

# Roman S Polishchuk

## List of Publications by Year in descending order

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117  
papers

12,514  
citations

30070

54  
h-index

25787

108  
g-index

122  
all docs

122  
docs citations

122  
times ranked

16801  
citing authors

#	ARTICLE	IF	CITATIONS
1	A Gene Network Regulating Lysosomal Biogenesis and Function. <i>Science</i> , 2009, 325, 473-477.	12.6	1,958
2	Transcriptional Activation of Lysosomal Exocytosis Promotes Cellular Clearance. <i>Developmental Cell</i> , 2011, 21, 421-430.	7.0	594
3	Connecting copper and cancer: from transition metal signalling to metalloplasia. <i>Nature Reviews Cancer</i> , 2022, 22, 102-113.	28.4	519
4	Rapid Cycling of Lipid Raft Markers between the Cell Surface and Golgi Complex. <i>Journal of Cell Biology</i> , 2001, 153, 529-542.	5.2	496
5	Maintenance of Golgi structure and function depends on the integrity of ER export. <i>Journal of Cell Biology</i> , 2001, 155, 557-570.	5.2	398
6	Glycosphingolipid synthesis requires FAPP2 transfer of glucosylceramide. <i>Nature</i> , 2007, 449, 62-67.	27.8	359
7	Golgi Membranes Are Absorbed into and Reemerge from the ER during Mitosis. <i>Cell</i> , 1999, 99, 589-601.	28.9	315
8	Correlative Light-Electron Microscopy Reveals the Tubular-Saccular Ultrastructure of Carriers Operating between Golgi Apparatus and Plasma Membrane. <i>Journal of Cell Biology</i> , 2000, 148, 45-58.	5.2	304
9	Secretory traffic triggers the formation of tubular continuities across Golgi sub-compartments. <i>Nature Cell Biology</i> , 2004, 6, 1071-1081.	10.3	283
10	Transcription factor EB (TFEB) is a new therapeutic target for Pompe disease. <i>EMBO Molecular Medicine</i> , 2013, 5, 691-706.	6.9	273
11	A tubular EHD1-containing compartment involved in the recycling of major histocompatibility complex class I molecules to the plasma membrane. <i>EMBO Journal</i> , 2002, 21, 2557-2567.	7.8	265
12	Transport through the Golgi Apparatus by Rapid Partitioning within a Two-Phase Membrane System. <i>Cell</i> , 2008, 133, 1055-1067.	28.9	256
13	Mitochondria Are Linked to Calcium Stores in Striated Muscle by Developmentally Regulated Tethering Structures. <i>Molecular Biology of the Cell</i> , 2009, 20, 1058-1067.	2.1	240
14	Wilson Disease Protein ATP7B Utilizes Lysosomal Exocytosis to Maintain Copper Homeostasis. <i>Developmental Cell</i> , 2014, 29, 686-700.	7.0	203
15	Effective delivery of large genes to the retina by dual AAV vectors. <i>EMBO Molecular Medicine</i> , 2014, 6, 194-211.	6.9	202
16	Delivery of raft-associated, GPI-anchored proteins to the apical surface of polarized MDCK cells by a transcytotic pathway. <i>Nature Cell Biology</i> , 2004, 6, 297-307.	10.3	192
17	Lysosomal fusion and SNARE function are impaired by cholesterol accumulation in lysosomal storage disorders. <i>EMBO Journal</i> , 2010, 29, 3607-3620.	7.8	192
18	Small cargo proteins and large aggregates can traverse the Golgi by a common mechanism without leaving the lumen of cisternae. <i>Journal of Cell Biology</i> , 2001, 155, 1225-1238.	5.2	185

