

Martin Lowe

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/6860661/martin-lowe-publications-by-year.pdf>

Version: 2024-04-29

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

45
papers

1,967
citations

25
h-index

44
g-index

106
ext. papers

2,429
ext. citations

7.4
avg, IF

5.16
L-index

#	Paper	IF	Citations
45	Supply chain logistics - the role of the Golgi complex in extracellular matrix production and maintenance.. <i>Journal of Cell Science</i> , 2022 , 135,	5.3	2
44	Basement membrane defects in CD151-associated glomerular disease.. <i>Pediatric Nephrology</i> , 2022 , 1	3.2	1
43	SdhA blocks disruption of the Legionella-containing vacuole by hijacking the OCRL phosphatase. <i>Cell Reports</i> , 2021 , 37, 109894	10.6	2
42	Membrane trafficking in health and disease. <i>DMM Disease Models and Mechanisms</i> , 2020 , 13,	4.1	25
41	Cell-Based Phenotypic Drug Screening Identifies Luteolin as Candidate Therapeutic for Nephropathic Cystinosis. <i>Journal of the American Society of Nephrology: JASN</i> , 2020 , 31, 1522-1537	12.7	5
40	The NLRP3-inflammasome as a sensor of organelle dysfunction. <i>Journal of Cell Biology</i> , 2020 , 219,	7.3	24
39	A role for OCRL in glomerular function and disease. <i>Pediatric Nephrology</i> , 2020 , 35, 641-648	3.2	4
38	PTEN reduces endosomal PtdIns(4,5)P in a phosphatase-independent manner via a PLC pathway. <i>Journal of Cell Biology</i> , 2019 , 218, 2198-2214	7.3	6
37	IPIP27 Coordinates PtdIns(4,5)P Homeostasis for Successful Cytokinesis. <i>Current Biology</i> , 2019 , 29, 775-789.e7	8.9	8
36	The Physiological Functions of the Golgin Vesicle Tethering Proteins. <i>Frontiers in Cell and Developmental Biology</i> , 2019 , 7, 94	5.7	22
35	Hypomorphic mutations of TRIP11 cause odontochondrodysplasia. <i>JCI Insight</i> , 2019 , 4,	9.9	18
34	GORAB scaffolds COPI at the trans-Golgi for efficient enzyme recycling and correct protein glycosylation. <i>Nature Communications</i> , 2019 , 10, 127	17.4	27
33	A common pathomechanism in GMAP-210- and LBR-related diseases. <i>JCI Insight</i> , 2018 , 3,	9.9	4
32	Impaired proteoglycan glycosylation, elevated TGF- β signaling, and abnormal osteoblast differentiation as the basis for bone fragility in a mouse model for geroderma osteodysplastica. <i>PLoS Genetics</i> , 2018 , 14, e1007242	6	25
31	Identification and functional analysis of a novel oculocerebrorenal syndrome of Lowe () gene variant in two pedigrees with varying phenotypes including isolated congenital cataract. <i>Molecular Vision</i> , 2018 , 24, 847-852	2.3	3
30	Genetic Renal Diseases: The Emerging Role of Zebrafish Models. <i>Cells</i> , 2018 , 7,	7.9	23
29	Cystinosis (ctns) zebrafish mutant shows pronephric glomerular and tubular dysfunction. <i>Scientific Reports</i> , 2017 , 7, 42583	4.9	23

