

Andreas Matouschek

List of Publications by Year in descending order

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86
papers

9,579
citations

53660

45
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58464

82
g-index

92
all docs

92
docs citations

92
times ranked

6614
citing authors

#	ARTICLE	IF	CITATIONS
1	Design principles that protect the proteasome from self-destruction. <i>Protein Science</i> , 2022, 31, 556-567.	3.1	2
2	Mechanisms of substrate recognition by the 26S proteasome. <i>Current Opinion in Structural Biology</i> , 2021, 67, 161-169.	2.6	34
3	Use of Multiple Ion Fragmentation Methods to Identify Protein Cross-Links and Facilitate Comparison of Data Interpretation Algorithms. <i>Journal of Proteome Research</i> , 2020, 19, 2758-2771.	1.8	3
4	The proteasome 19S cap and its ubiquitin receptors provide a versatile recognition platform for substrates. <i>Nature Communications</i> , 2020, 11, 477.	5.8	101
5	A masked initiation region in retinoblastoma protein regulates its proteasomal degradation. <i>Nature Communications</i> , 2020, 11, 2019.	5.8	33
6	Mechanical unfolding of spectrin reveals a super-exponential dependence of unfolding rate on force. <i>Scientific Reports</i> , 2019, 9, 11101.	1.6	9
7	Substrate selection by the proteasome through initiation regions. <i>Protein Science</i> , 2019, 28, 1222-1232.	3.1	26
8	Decoding without the cipher. <i>Nature Chemical Biology</i> , 2019, 15, 210-212.	3.9	2
9	Scalable In Vitro Proteasome Activity Assay. <i>Methods in Molecular Biology</i> , 2018, 1844, 321-341.	0.4	7
10	Recognition of Client Proteins by the Proteasome. <i>Annual Review of Biophysics</i> , 2017, 46, 149-173.	4.5	99
11	Mouse Mammary Tumor Virus Signal Peptide Uses a Novel p97-Dependent and Derlin-Independent Retrotranslocation Mechanism To Escape Proteasomal Degradation. <i>MBio</i> , 2017, 8, .	1.8	12
12	An Inducible System for Rapid Degradation of Specific Cellular Proteins Using Proteasome Adaptors. <i>PLoS ONE</i> , 2016, 11, e0152679.	1.1	25
13	Conserved Sequence Preferences Contribute to Substrate Recognition by the Proteasome. <i>Journal of Biological Chemistry</i> , 2016, 291, 14526-14539.	1.6	56
14	Ubiquitin-like domains can target to the proteasome but proteolysis requires a disordered region. <i>EMBO Journal</i> , 2016, 35, 1522-1536.	3.5	52
15	An assay for 26S proteasome activity based on fluorescence anisotropy measurements of dye-labeled protein substrates. <i>Analytical Biochemistry</i> , 2016, 509, 50-59.	1.1	22
16	Ramping up degradation for proliferation. <i>Nature Cell Biology</i> , 2016, 18, 141-142.	4.6	3
17	A Rapid and Versatile Method for Generating Proteins with Defined Ubiquitin Chains. <i>Biochemistry</i> , 2016, 55, 1898-1908.	1.2	36
18	Sequence composition of disordered regions fine-tunes protein half-life. <i>Nature Structural and Molecular Biology</i> , 2015, 22, 214-221.	3.6	109

