

Laura C Greaves

List of Publications by Year in descending order

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Version: 2024-02-01

55
papers

4,615
citations

159358

30
h-index

168136

53
g-index

57
all docs

57
docs citations

57
times ranked

6531
citing authors

#	ARTICLE	IF	CITATIONS
1	Mitochondrial complex I subunit deficiency promotes pancreatic $\hat{\pm}$ -cell proliferation. <i>Molecular Metabolism</i> , 2022, 60, 101489.	3.0	1
2	Automated quantitative high-throughput multiplex immunofluorescence pipeline to evaluate OXPHOS defects in formalin-fixed human prostate tissue. <i>Scientific Reports</i> , 2022, 12, 6660.	1.6	2
3	Mitochondrial <scp>DNA</scp> mutations in ageing and cancer. <i>Molecular Oncology</i> , 2022, 16, 3276-3294.	2.1	18
4	Age-associated mitochondrial complex I deficiency is linked to increased stem cell proliferation rates in the mouse colon. <i>Aging Cell</i> , 2021, 20, e13321.	3.0	8
5	Aberrant mitochondrial function in ageing and cancer. <i>Biogerontology</i> , 2020, 21, 445-459.	2.0	17
6	Mitochondrial dysfunction impairs osteogenesis, increases osteoclast activity, and accelerates age related bone loss. <i>Scientific Reports</i> , 2020, 10, 11643.	1.6	58
7	Age-associated mitochondrial DNA mutations cause metabolic remodeling that contributes to accelerated intestinal tumorigenesis. <i>Nature Cancer</i> , 2020, 1, 976-989.	5.7	69
8	The rise and rise of mitochondrial DNA mutations. <i>Open Biology</i> , 2020, 10, 200061.	1.5	89
9	Design and baseline characteristics of the Biomarkers Of Risk In Colorectal Cancer (BORICC) Follow-Up study: A 12+ years follow-up. <i>Nutrition and Health</i> , 2019, 25, 231-238.	0.6	2
10	Effects of obesity and weight loss on mitochondrial structure and function and implications for colorectal cancer risk. <i>Proceedings of the Nutrition Society</i> , 2019, 78, 426-437.	0.4	17
11	A Bioreactor Technology for Modeling Fibrosis in Human and Rodent Precision-Cut Liver Slices. <i>Hepatology</i> , 2019, 70, 1377-1391.	3.6	66
12	Length-independent telomere damage drives post-mitotic cardiomyocyte senescence. <i>EMBO Journal</i> , 2019, 38, .	3.5	307
13	A novel histochemistry assay to assess and quantify focal cytochrome <i>c</i> oxidase deficiency. <i>Journal of Pathology</i> , 2018, 245, 311-323.	2.1	17
14	Predominant Asymmetrical Stem Cell Fate Outcome Limits the Rate of Niche Succession in Human Colonic Crypts. <i>EBioMedicine</i> , 2018, 31, 166-173.	2.7	19
15	Impact of Age-Related Mitochondrial Dysfunction and Exercise on Intestinal Microbiota Composition. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2018, 73, 571-578.	1.7	28
16	Roles of Mitochondrial DNA Mutations in Stem Cell Ageing. <i>Genes</i> , 2018, 9, 182.	1.0	19
17	Inherited pathogenic mitochondrial DNA mutations and gastrointestinal stem cell populations. <i>Journal of Pathology</i> , 2018, 246, 427-432.	2.1	13
18	Multipotent Basal Stem Cells, Maintained in Localized Proximal Niches, Support Directed Long-Ranging Epithelial Flows in Human Prostates. <i>Cell Reports</i> , 2017, 20, 1609-1622.	2.9	64

