Carl P Blobel

List of Publications by Year
in descending order

112 papers	13,911 citations	20817 60 h-index	26613 107 g-index
114 all docs	114 docs citations	114 times ranked	11352 citing authors

#	Article	IF	Citations
1	ADAMs and ADAMTSs. , 2022, , 568-573.		3
2	ADAMs Regulate Cell-Cell Interactions by Controlling the Function of the EGF-Receptor, TNFÎ± and Notch. , 2022, , .		0
3	ADAM10 and ADAM17 promote SARSâ€CoVâ€2 cell entry and spike proteinâ€mediated lung cell fusion. EMBO Reports, 2022, 23, e54305.	4.5	57
4	Targeted truncation of the ADAM17 cytoplasmic domain in mice results in protein destabilization and a hypomorphic phenotype. Journal of Biological Chemistry, 2021, 296, 100733.	3.4	9
5	Analysis of the Conditions That Affect the Selective Processing of Endogenous Notch1 by ADAM10 and ADAM17. International Journal of Molecular Sciences, 2021, 22, 1846.	4.1	10
6	The pseudoprotease iRhom1 controls ectodomain shedding of membrane proteins in the nervous system. FASEB Journal, 2021, 35, e21962.	0.5	5
7	Role of iRhoms 1 and 2 in Endochondral Ossification. International Journal of Molecular Sciences, 2020, 21, 8732.	4.1	4
8	Endothelial deletion of ADAM10, a key regulator of Notch signaling, causes impaired decidualization and reduced fertility in female mice. Angiogenesis, 2020, 23, 443-458.	7.2	15
9	ADAM17 stabilizes its interacting partner inactive Rhomboid 2 (iRhom2) but not inactive Rhomboid 1 (iRhom1). Journal of Biological Chemistry, 2020, 295, 4350-4358.	3.4	12
10	Substrateâ€selective protein ectodomain shedding by ADAM17 and iRhom2 depends on their juxtamembrane and transmembrane domains. FASEB Journal, 2020, 34, 4956-4969.	0.5	22
11	The Threshold Effect: Lipopolysaccharide-Induced Inflammatory Responses in Primary Macrophages Are Differentially Regulated in an iRhom2-Dependent Manner. Frontiers in Cellular and Infection Microbiology, 2020, 10, 620392.	3.9	1
12	ADAM10 controls the differentiation of the coronary arterial endothelium. Angiogenesis, 2019, 22, 237-250.	7.2	36
13	Glomerular endothelial cell maturation depends on ADAM10, a key regulator of Notch signaling. Angiogenesis, 2018, 21, 335-347.	7.2	31
14	Macrocyclic Î¸-defensins suppress tumor necrosis factor-Î± (TNF-Î±) shedding by inhibition of TNF-Î±â€“converting enzyme. Journal of Biological Chemistry, 2018, 293, 2725-2734.	3.4	28
15	The xenoestrogens biphenolâ€A and nonylphenol differentially regulate metalloproteaseâ€mediated shedding of EGFR ligands. Journal of Cellular Physiology, 2018, 233, 2247-2256.	4.1	16
16	Blood-induced bone loss in murine hemophilic arthropathy is prevented by blocking the iRhom2/ADAM17/TNF-Î± pathway. Blood, 2018, 132, 1064-1074.	1.4	38
17	Intriguing Roles for Endothelial ADAM10/Notch Signaling in the Development of Organ-Specific Vascular Beds. Physiological Reviews, 2018, 98, 2025-2061.	28.8	37
18	The metalloprotease ADAM10 (a disintegrin and metalloprotease 10) undergoes rapid, postlysis autocatalytic degradation. FASEB Journal, 2018, 32, 3560-3573.	0.5	26

