

38  
all docs

38  
docs citations

38  
times ranked

1736  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic Fate of Recombinant Adeno-Associated Virus Vector Genomes in Muscle. <i>Journal of Virology</i> , 2003, 77, 3495-3504.	3.4	227
2	Actin-dependent mitochondrial internalization in cardiomyocytes: evidence for rescue of mitochondrial function. <i>Biology Open</i> , 2015, 4, 622-626.	1.2	125
3	Infiltrative and drug-resistant slow-cycling cells support metabolic heterogeneity in glioblastoma. <i>EMBO Journal</i> , 2018, 37, .	7.8	118
4	Tissue specific promoters improve specificity of AAV9 mediated transgene expression following intra-vascular gene delivery in neonatal mice. <i>Genetic Vaccines and Therapy</i> , 2008, 6, 13.	1.5	80
5	Rapid Isolation And Purification Of Mitochondria For Transplantation By Tissue Dissociation And Differential Filtration. <i>Journal of Visualized Experiments</i> , 2014, , e51682.	0.3	61
6	Long-term Skeletal Muscle Protection After Gene Transfer in a Mouse Model of LGMD-2D. <i>Molecular Therapy</i> , 2007, 15, 1775-1781.	8.2	45
7	AAV-Mediated TAZ Gene Replacement Restores Mitochondrial and Cardioskeletal Function in Barth Syndrome. <i>Human Gene Therapy</i> , 2019, 30, 139-154.	2.7	40
8	Impaired cardiac and skeletal muscle bioenergetics in children, adolescents, and young adults with Barth syndrome. <i>Physiological Reports</i> , 2017, 5, e13130.	1.7	33
9	Consequences of MEGF10 deficiency on myoblast function and Notch1 interactions. <i>Human Molecular Genetics</i> , 2017, 26, 2984-3000.	2.9	30
10	Blunted fat oxidation upon submaximal exercise is partially compensated by enhanced glucose metabolism in children, adolescents, and young adults with Barth syndrome. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 480-493.	3.6	24
11	Silencing of drpr Leads to Muscle and Brain Degeneration in Adult Drosophila. <i>American Journal of Pathology</i> , 2014, 184, 2653-2661.	3.8	23
12	Ultrarapid Purification of Collagen Type I for Tissue Engineering Applications. <i>Tissue Engineering - Part C: Methods</i> , 2011, 17, 879-885.	2.1	21
13	Myocardial glucose and fatty acid metabolism is altered and associated with lower cardiac function in young adults with Barth syndrome. <i>Journal of Nuclear Cardiology</i> , 2021, 28, 1649-1659.	2.1	21
14	18 F-labeled rhodamines as potential myocardial perfusion agents: comparison of pharmacokinetic properties of several rhodamines. <i>Nuclear Medicine and Biology</i> , 2015, 42, 796-803.	0.6	20
15	AAV9-TAZ Gene Replacement Ameliorates Cardiac TMT Proteomic Profiles in a Mouse Model of Barth Syndrome. <i>Molecular Therapy - Methods and Clinical Development</i> , 2019, 13, 167-179.	4.1	17
16	Impact of PYROXD1 deficiency on cellular respiration and correlations with genetic analyses of limb-girdle muscular dystrophy in Saudi Arabia and Sudan. <i>Physiological Genomics</i> , 2018, 50, 929-939.	2.3	15
17	A form of muscular dystrophy associated with pathogenic variants in JAG2. <i>American Journal of Human Genetics</i> , 2021, 108, 840-856.	6.2	15
18	Bioprocessing of Human Mesenchymal Stem Cells: From Planar Culture to Microcarrier-Based Bioreactors. <i>Bioengineering</i> , 2021, 8, 96.	3.5	15