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19	Mitochondrial DNA changes in pedunculopontine cholinergic neurons in Parkinson disease. <i>Annals of Neurology</i> , 2017, 82, 1016-1021.	2.8	45
20	Mitochondria are required for pro-ageing features of the senescent phenotype. <i>EMBO Journal</i> , 2016, 35, 724-742.	3.5	527
21	A Phenotype-Driven Approach to Generate Mouse Models with Pathogenic mtDNA Mutations Causing Mitochondrial Disease. <i>Cell Reports</i> , 2016, 16, 2980-2990.	2.9	102
22	Involving older people in the design, development, and delivery of an innovative module on aging for undergraduate students. <i>Educational Gerontology</i> , 2016, 42, 698-705.	0.7	8
23	Unique quadruple immunofluorescence assay demonstrates mitochondrial respiratory chain dysfunction in osteoblasts of aged and PolgA ^{tm1.1j} mice. <i>Scientific Reports</i> , 2016, 6, 31907.	1.6	13
24	Novel <i>MTND1</i> mutations cause isolated exercise intolerance, complex I deficiency and increased assembly factor expression. <i>Clinical Science</i> , 2015, 128, 895-904.	1.8	21
25	The presence of highly disruptive 16S rRNA mutations in clinical samples indicates a wider role for mutations of the mitochondrial ribosome in human disease. <i>Mitochondrion</i> , 2015, 25, 17-27.	1.6	29
26	SCNT-Derived ESCs with Mismatched Mitochondria Trigger an Immune Response in Allogeneic Hosts. <i>Cell Stem Cell</i> , 2015, 16, 33-38.	5.2	52
27	Clonal Expansion of Early to Mid-Life Mitochondrial DNA Point Mutations Drives Mitochondrial Dysfunction during Human Ageing. <i>PLoS Genetics</i> , 2014, 10, e1004620.	1.5	115
28	Human stem cell aging: do mitochondrial DNA mutations have a causal role?. <i>Aging Cell</i> , 2014, 13, 201-205.	3.0	30
29	Bmi1 enhances skeletal muscle regeneration through MT1-mediated oxidative stress protection in a mouse model of dystrophinopathy. <i>Journal of Experimental Medicine</i> , 2014, 211, 2617-2633.	4.2	34
30	The role of the mitochondrial ribosome in human disease: searching for mutations in 12S mitochondrial rRNA with high disruptive potential. <i>Human Molecular Genetics</i> , 2014, 23, 949-967.	1.4	35
31	Chronic inflammation induces telomere dysfunction and accelerates ageing in mice. <i>Nature Communications</i> , 2014, 5, 4172.	5.8	596
32	Similar patterns of clonally expanded somatic mtDNA mutations in the colon of heterozygous mtDNA mutator mice and ageing humans. <i>Mechanisms of Ageing and Development</i> , 2014, 139, 22-30.	2.2	33
33	Bmi1 enhances skeletal muscle regeneration through MT1-mediated oxidative stress protection in a mouse model of dystrophinopathy. <i>Journal of Cell Biology</i> , 2014, 207, 2075OIA222.	2.3	0
34	Comparison of Mitochondrial Mutation Spectra in Ageing Human Colonic Epithelium and Disease: Absence of Evidence for Purifying Selection in Somatic Mitochondrial DNA Point Mutations. <i>PLoS Genetics</i> , 2012, 8, e1003082.	1.5	61
35	Mitochondrial DNA and disease. <i>Journal of Pathology</i> , 2012, 226, 274-286.	2.1	239
36	Differences in the accumulation of mitochondrial defects with age in mice and humans. <i>Mechanisms of Ageing and Development</i> , 2011, 132, 588-591.	2.2	26

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37	<i>In situ</i> lineage tracking of human prostatic epithelial stem cell fate reveals a common clonal origin for basal and luminal cells. <i>Journal of Pathology</i> , 2011, 225, 181-188.	2.1	62
38	Defects in multiple complexes of the respiratory chain are present in ageing human colonic crypts. <i>Experimental Gerontology</i> , 2010, 45, 573-579.	1.2	52
39	Age-associated mitochondrial DNA mutations lead to small but significant changes in cell proliferation and apoptosis in human colonic crypts. <i>Aging Cell</i> , 2010, 9, 96-99.	3.0	56
40	Somatic Mitochondrial DNA Deletions Accumulate to High Levels in Aging Human Extraocular Muscles. , 2010, 51, 3347.		48
41	Mitochondrial DNA Defects and Selective Extraocular Muscle Involvement in CPEO. , 2010, 51, 3340.		58
42	Modelling mitochondrial DNA mutations in bacterial cytochrome <i>c</i> oxidase: Link to colon cancer?. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, E57.	3.3	1
43	Locating the stem cell niche and tracing hepatocyte lineages in human liver. <i>Hepatology</i> , 2009, 49, 1655-1663.	3.6	135
44	A Methodological Approach to Tracing Cell Lineage in Human Epithelial Tissues. <i>Stem Cells</i> , 2009, 27, 1410-1420.	1.4	72
45	Quantification of mitochondrial DNA mutation load. <i>Aging Cell</i> , 2009, 8, 566-572.	3.0	36
46	Detection of cytochrome c oxidase activity and mitochondrial proteins in single cells. <i>Journal of Neuroscience Methods</i> , 2009, 184, 310-319.	1.3	30
47	Mitochondrial DNA mutations and ageing. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2009, 1790, 1015-1020.	1.1	45
48	Mechanisms of Field Cancerization in the Human Stomach: The Expansion and Spread of Mutated Gastric Stem Cells. <i>Gastroenterology</i> , 2008, 134, 500-510.	0.6	222
49	The ageing mitochondrial genome. <i>Nucleic Acids Research</i> , 2007, 35, 7399-7405.	6.5	76
50	Mitochondrial DNA Mutations and Aging. <i>Annals of the New York Academy of Sciences</i> , 2007, 1100, 227-240.	1.8	50
51	Mitochondrial DNA mutations in human disease. <i>IUBMB Life</i> , 2006, 58, 143-151.	1.5	37
52	Interactions of skin thickness and physicochemical properties of test compounds in percutaneous penetration studies. <i>International Archives of Occupational and Environmental Health</i> , 2006, 79, 405-413.	1.1	85
53	Clonal Expansion in the Human Gut: Mitochondrial DNA Mutations Show Us the Way. <i>Cell Cycle</i> , 2006, 5, 808-811.	1.3	43
54	Mitochondrial DNA mutations are established in human colonic stem cells, and mutated clones expand by crypt fission. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 714-719.	3.3	269

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55	Mitochondrial DNA mutations in human colonic crypt stem cells. <i>Journal of Clinical Investigation</i> , 2003, 112, 1351-1360.	3.9	454