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19	A protective Langerhans cellâ€“keratinocyte axis that is dysfunctional in photosensitivity. Science Translational Medicine, 2018, 10, .	12.4	48
20	iRhom2 promotes lupus nephritis through TNF-Î± and EGFR signaling. Journal of Clinical Investigation, 2018, 128, 1397-1412.	8.2	66
21	Structural modeling defines transmembrane residues in ADAM17 that are crucial for Rhbdf2/ADAM17-dependent proteolysis. Journal of Cell Science, 2017, 130, 868-878.	2.0	34
22	Characterization of the catalytic properties of the membrane-anchored metalloproteinase ADAM9 in cell-based assays. Biochemical Journal, 2017, 474, 1467-1479.	3.7	16
23	ADAM10-Dependent Signaling Through Notch1 and Notch4 Controls Development of Organ-Specific Vascular Beds. Circulation Research, 2016, 119, 519-531.	4.5	39
24	iRhom2 regulates CSF1R cell surface expression and nonâ€steady state myelopoiesis in mice. European Journal of Immunology, 2016, 46, 2737-2748.	2.9	14
25	Evidence for cadherin-11 cleavage in the synovium and partial characterization of its mechanism. Arthritis Research and Therapy, 2015, 17, 126.	3.5	18
26	The Functional Maturation of A Disintegrin and Metalloproteinase (ADAM) 9, 10, and 17 Requires Processing at a Newly Identified Proprotein Convertase (PC) Cleavage Site. Journal of Biological Chemistry, 2015, 290, 12135-12146.	3.4	59
27	The Cytoplasmic Domain of A Disintegrin and Metalloproteinase 10 (ADAM10) Regulates Its Constitutive Activity but Is Dispensable for Stimulated ADAM10-dependent Shedding. Journal of Biological Chemistry, 2015, 290, 7416-7425.	3.4	34
28	Blood-Induced Arthropathy in Hemophilia: Mechanisms and Heterogeneity. Seminars in Thrombosis and Hemostasis, 2015, 41, 832-837.	2.7	31
29	iRhoms 1 and 2 are essential upstream regulators of ADAM17-dependent EGFR signaling. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 6080-6085.	7.1	121
30	iRhoms in the brain â€“ a new frontier?. Cell Cycle, 2015, 14, 3003-3004.	2.6	10
31	Deletions in the cytoplasmic domain of iRhom1 and iRhom2 promote shedding of the TNF receptor by the protease ADAM17. Science Signaling, 2015, 8, ra109.	3.6	60
32	Lack of ADAM10 in endothelial cells affects osteoclasts at the chondroâ€osseus junction. Journal of Orthopaedic Research, 2014, 32, 224-230.	2.3	10
33	Characterization of Oxygen-Induced Retinopathy in Mice Carrying an Inactivating Point Mutation in the Catalytic Site of ADAM15. Investigative Ophthalmology and Visual Science, 2014, 55, 6774-6782.	3.3	10
34	A Murine Model for Retinopathy of Prematurity Identifies Endothelial Cell Proliferation as a Potential Mechanism for Plus Disease. , 2013, 54, 5294.		21
35	Epidermal ADAM17 Is Dispensable for Notch Activation. Journal of Investigative Dermatology, 2013, 133, 2286-2288.	0.7	24
36	ADAM12 is expressed in the tumour vasculature and mediates ectodomain shedding of several membrane-anchored endothelial proteins. Biochemical Journal, 2013, 452, 97-109.	3.7	48

