

Susana S Lopes

List of Publications by Year in descending order

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Version: 2024-02-01

39
papers

2,160
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361413

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docs citations

44
times ranked

2671
citing authors

#	ARTICLE	IF	CITATIONS
1	Crosstalk between cilia and autophagy: implication for human diseases. <i>Autophagy</i> , 2023, 19, 24-43.	9.1	10
2	Current methods to analyze lysosome morphology, positioning, motility and function. <i>Traffic</i> , 2022, 23, 238-269.	2.7	37
3	CiliarMove: new software for evaluating ciliary beat frequency helps find novel mutations by a Portuguese multidisciplinary team on primary ciliary dyskinesia. <i>ERJ Open Research</i> , 2021, 7, 00792-2020.	2.6	15
4	Pkd2 Affects Cilia Length and Impacts LR Flow Dynamics and Dand5. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 624531.	3.7	5
5	Primary ciliary dyskinesia due to CCNO mutationsâ€”A genotypeâ€”phenotype correlation contribution. <i>Pediatric Pulmonology</i> , 2021, 56, 2776-2779.	2.0	4
6	Nutritional and toxicity profiles of two species of land snail, <i>Theba pisana</i> and <i>Otala lactea</i> , from Morocco. <i>Journal of Food Composition and Analysis</i> , 2021, 100, 103893.	3.9	4
7	Zebrafish Model as a Screen to Prevent Cyst Inflation in Autosomal Dominant Polycystic Kidney Disease. <i>International Journal of Molecular Sciences</i> , 2021, 22, 9013.	4.1	1
8	Zebrafish Motile Cilia as a Model for Primary Ciliary Dyskinesia. <i>International Journal of Molecular Sciences</i> , 2021, 22, 8361.	4.1	8
9	Editorial: The Cytoskeleton and Cellular Compartmentation: Cilia as Specialized Cellular Domains. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 777758.	3.7	0
10	Clinical utility of NGS diagnosis and disease stratification in a multiethnic primary ciliary dyskinesia cohort. <i>Journal of Medical Genetics</i> , 2020, 57, 322-330.	3.2	50
11	The Zebrafish Kupfferâ€™s Vesicle: A Special Organ in a Model Organism to Study Human Diseases. , 2020, , .		2
12	Unmasking the relevance of hemispheric asymmetriesâ€”Break on through (to the other side). <i>Progress in Neurobiology</i> , 2020, 192, 101823.	5.7	29
13	Symmetry-Breaking Cilia-Driven Flow in Embryogenesis. <i>Annual Review of Fluid Mechanics</i> , 2019, 51, 105-128.	25.0	31
14	Rab35 controls cilium length, function and membrane composition. <i>EMBO Reports</i> , 2019, 20, e47625.	4.5	35
15	Wall stress enhanced exocytosis of extracellular vesicles as a possible mechanism of left-right symmetry-breaking in vertebrate development. <i>Journal of Theoretical Biology</i> , 2019, 460, 220-226.	1.7	7
16	Imbalanced mitochondrial function provokes heterotaxy via aberrant ciliogenesis. <i>Journal of Clinical Investigation</i> , 2019, 129, 2841-2855.	8.2	43
17	Usefulness of zebrafish larvae to evaluate drug-induced functional and morphological renal tubular alterations. <i>Archives of Toxicology</i> , 2018, 92, 411-423.	4.2	39
18	Zebrafish Larvae Are a Suitable Model to Investigate the Metabolic Phenotype of Drug-Induced Renal Tubular Injury. <i>Frontiers in Pharmacology</i> , 2018, 9, 1193.	3.5	13

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19	Dynamics of cilia length in left-right development. Royal Society Open Science, 2017, 4, 161102.	2.4	19
20	Notch/Her12 signalling modulates, motile/immotile cilia ratio downstream of Foxj1a in zebrafish left-right organizer. ELife, 2017, 6, .	6.0	26
21	SPO22THE CROSSTALK BETWEEN POLYCYSTIN-2 AND CFTR IN AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE. Nephrology Dialysis Transplantation, 2016, 31, i94-i94.	0.7	0
22	Three-dimensional flow in Kupffer's Vesicle. Journal of Mathematical Biology, 2016, 73, 705-725.	1.9	18
23	The zebrafish Kupffer's vesicle as a model system for the molecular mechanisms by which the lack of Polycystin-2 leads to stimulation of CFTR. Biology Open, 2015, 4, 1356-1366.	1.2	24
24	Arl13b interferes with β -tubulin acetylation. Cilia, 2015, 4, .	1.8	2
25	Early steps in primary cilium assembly require EHD1/EHD3-dependent ciliary vesicle formation. Nature Cell Biology, 2015, 17, 228-240.	10.3	221
26	Organized chaos in Kupffer's vesicle: How a heterogeneous structure achieves consistent left-right patterning. Bioarchitecture, 2014, 4, 119-125.	1.5	22
27	Paraoxonase as part of endogenous free-radical scavenging system in zebrafish. Toxicology Letters, 2014, 229, S41.	0.8	0
28	Arl13b and the non-muscle myosin heavy chain IIA are required for circular dorsal ruffle formation and cell migration. Journal of Cell Science, 2014, 127, 2709-22.	2.0	33
29	Left-Right Organizer Flow Dynamics: How Much Cilia Activity Reliably Yields Laterality?. Developmental Cell, 2014, 29, 716-728.	7.0	85
30	The Importance of Zebrafish in Biomedical Research. Acta Medica Portuguesa, 2013, 26, 583-592.	0.4	56
31	The importance of Zebrafish in biomedical research. Acta Medica Portuguesa, 2013, 26, 583-92.	0.4	36
32	Notch signalling regulates left-right asymmetry through ciliary length control. Development (Cambridge), 2010, 137, 3625-3632.	2.5	107
33	Left-Right Function of dmrt2 Genes Is Not Conserved between Zebrafish and Mouse. PLoS ONE, 2010, 5, e14438.	2.5	39
34	Dll1 and Dll4 function sequentially in the retina and pV2 domain of the spinal cord to regulate neurogenesis and create cell diversity. Developmental Biology, 2009, 328, 54-65.	2.0	63
35	16-P010 A novel role for notch signalling in left-right determination through ciliary length control. Mechanisms of Development, 2009, 126, S265.	1.7	0
36	Leukocyte Tyrosine Kinase Functions in Pigment Cell Development. PLoS Genetics, 2008, 4, e1000026.	3.5	137

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37	Zebrafish <i>colourless</i> encodes <i>sox10</i> and specifies non-ectomesenchymal neural crest fates. <i>Development (Cambridge)</i> , 2001, 128, 4113-4125.	2.5	449
38	Zebrafish <i>colourless</i> encodes <i>sox10</i> and specifies non-ectomesenchymal neural crest fates. <i>Development (Cambridge)</i> , 2001, 128, 4113-25.	2.5	218
39	Mutational Analysis of Endothelin Receptor b1 (<i>rose</i>) during Neural Crest and Pigment Pattern Development in the Zebrafish <i>Danio rerio</i> . <i>Developmental Biology</i> , 2000, 227, 294-306.	2.0	209