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19	Mechanism of Constitutive Export from the Golgi: Bulk Flow via the Formation, Protrusion, and En Bloc Cleavage of large trans-Golgi Network Tubular Domains. <i>Molecular Biology of the Cell</i> , 2003, 14, 4470-4485.	2.1	177
20	A traffic-activated Golgi-based signalling circuit coordinates the secretory pathway. <i>Nature Cell Biology</i> , 2008, 10, 912-922.	10.3	175
21	FGF signalling regulates bone growth through autophagy. <i>Nature</i> , 2015, 528, 272-275.	27.8	170
22	OCRL controls trafficking through early endosomes via PtdIns4,5P <sub>2</sub> -dependent regulation of endosomal actin. <i>EMBO Journal</i> , 2011, 30, 4970-4985.	7.8	158
23	Activation of Autophagy, Observed in Liver Tissues From Patients With Wilson Disease and From ATP7B-Deficient Animals, Protects Hepatocytes From Copper-Induced Apoptosis. <i>Gastroenterology</i> , 2019, 156, 1173-1189.e5.	1.3	150
24	Gene transfer of master autophagy regulator TFEB results in clearance of toxic protein and correction of hepatic disease in alpha <sub>1</sub> -antitrypsin deficiency. <i>EMBO Molecular Medicine</i> , 2013, 5, 397-412.	6.9	134
25	The Golgi-associated Protein GRASP65 Regulates Spindle Dynamics and Is Essential for Cell Division. <i>Molecular Biology of the Cell</i> , 2005, 16, 3211-3222.	2.1	126
26	Dimeric PKD regulates membrane fission to form transport carriers at the TGN. <i>Journal of Cell Biology</i> , 2007, 179, 1123-1131.	5.2	121
27	A role for Arf1 in mitotic Golgi disassembly, chromosome segregation, and cytokinesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 13314-13319.	7.1	110
28	Intein-mediated protein trans-splicing expands adeno-associated virus transfer capacity in the retina. <i>Science Translational Medicine</i> , 2019, 11, .	12.4	109
29	COPI acts in both vesicular and tubular transport. <i>Nature Cell Biology</i> , 2011, 13, 996-1003.	10.3	108
30	Actin remodeling by ADF/cofilin is required for cargo sorting at the trans-Golgi network. <i>Journal of Cell Biology</i> , 2009, 187, 1055-1069.	5.2	98
31	Golgi in copper homeostasis: a view from the membrane trafficking field. <i>Histochemistry and Cell Biology</i> , 2013, 140, 285-295.	1.7	97
32	Polycystin-1 Induces Cell Migration by Regulating Phosphatidylinositol 3-kinase-dependent Cytoskeletal Rearrangements and GSK3 $\beta$ -dependent Cellâ€”Cell Mechanical Adhesion. <i>Molecular Biology of the Cell</i> , 2007, 18, 4050-4061.	2.1	96
33	Molecular determinants of ERâ€”Golgi contacts identified through a new FRETâ€”FLIM system. <i>Journal of Cell Biology</i> , 2019, 218, 1055-1065.	5.2	94
34	Defective Intracellular Trafficking of Uromodulin Mutant Isoforms. <i>Traffic</i> , 2006, 7, 1567-1579.	2.7	93
35	Golgi Enzymes Are Enriched in Perforated Zones of Golgi Cisternae but Are Depleted in COPI Vesicles. <i>Molecular Biology of the Cell</i> , 2004, 15, 4710-4724.	2.1	90
36	A new class of carriers that transport selective cargo from the trans Golgi network to the cell surface. <i>EMBO Journal</i> , 2012, 31, 3976-3990.	7.8	88

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37	The role of NSP6 in the biogenesis of the SARS-CoV-2 replication organelle. <i>Nature</i> , 2022, 606, 761-768.	27.8	87
38	Activation of the transcription factor EB rescues lysosomal abnormalities in cystinotic kidney cells. <i>Kidney International</i> , 2016, 89, 862-873.	5.2	85
39	Group IV Phospholipase A2 $\pm$ Controls the Formation of Inter-Cisternal Continuities Involved in Intra-Golgi Transport. <i>PLoS Biology</i> , 2009, 7, e1000194.	5.6	81
40	A 14-3-3 $\beta$ dimer-based scaffold bridges CtBP1-S/BARS to PI(4)KIII $\beta$ to regulate post-Golgi carrier formation. <i>Nature Cell Biology</i> , 2012, 14, 343-354.	10.3	79
41	The Role of GRASP55 in Golgi Fragmentation and Entry of Cells into Mitosis. <i>Molecular Biology of the Cell</i> , 2008, 19, 2579-2587.	2.1	78
42	The E3-Ubiquitin Ligase TRIM50 Interacts with HDAC6 and p62, and Promotes the Sequestration and Clearance of Ubiquitinated Proteins into the Aggresome. <i>PLoS ONE</i> , 2012, 7, e40440.	2.5	76
43	T-Cell Antigen Receptor-Induced Signaling Complexes: Internalization Via a Cholesterol-Dependent Endocytic Pathway. <i>Traffic</i> , 2006, 7, 1143-1162.	2.7	74
44	Sphingomyelin organization is required for vesicle biogenesis at the Golgi complex. <i>EMBO Journal</i> , 2012, 31, 4535-4546.	7.8	74
45	Defective autophagy in spastizin mutated patients with hereditary spastic paraparesis type 15. <i>Brain</i> , 2013, 136, 3119-3139.	7.6	74
46	Transport of soluble proteins through the Golgi occurs by diffusion via continuities across cisternae. <i>ELife</i> , 2014, 3, .	6.0	74
47	Faciogenital Dysplasia Protein Fgd1 Regulates Invadopodia Biogenesis and Extracellular Matrix Degradation and Is Up-regulated in Prostate and Breast Cancer. <i>Cancer Research</i> , 2009, 69, 747-752.	0.9	73
48	Improved dual AAV vectors with reduced expression of truncated proteins are safe and effective in the retina of a mouse model of Stargardt disease. <i>Human Molecular Genetics</i> , 2015, 24, 6811-6825.	2.9	73
49	Metformin restores the mitochondrial network and reverses mitochondrial dysfunction in Down syndrome cells. <i>Human Molecular Genetics</i> , 2017, 26, ddx016.	2.9	70
50	NCX3 regulates mitochondrial calcium handling through AKAP121-anchored signaling complex and prevents hypoxia-induced cell death. <i>Journal of Cell Science</i> , 2013, 126, 5566-77.	2.0	64
51	The emerging role of lysosomes in copper homeostasis. <i>Metallomics</i> , 2016, 8, 853-862.	2.4	64
52	Golgi membrane fission requires the CtBP1-S/BARS-induced activation of lysophosphatidic acid acyltransferase $\beta$ . <i>Nature Communications</i> , 2016, 7, 12148.	12.8	63
53	Ultrastructure of Long-Range Transport Carriers Moving from the trans Golgi Network to Peripheral Endosomes. <i>Traffic</i> , 2006, 7, 1092-1103.	2.7	62
54	Connexin-Mediated Signaling in Nonsensory Cells Is Crucial for the Development of Sensory Inner Hair Cells in the Mouse Cochlea. <i>Journal of Neuroscience</i> , 2017, 37, 258-268.	3.6	61