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37	iRhom2 controls the substrate selectivity of stimulated ADAM17-dependent ectodomain shedding. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 11433-11438.	7.1	138
38	ADAM17 Controls Endochondral Ossification by Regulating Terminal Differentiation of Chondrocytes. Molecular and Cellular Biology, 2013, 33, 3077-3090.	2.3	47
39	iRHOM2 is a critical pathogenic mediator of inflammatory arthritis. Journal of Clinical Investigation, 2013, 123, 928-32.	8.2	129
40	Epidermal ADAM17 maintains the skin barrier by regulating EGFR ligandâ€“dependent terminal keratinocyte differentiation. Journal of Experimental Medicine, 2012, 209, 1105-1119.	8.5	161
41	iRhom2 Regulation of TACE Controls TNF-Mediated Protection Against Listeria and Responses to LPS. Science, 2012, 335, 229-232.	12.6	292
42	Interleukin-1 Stimulates ADAM17 through a Mechanism Independent of its Cytoplasmic Domain or Phosphorylation at Threonine 735. PLoS ONE, 2012, 7, e31600.	2.5	43
43	The disintegrin/metalloproteinase Adam10 is essential for epidermal integrity and Notch-mediated signaling. Development (Cambridge), 2011, 138, 495-505.	2.5	130
44	Deletion of Adam10 in endothelial cells leads to defects in organ-specific vascular structures. Blood, 2011, 118, 1163-1174.	1.4	69
45	Migration of growth factor-stimulated epithelial and endothelial cells depends on EGFR transactivation by ADAM17. Nature Communications, 2011, 2, 229.	12.8	128
46	TNF-Î±â€“Converting Enzyme/A Disintegrin and Metalloproteaseâ”17 Mediates Mechanotransduction in Murine Tracheal Epithelial Cells. American Journal of Respiratory Cell and Molecular Biology, 2011, 45, 376-385.	2.9	24
47	Neuronal Brain-derived Neurotrophic Factor Is Synthesized in Excess, with Levels Regulated by Sortilin-mediated Trafficking and Lysosomal Degradation. Journal of Biological Chemistry, 2011, 286, 29556-29567.	3.4	91
48	Shedding of the Mer Tyrosine Kinase Receptor Is Mediated by ADAM17 Protein through a Pathway Involving Reactive Oxygen Species, Protein Kinase CÎ´, and p38 Mitogen-activated Protein Kinase (MAPK). Journal of Biological Chemistry, 2011, 286, 33335-33344.	3.4	228
49	TACE (ADAM17) inhibits Schwann cell myelination. Nature Neuroscience, 2011, 14, 857-865.	14.8	136
50	3D trumps 2D when studying endothelial cells. Blood, 2010, 115, 5128-5130.	1.4	10
51	ADAM8 is a negative regulator of retinal neovascularization and of the growth of heterotopically injected tumor cells in mice. Journal of Molecular Medicine, 2010, 88, 497-505.	3.9	49
52	Stimulation of Platelet-derived Growth Factor Receptor Î² (PDGFRÎ²) Activates ADAM17 and Promotes Metalloproteinase-dependent Cross-talk between the PDGFRÎ² and Epidermal Growth Factor Receptor (EGFR) Signaling Pathways. Journal of Biological Chemistry, 2010, 285, 25024-25032.	3.4	45
53	Pathological Neovascularization Is Reduced by Inactivation of ADAM17 in Endothelial Cells but Not in Pericytes. Circulation Research, 2010, 106, 932-940.	4.5	132
54	MyD88 signaling in nonhematopoietic cells protects mice against induced colitis by regulating specific EGF receptor ligands. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 19967-19972.	7.1	134

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55	The Disintegrin/Metalloproteinase ADAM10 Is Essential for the Establishment of the Brain Cortex. Journal of Neuroscience, 2010, 30, 4833-4844.	3.6	327
56	ADAM17 is regulated by a rapid and reversible mechanism that controls access to its catalytic site. Journal of Cell Science, 2010, 123, 3913-3922.	2.0	165
57	Ectodomain shedding of EGFR ligands and TNFR1 dictates hepatocyte apoptosis during fulminant hepatitis in mice. Journal of Clinical Investigation, 2010, 120, 2731-2744.	8.2	76
58	Conditional Inactivation of TACE by a Sox9 Promoter Leads to Osteoporosis and Increased Granulopoiesis via Dysregulation of IL-17 and G-CSF. Journal of Immunology, 2009, 182, 2093-2101.	0.8	57
59	Ectodomain Shedding of FLT3 Ligand Is Mediated by TNF-Î± Converting Enzyme. Journal of Immunology, 2009, 182, 7408-7414.	0.8	29
60	Src Stimulates Fibroblast Growth Factor Receptor-2 Shedding by an ADAM15 Splice Variant Linked to Breast Cancer. Cancer Research, 2009, 69, 4573-4576.	0.9	30
61	Shedding of Collagen XVII/BP180 in Skin Depends on Both ADAM10 and ADAM9. Journal of Biological Chemistry, 2009, 284, 23386-23396.	3.4	105
62	ADAM9 Is Involved in Pathological Retinal Neovascularization. Molecular and Cellular Biology, 2009, 29, 2694-2703.	2.3	85
63	ADAMs 10 and 17 Represent Differentially Regulated Components of a General Shedding Machinery for Membrane Proteins Such as Transforming Growth Factor Î±, L-Selectin, and Tumor Necrosis Factor Î±. Molecular Biology of the Cell, 2009, 20, 1785-1794.	2.1	230
64	The role of protease activity in ErbB biology. Experimental Cell Research, 2009, 315, 671-682.	2.6	75
65	Characterization of the catalytic activity of the membrane-anchored metalloproteinase ADAM15 in cell-based assays. Biochemical Journal, 2009, 420, 105-113.	3.7	48
66	ADAM12: a potential target for the treatment of chronic wounds. Journal of Molecular Medicine, 2008, 86, 961-969.	3.9	50
67	ADAM10 Regulates Endothelial Permeability and T-Cell Transmigration by Proteolysis of Vascular Endothelial Cadherin. Circulation Research, 2008, 102, 1192-1201.	4.5	264
68	VEGF-A Stimulates ADAM17-Dependent Shedding of VEGFR2 and Crosstalk Between VEGFR2 and ERK Signaling. Circulation Research, 2008, 103, 916-918.	4.5	146
69	Different ADAMs have distinct influences on Kit ligand processing: phorbol-ester-stimulated ectodomain shedding of Kitl1 by ADAM17 is reduced by ADAM19. Journal of Cell Science, 2007, 120, 943-952.	2.0	56
70	Cutting Edge: TNF-Î±-Converting Enzyme (TACE/ADAM17) Inactivation in Mouse Myeloid Cells Prevents Lethality from Endotoxin Shock. Journal of Immunology, 2007, 179, 2686-2689.	0.8	287
71	Cell Surface Colony-Stimulating Factor 1 Can Be Cleaved by TNF-Î± Converting Enzyme or Endocytosed in a Clathrin-Dependent Manner. Journal of Immunology, 2007, 179, 6715-6724.	0.8	42
72	Substrate Selectivity of Epidermal Growth Factor-Receptor Ligand Sheddases and their Regulation by Phorbol Esters and Calcium Influx. Molecular Biology of the Cell, 2007, 18, 176-188.	2.1	276