#	ARTICLE	IF	CITATIONS
19	Superparamagnetic Iron Oxide Nanoparticles Function as a Long-Term, Multi-Modal Imaging Label for Non-Invasive Tracking of Implanted Progenitor Cells. <i>PLoS ONE</i> , 2014, 9, e108695.	2.5	14
20	An Improved Method for the Preparation of Type I Collagen From Skin. <i>Journal of Visualized Experiments</i> , 2014, , e51011.	0.3	12
21	Longitudinal Patterns of Thalidomide Neuropathy in Children and Adolescents. <i>Journal of Pediatrics</i> , 2016, 178, 227-232.	1.8	12
22	Transfer of Therapeutic Genes into Fetal Rhesus Monkeys Using Recombinant Adeno-Associated Type I Viral Vectors. <i>Human Gene Therapy Clinical Development</i> , 2016, 27, 152-159.	3.1	12
23	Straightening of curved pattern of collagen fibers under load controls aortic valve shape. <i>Journal of Biomechanics</i> , 2014, 47, 341-346.	2.1	11
24	PDK4 Deficiency Induces Intrinsic Apoptosis in Response to Starvation in Fibroblasts from Doberman Pinschers with Dilated Cardiomyopathy. <i>BioResearch Open Access</i> , 2017, 6, 182-191.	2.6	10
25	Functional Consequences of PDK4 Deficiency in Doberman Pinscher Fibroblasts. <i>Scientific Reports</i> , 2020, 10, 3930.	3.3	10
26	Increased mtDNA Abundance and Improved Function in Human Barth Syndrome Patient Fibroblasts Following AAV-TAZ Gene Delivery. <i>International Journal of Molecular Sciences</i> , 2019, 20, 3416.	4.1	9
27	Identification of a pathogenic mutation in ATP2A1 via in silico analysis of exome data for cryptic aberrant splice sites. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e552.	1.2	9
28	Megf10 deficiency impairs skeletal muscle stem cell migration and muscle regeneration. <i>FEBS Open Bio</i> , 2021, 11, 114-123.	2.3	8
29	Selective serotonin reuptake inhibitors ameliorate MEGF10 myopathy. <i>Human Molecular Genetics</i> , 2019, 28, 2365-2377.	2.9	7
30	Fabrication of Myogenic Engineered Tissue Constructs. <i>Journal of Visualized Experiments</i> , 2009, , .	0.3	5
31	Microcarrier-Based Expansion of Adult Murine Side Population Stem Cells. <i>PLoS ONE</i> , 2013, 8, e55187.	2.5	4
32	Generation of Induced Pluripotent Stem Cells from a Female Patient with a Xq27.3-q28 Deletion to Establish Disease Models and Identify Therapies. <i>Cellular Reprogramming</i> , 2020, 22, 179-188.	0.9	3
33	The past, present, and future of modeling Cockayne Syndrome – A commentary on – Rat Model of Cockayne Syndrome Neurological Disease – DNA Repair, 2020, 88, 102788.	2.8	3
34	A review of the underlying genetics and emerging therapies for canine cardiomyopathies. <i>Journal of Veterinary Cardiology</i> , 2022, 40, 2-14.	0.9	3
35	Growth of Bone Marrow and Skeletal Muscle Side Population Stem Cells in Suspension Culture. <i>Methods in Molecular Biology</i> , 2014, 1210, 51-61.	0.9	3
36	TMT Sample Preparation for Proteomics Facility Submission and Subsequent Data Analysis. <i>Journal of Visualized Experiments</i> , 2020, , .	0.3	2

#	ARTICLE	IF	CITATIONS
37	The End of the Beginning: The Journey to Molecular Therapies for Spinal Muscular Atrophy. <i>Pediatric Neurology</i> , 2020, 102, 1-2.	2.1	1
38	Noninvasive Tracking of Implanted Cells: Superparamagnetic Iron Oxide Nanoparticles as a Long-Term, Multimodal Imaging Label. <i>Methods in Molecular Biology</i> , 2020, 2126, 167-175.	0.9	1