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55	Visualizing membrane traffic in vivo by combined video fluorescence and 3D electron microscopy. <i>Trends in Cell Biology</i> , 2000, 10, 349-353.	7.9	60
56	The Menkes and Wilson disease genes counteract in copper toxicosis in Labrador retrievers: a new canine model for copper-metabolism disorders. <i>DMM Disease Models and Mechanisms</i> , 2016, 9, 25-38.	2.4	60
57	Activity and Trafficking of Copper-Transporting ATPases in Tumor Development and Defense against Platinum-Based Drugs. <i>Cells</i> , 2019, 8, 1080.	4.1	58
58	Morphogenesis of post-Golgi transport carriers. <i>Histochemistry and Cell Biology</i> , 2008, 129, 153-161.	1.7	57
59	Molecular Events Initiating Exit of a Copper-transporting ATPase ATP7B from the Trans-Golgi Network. <i>Journal of Biological Chemistry</i> , 2012, 287, 36041-36050.	3.4	53
60	Faciogenital Dysplasia Protein (FGD1) Regulates Export of Cargo Proteins from the Golgi Complex via Cdc42 Activation. <i>Molecular Biology of the Cell</i> , 2009, 20, 2413-2427.	2.1	52
61	Uromodulin is expressed in renal primary cilia and UMOD mutations result in decreased ciliary uromodulin expression. <i>Human Molecular Genetics</i> , 2010, 19, 1985-1997.	2.9	52
62	Role of NAD <sup>+</sup> and ADP-Ribosylation in the Maintenance of the Golgi Structure. <i>Journal of Cell Biology</i> , 1997, 139, 1109-1118.	5.2	50
63	A High-Calorie Diet Aggravates Mitochondrial Dysfunction and Triggers Severe Liver Damage in Wilson Disease Rats. <i>Cellular and Molecular Gastroenterology and Hepatology</i> , 2019, 7, 571-596.	4.5	50
64	Hypothyroidism induced by loss of the manganese efflux transporter SLC30A10 may be explained by reduced thyroxine production. <i>Journal of Biological Chemistry</i> , 2017, 292, 16605-16615.	3.4	46
65	D-Aspartate treatment attenuates myelin damage and stimulates myelin repair. <i>EMBO Molecular Medicine</i> , 2019, 11, .	6.9	44
66	Coalescence of Golgi fragments in microtubule-deprived living cells. <i>European Journal of Cell Biology</i> , 1999, 78, 170-185.	3.6	43
67	Dicumarol, an inhibitor of ADP-ribosylation of CtBP3/BARS, fragments Golgi non-compact tubular zones and inhibits intra-Golgi transport. <i>European Journal of Cell Biology</i> , 2004, 83, 263-279.	3.6	43
68	Correlation of 4Pi and Electron Microscopy to Study Transport Through Single Golgi Stacks in Living Cells with Super Resolution. <i>Traffic</i> , 2009, 10, 379-391.	2.7	43
69	Correction of CNS defects in the MPSII mouse model via systemic enzyme replacement therapy. <i>Human Molecular Genetics</i> , 2010, 19, 4871-4885.	2.9	43
70	Identification of p38 MAPK and JNK as new targets for correction of Wilson disease-causing ATP7B mutants. <i>Hepatology</i> , 2016, 63, 1842-1859.	7.3	42
71	Localization and age-dependent expression of the inward rectifier K <sup>+</sup> channel subunit Kir 5.1 in a mammalian reproductive system. <i>FEBS Letters</i> , 1999, 449, 146-152.	2.8	41
72	Intra-Golgi transport: A way to a new paradigm?. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2005, 1744, 340-350.	4.1	40