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73	Proteolytic Processing of Delta-like 1 by ADAM Proteases. Journal of Biological Chemistry, 2007, 282, 436-444.	3.4	117
74	Ectodomain shedding of the EGF-receptor ligand epigen is mediated by ADAM17. FEBS Letters, 2007, 581, 41-44.	2.8	101
75	In search of partners: linking extracellular proteases to substrates. Nature Reviews Molecular Cell Biology, 2007, 8, 245-257.	37.0	326
76	Metalloproteases regulate T-cell proliferation and effector function via LAG-3. EMBO Journal, 2007, 26, 494-504.	7.8	203
77	A Sensitive Method to Monitor Ectodomain Shedding of Ligands of the Epidermal Growth Factor Receptor. , 2006, 327, 99-114.		46
78	ADAM10 is a principal 'sheddase' of the low-affinity immunoglobulin E receptor CD23. Nature Immunology, 2006, 7, 1293-1298.	14.5	189
79	ADAMs: key components in EGFR signalling and development. Nature Reviews Molecular Cell Biology, 2005, 6, 32-43.	37.0	989
80	Metalloprotease-disintegrin ADAM8: Expression analysis and targeted deletion in mice. Developmental Dynamics, 2005, 232, 221-231.	1.8	107
81	Homeostatic effects of the metalloproteinase disintegrin ADAM15 in degenerative cartilage remodeling. Arthritis and Rheumatism, 2005, 52, 1100-1109.	6.7	57
82	Studies from ADAM Knockout Mice. , 2005, , 29-64.		2
83	Critical Function for ADAM9 in Mouse Prostate Cancer. Cancer Research, 2005, 65, 9312-9319.	0.9	100
84	L1 Is Sequentially Processed by Two Differently Activated Metalloproteases and Presenilin/Î³-Secretase and Regulates Neural Cell Adhesion, Cell Migration, and Neurite Outgrowth. Molecular and Cellular Biology, 2005, 25, 9040-9053.	2.3	212
85	Adam Meets Eph: An ADAM Substrate Recognition Module Acts as a Molecular Switch for Ephrin Cleavage In trans. Cell, 2005, 123, 291-304.	28.9	407
86	Evaluation of the contributions of ADAMs 9, 12, 15, 17, and 19 to heart development and ectodomain shedding of neuregulins Î²1 and Î²2. Developmental Biology, 2005, 283, 459-471.	2.0	147
87	Evaluation of the Contribution of Different ADAMs to Tumor Necrosis Factor Î± (TNFÎ±) Shedding and of the Function of the TNFÎ± Ectodomain in Ensuring Selective Stimulated Shedding by the TNFÎ± Convertase (TACE/ADAM17). Journal of Biological Chemistry, 2004, 279, 42898-42906.	3.4	135
88	Essential Role for ADAM19 in Cardiovascular Morphogenesis. Molecular and Cellular Biology, 2004, 24, 96-104.	2.3	118
89	Distinct roles for ADAM10 and ADAM17 in ectodomain shedding of six EGFR ligands. Journal of Cell Biology, 2004, 164, 769-779.	5.2	895
90	Evidence for a Critical Role of the Tumor Necrosis Factor Î± Convertase (TACE) in Ectodomain Shedding of the p75 Neurotrophin Receptor (p75NTR). Journal of Biological Chemistry, 2004, 279, 4241-4249.	3.4	134