28	Recognition and tethering of transport vesicles at the Golgi apparatus. <i>Current Opinion in Cell Biology</i> , 2017 , 47, 16-23	9	34
27	Loss of the golgin GM130 causes Golgi disruption, Purkinje neuron loss, and ataxia in mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, 346-351	11.5	66
26	OCRL1 engages with the F-BAR protein pacsin 2 to promote biogenesis of membrane-trafficking intermediates. <i>Molecular Biology of the Cell</i> , 2016 , 27, 90-107	3.5	21
25	Role of reverse phenotyping in interpretation of next generation sequencing data and a review of INPP5E related disorders. <i>European Journal of Paediatric Neurology</i> , 2016 , 20, 286-295	3.8	25
24	Autophagosome-lysosome fusion triggers a lysosomal response mediated by TLR9 and controlled by OCRL. <i>Nature Cell Biology</i> , 2016 , 18, 839-850	23.4	103
23	The golgin GMAP-210 is required for efficient membrane trafficking in the early secretory pathway. <i>Journal of Cell Science</i> , 2015 , 128, 1595-606	5.3	32
22	Coupling of vesicle tethering and Rab binding is required for in vivo functionality of the golgin GMAP-210. <i>Molecular Biology of the Cell</i> , 2015 , 26, 537-53	3.5	36
21	Lamellipodin Is Important for Cell-to-Cell Spread and Actin-Based Motility in <i>Listeria monocytogenes</i> . <i>Infection and Immunity</i> , 2015 , 83, 3740-8	3.7	11
20	The Lowe syndrome protein OCRL1 is required for endocytosis in the zebrafish pronephric tubule. <i>PLoS Genetics</i> , 2015 , 11, e1005058	6	50
19	The Golgin Family of Coiled-Coil Tethering Proteins. <i>Frontiers in Cell and Developmental Biology</i> , 2015 , 3, 86	5.7	62
18	The cellular and physiological functions of the Lowe syndrome protein OCRL1. <i>Traffic</i> , 2014 , 15, 471-87	5.7	76
17	Inositol lipid phosphatases in membrane trafficking and human disease. <i>Biochemical Journal</i> , 2014 , 461, 159-75	3.8	49
16	The Lowe syndrome protein OCRL1 is involved in primary cilia assembly. <i>Human Molecular Genetics</i> , 2012 , 21, 1835-47	5.6	71
15	Impaired neural development in a zebrafish model for Lowe syndrome. <i>Human Molecular Genetics</i> , 2012 , 21, 1744-59	5.6	55
14	Lowe Syndrome protein OCRL1 supports maturation of polarized epithelial cells. <i>PLoS ONE</i> , 2011 , 6, e24044	3.7	18
13	Structural organization of the Golgi apparatus. <i>Current Opinion in Cell Biology</i> , 2011 , 23, 85-93	9	95
12	The PH domain proteins IPIP27A and B link OCRL1 to receptor recycling in the endocytic pathway. <i>Molecular Biology of the Cell</i> , 2011 , 22, 606-23	3.5	52
11	Lowe syndrome patient fibroblasts display Ocr1-specific cell migration defects that cannot be rescued by the homologous Inpp5b phosphatase. <i>Human Molecular Genetics</i> , 2009 , 18, 4478-91	5.6	46

10	Differential clathrin binding and subcellular localization of OCRL1 splice isoforms. <i>Journal of Biological Chemistry</i> , 2009 , 284, 9965-73	5-4	56
9	Inheritance and biogenesis of organelles in the secretory pathway. <i>Nature Reviews Molecular Cell Biology</i> , 2007 , 8, 429-39	48-7	91
8	Dissecting the role of the ARF guanine nucleotide exchange factor GBF1 in Golgi biogenesis and protein trafficking. <i>Journal of Cell Science</i> , 2007 , 120, 3929-40	5-3	83
7	Membrane targeting and activation of the Lowe syndrome protein OCRL1 by rab GTPases. <i>EMBO Journal</i> , 2006 , 25, 3750-61	13	126
6	Structure and function of the Lowe syndrome protein OCRL1. <i>Traffic</i> , 2005 , 6, 711-9	5-7	125
5	Phosphoinositides and membrane traffic at the trans-Golgi network. <i>Biochemical Society Symposia</i> , 2005 , 72, 31-8		12
4	Lowe syndrome protein OCRL1 interacts with clathrin and regulates protein trafficking between endosomes and the trans-Golgi network. <i>Molecular Biology of the Cell</i> , 2005 , 16, 3467-79	3-5	152
3	The coiled-coil membrane protein golgin-84 is a novel rab effector required for Golgi ribbon formation. <i>Journal of Cell Biology</i> , 2003 , 160, 201-12	7-3	193
2	A novel nanoluciferase transgenic reporter to measure proteinuria in zebrafish		3
1	NLRP3 activation in response to disrupted endocytic traffic		1