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73	Golgi-modifying properties of macfarlandin E and the synthesis and evaluation of its 2,7-dioxabicyclo[3.2.1]octan-3-one core. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 6158-6163.	7.1	40
74	AAV-mediated transcription factor EB (TFEB) gene delivery ameliorates muscle pathology and function in the murine model of Pompe Disease. Scientific Reports, 2017, 7, 15089.	3.3	40
75	A novel murine model for arrhythmogenic cardiomyopathy points to a pathogenic role of Wnt signalling and miRNA dysregulation. Cardiovascular Research, 2019, 115, 739-751.	3.8	40
76	Akap1 Deficiency Promotes Mitochondrial Aberrations and Exacerbates Cardiac Injury Following Permanent Coronary Ligation via Enhanced Mitophagy and Apoptosis. PLoS ONE, 2016, 11, e0154076.	2.5	39
77	PERK-Mediated Unfolded Protein Response Activation and Oxidative Stress in PARK20 Fibroblasts. Frontiers in Neuroscience, 2019, 13, 673.	2.8	38
78	Silver Ions as a Tool for Understanding Different Aspects of Copper Metabolism. Nutrients, 2019, 11, 1364.	4.1	38
79	Pharmacological read-through of nonsense ARSB mutations as a potential therapeutic approach for mucopolysaccharidosis VI. Journal of Inherited Metabolic Disease, 2013, 36, 363-371.	3.6	36
80	Rab6 and myosin II at the cutting edge of membrane fission. Nature Cell Biology, 2010, 12, 635-638.	10.3	35
81	Characterization of the most frequent ATP7B mutation causing Wilson disease in hepatocytes from patient induced pluripotent stem cells. Scientific Reports, 2018, 8, 6247.	3.3	35
82	Atherosclerosis-prone branch regions in human aorta: microarchitecture and cell composition of intima. Atherosclerosis, 1996, 122, 173-189.	0.8	31
83	The cytosolic chaperone $\beta$ -Crystallin B rescues appropriate folding and compartmentalization of misfolded multispan transmembrane proteins. Journal of Cell Science, 2013, 126, 4160-72.	2.0	31
84	Myosin7a Deficiency Results in Reduced Retinal Activity Which Is Improved by Gene Therapy. PLoS ONE, 2013, 8, e72027.	2.5	29
85	Cytosolic phospholipase A2 $\mu$ drives recycling in the clathrin-independent endocytic route. Journal of Cell Science, 2014, 127, 977-93.	2.0	26
86	CREB3L1-mediated functional and structural adaptation of the secretory pathway in hormone-stimulated thyroid cells. Journal of Cell Science, 2017, 130, 4155-4167.	2.0	26
87	Synthetic Lethality Screening Identifies FDA-Approved Drugs that Overcome ATP7B-Mediated Tolerance of Tumor Cells to Cisplatin. Cancers, 2020, 12, 608.	3.7	25
88	Changes in Muscle Cell Metabolism and Mechanotransduction Are Associated with Myopathic Phenotype in a Mouse Model of Collagen VI Deficiency. PLoS ONE, 2013, 8, e56716.	2.5	23
89	Shaping tubular carriers for intracellular membrane transport. FEBS Letters, 2009, 583, 3847-3856.	2.8	22
90	Pre-embedding labeling for subcellular detection of molecules with electron microscopy. Tissue and Cell, 2019, 57, 103-110.	2.2	22