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91	Biochemical properties and functions of membrane-anchored metalloprotease-disintegrin proteins (ADAMs). Current Topics in Developmental Biology, 2003, 54, 101-123.	2.2	81
92	Catalytic Properties of ADAM19. Journal of Biological Chemistry, 2003, 278, 22331-22340.	3.4	114
93	Potential Role for ADAM15 in Pathological Neovascularization in Mice. Molecular and Cellular Biology, 2003, 23, 5614-5624.	2.3	170
94	Tumor Necrosis Factor-Î± Converting Enzyme/ADAM 17 Mediates MUC1 Shedding. Journal of Biological Chemistry, 2003, 278, 3386-3394.	3.4	177
95	Mice Lacking the Metalloprotease-Disintegrin MDC9 (ADAM9) Have No Evident Major Abnormalities during Development or Adult Life. Molecular and Cellular Biology, 2002, 22, 1537-1544.	2.3	183
96	Evidence for Regulation of the Tumor Necrosis Factor Î±-Convertase (TACE) by Protein-tyrosine Phosphatase PTPH1. Journal of Biological Chemistry, 2002, 277, 42463-42470.	3.4	116
97	The enzymatic activity of ADAM8 and ADAM9 is not regulated by TIMPs. FEBS Letters, 2002, 524, 154-158.	2.8	128
98	Catalytic activity of ADAM28. FEBS Letters, 2001, 498, 82-86.	2.8	63
99	Tumor Necrosis Factor-Î±-converting Enzyme (ADAM17) Mediates the Cleavage and Shedding of Fractalkine (CX3CL1). Journal of Biological Chemistry, 2001, 276, 37993-38001.	3.4	551
100	Biochemical and Pharmacological Criteria Define Two Shedding Activities for TRANCE/OPGL That Are Distinct from the Tumor Necrosis Factor Î± Convertase. Journal of Biological Chemistry, 2001, 276, 14665-14674.	3.4	121
101	Intracellular maturation and localization of the tumour necrosis factor Î± convertase (TACE). Biochemical Journal, 2000, 347, 131.	3.7	89
102	Cloning and characterization of ADAM28: evidence for autocatalytic pro-domain removal and for cell surface localization of mature ADAM28. Biochemical Journal, 2000, 348, 21-27.	3.7	116
103	Intracellular maturation and localization of the tumour necrosis factor Î± convertase (TACE). Biochemical Journal, 2000, 347, 131-138.	3.7	320
104	Interaction of the Metalloprotease Disintegrins MDC9 and MDC15 with Two SH3 Domain-containing Proteins, Endophilin I and SH3PX1. Journal of Biological Chemistry, 1999, 274, 31693-31699.	3.4	157
105	Metalloprotease-Disintegrin MDC9: Intracellular Maturation and Catalytic Activity. Journal of Biological Chemistry, 1999, 274, 3531-3540.	3.4	284
106	Evidence for an interaction of the metalloproteaseâ€“disintegrin tumour necrosis factor Î± convertase (TACE) with mitotic arrest deficient 2 (MAD2), and of the metalloproteaseâ€“disintegrin MDC9 with a novel MAD2-related protein, MAD2Î². Biochemical Journal, 1999, 343, 673-680.	3.7	80
107	Evidence for an interaction of the metalloproteaseâ€™disintegrin tumour necrosis factor Î± convertase (TACE) with mitotic arrest deficient 2 (MAD2), and of the metalloproteaseâ€™disintegrin MDC9 with a novel MAD2-related protein, MAD2Î². Biochemical Journal, 1999, 343, 673.	3.7	50
108	Intracellular Maturation of the Mouse Metalloprotease Disintegrin MDC15. Journal of Biological Chemistry, 1998, 273, 26236-26247.	3.4	145

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109	Cloning and Initial Characterization of Mouse Meltrin Î² and Analysis of the Expression of Four MetalloproteaseDisintegrins in Bone Cells. Journal of Biological Chemistry, 1998, 273, 4180-4187.	3.4	100
110	Metalloprotease-Disintegrins: Links to Cell Adhesion and Cleavage of TNFÎ± and Notch. Cell, 1997, 90, 589-592.	28.9	371
111	Metargidin, a Membrane-anchored Metalloprotease-Disintegrin Protein with an RGD Integrin Binding Sequence. Journal of Biological Chemistry, 1996, 271, 4593-4596.	3.4	154
112	A potential fusion peptide and an integrin ligand domain in a protein active in spermâ€“egg fusion. Nature, 1992, 356, 248-252.	27.8	708