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91	Visualizing Live Dynamics and Ultrastructure of Intracellular Organelles with Preembedding Correlative Light-Electron Microscopy. <i>Methods in Cell Biology</i> , 2012, 111, 21-35.	1.1	21
92	Helical organization of microtubules occurs in a minority of tunneling membrane nanotubes in normal and cancer urothelial cells. <i>Scientific Reports</i> , 2018, 8, 17133.	3.3	21
93	A systems biology approach reveals new endoplasmic reticulum-associated targets for the correction of the ATP7B mutant causing Wilson disease. <i>Metallomics</i> , 2016, 8, 920-930.	2.4	19
94	Abnormal cell-clearance and accumulation of autophagic vesicles in lymphocytes from patients affected with Ataxia-Teleangiectasia. <i>Clinical Immunology</i> , 2017, 175, 16-25.	3.2	19
95	Uroplakin traffic through the Golgi apparatus induces its fragmentation: new insights from novel in vitro models. <i>Scientific Reports</i> , 2017, 7, 12842.	3.3	19
96	Mitochondrial Dynamics of Proximal Tubular Epithelial Cells in Nephropathic Cystinosis. <i>International Journal of Molecular Sciences</i> , 2020, 21, 192.	4.1	19
97	Correlative Light-Electron Microscopy as a Tool to Study In Vivo Dynamics and Ultrastructure of Intracellular Structures. <i>Methods in Molecular Biology</i> , 2012, 931, 413-422.	0.9	18
98	Golgi-Dependent Copper Homeostasis Sustains Synaptic Development and Mitochondrial Content. <i>Journal of Neuroscience</i> , 2021, 41, 215-233.	3.6	17
99	Chapter 5 Combined Video Fluorescence and 3D Electron Microscopy. <i>Methods in Cell Biology</i> , 2008, 88, 83-95.	1.1	16
100	From and to the Golgi – defining the Wilson disease protein road map. <i>FEBS Letters</i> , 2019, 593, 2341-2350.	2.8	16
101	Correlative Video Light/Electron Microscopy. <i>Current Protocols in Cell Biology</i> , 2001, 11, Unit 4.8.	2.3	13
102	Correction of oxidative stress enhances enzyme replacement therapy in Pompe disease. <i>EMBO Molecular Medicine</i> , 2021, 13, e14434.	6.9	13
103	Visualizing Intracellular Events In Vivo by Combined Video Fluorescence and 3D Electron Microscopy. <i>Methods in Enzymology</i> , 2005, 404, 43-57.	1.0	12
104	Myopalladin knockout mice develop cardiac dilation and show a maladaptive response to mechanical pressure overload. <i>ELife</i> , 2021, 10, .	6.0	12
105	Analysis of Golgi Complex Function Using Correlative Light-Electron Microscopy. <i>Methods in Cell Biology</i> , 2013, 118, 243-258.	1.1	11
106	Emerging role of Cdc42-specific guanine nucleotide exchange factors as regulators of membrane trafficking in health and disease. <i>Tissue and Cell</i> , 2017, 49, 157-162.	2.2	11
107	ER/Golgi trafficking is facilitated by unbranched actin filaments containing Tpm4.2. <i>Cytoskeleton</i> , 2017, 74, 379-389.	2.0	11
108	TFEB Regulates ATP7B Expression to Promote Platinum Chemoresistance in Human Ovarian Cancer Cells. <i>Cells</i> , 2022, 11, 219.	4.1	10

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109	Cystinosin-LKG rescues cystine accumulation and decreases apoptosis rate in cystinotic proximal tubular epithelial cells. <i>Pediatric Research</i> , 2017, 81, 113-119.	2.3	9
110	Pharmacoproteomics pinpoints HSP70 interaction for correction of the most frequent Wilson disease-causing mutant of ATP7B. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 32453-32463.	7.1	9
111	Skin fibroblasts of patients with geleophysic dysplasia due to <i>FBN1</i> mutations have lysosomal inclusions and losartan improves their microfibril deposition defect. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e844.	1.2	8
112	Identification of CDC42 Effectors Operating in FGD1-Dependent Trafficking at the Golgi. <i>Frontiers in Cell and Developmental Biology</i> , 2019, 7, 7.	3.7	8
113	Cellular Function of ATP7B (Wilson ATPase). , 2019, , 45-56.		2
114	Protein transport from the trans-Golgi network to endosomes. , 2008, , 388-401.		1
115	Down Syndrome Fetal Fibroblasts Display Alterations of Endosomal Trafficking Possibly due to SYNJ1 Overexpression. <i>Frontiers in Genetics</i> , 2022, 13, .	2.3	1
116	Toxicological Assessment Via Gene Network Analysis. <i>Methods in Pharmacology and Toxicology</i> , 2015, , 161-180.	0.2	0
117	Golgi-to-PM transport. , 2008, , 375-387.		0