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19	Top-Down 193-nm Ultraviolet Photodissociation Mass Spectrometry for Simultaneous Determination of Polyubiquitin Chain Length and Topology. <i>Analytical Chemistry</i> , 2015, 87, 1812-1820.	3.2	41
20	Regulation of Proteasomal Degradation by Modulating Proteasomal Initiation Regions. <i>ACS Chemical Biology</i> , 2015, 10, 2537-2543.	1.6	13
21	Intrinsically Disordered Segments Affect Protein Half-Life in the Cell and during Evolution. <i>Cell Reports</i> , 2014, 8, 1832-1844.	2.9	192
22	Paradigms of protein degradation by the proteasome. <i>Current Opinion in Structural Biology</i> , 2014, 24, 156-164.	2.6	102
23	Disordered Proteinaceous Machines. <i>Chemical Reviews</i> , 2014, 114, 6806-6843.	23.0	109
24	Regulated protein turnover: snapshots of the proteasome in action. <i>Nature Reviews Molecular Cell Biology</i> , 2014, 15, 122-133.	16.1	212
25	Chance, Destiny, and the Inner Workings of ClpXP. <i>Cell</i> , 2014, 158, 479-480.	13.5	2
26	An Ancient Portal to Proteolysis. <i>Science</i> , 2012, 337, 813-814.	6.0	5
27	Sequence- and Species-Dependence of Proteasomal Processivity. <i>ACS Chemical Biology</i> , 2012, 7, 1444-1453.	1.6	50
28	Proteasomal Degradation from Internal Sites Favors Partial Proteolysis <i>via</i> Remote Domain Stabilization. <i>ACS Chemical Biology</i> , 2011, 6, 1087-1095.	1.6	27
29	How ClpX Unfolds GFP in Stages by Pulling. <i>Journal of Molecular Biology</i> , 2011, 413, 1-3.	2.0	0
30	Defining the geometry of the two-component proteasome degron. <i>Nature Chemical Biology</i> , 2011, 7, 161-167.	3.9	149
31	Rad23 escapes degradation because it lacks a proteasome initiation region. <i>Nature Communications</i> , 2011, 2, 192.	5.8	87
32	A Three-part Signal Governs Differential Processing of Gli1 and Gli3 Proteins by the Proteasome. <i>Journal of Biological Chemistry</i> , 2011, 286, 39051-39058.	1.6	33
33	Making It Easier to Regulate Protein Stability. <i>Chemistry and Biology</i> , 2010, 17, 917-918.	6.2	3
34	Pup grows up: in vitro characterization of the degradation of pupylated proteins. <i>EMBO Journal</i> , 2010, 29, 1163-1164.	3.5	3
35	Substrate selection by the proteasome during degradation of protein complexes. <i>Nature Chemical Biology</i> , 2009, 5, 29-36.	3.9	108
36	Targeting proteins for degradation. <i>Nature Chemical Biology</i> , 2009, 5, 815-822.	3.9	260

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37	ATP-dependent Proteases Differ Substantially in Their Ability to Unfold Globular Proteins. <i>Journal of Biological Chemistry</i> , 2009, 284, 18674-18684.	1.6	69
38	How to pick a protein and pull at it. <i>Nature Structural and Molecular Biology</i> , 2008, 15, 1135-1136.	3.6	7
39	Protein targeting to ATP-dependent proteases. <i>Current Opinion in Structural Biology</i> , 2008, 18, 43-51.	2.6	36
40	Controlling a Single Protein in a Nanopore through Electrostatic Traps. <i>Journal of the American Chemical Society</i> , 2008, 130, 4081-4088.	6.6	109
41	1P041 SELECTING PROTEINS FOR DEGRADATION: THE INITIATION STEP(Proteins-functions, methodology,) Tj ETQq1_1 0.784314 rgBT	0.0	0
42	To degrade or release: ubiquitin-chain remodeling. <i>Trends in Cell Biology</i> , 2007, 17, 419-421.	3.6	29
43	Where to start and when to stop. <i>Nature Structural and Molecular Biology</i> , 2006, 13, 668-670.	3.6	10
44	A conserved processing mechanism regulates the activity of transcription factors Cubitus interruptus and NF- κ B. <i>Nature Structural and Molecular Biology</i> , 2005, 12, 1045-1053.	3.6	106
45	Effect of protein structure on mitochondrial import. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 15435-15440.	3.3	94
46	\hat{I}^2 -Synuclein Reduces Proteasomal Inhibition by \hat{I}^1 -Synuclein but Not \hat{I}^3 -Synuclein. <i>Journal of Biological Chemistry</i> , 2005, 280, 7562-7569.	1.6	49
47	Inefficient degradation of truncated polyglutamine proteins by the proteasome. <i>EMBO Journal</i> , 2004, 23, 4307-4318.	3.5	258
48	An unstructured initiation site is required for efficient proteasome-mediated degradation. <i>Nature Structural and Molecular Biology</i> , 2004, 11, 830-837.	3.6	404
49	Protein unfolding in the cell. <i>Trends in Biochemical Sciences</i> , 2004, 29, 593-600.	3.7	125
50	The Force Exerted by the Membrane Potential during Protein Import into the Mitochondrial Matrix. <i>Biophysical Journal</i> , 2004, 86, 3647-3652.	0.2	38
51	Protein unfolding " an important process in vivo?. <i>Current Opinion in Structural Biology</i> , 2003, 13, 98-109.	2.6	153
52	Finding a protein's Achilles heel. <i>Nature Structural and Molecular Biology</i> , 2003, 10, 674-676.	3.6	27
53	Lack of a Robust Unfoldase Activity Confers a Unique Level of Substrate Specificity to the Universal AAA Protease FtsH. <i>Molecular Cell</i> , 2003, 11, 659-669.	4.5	163
54	Aggregated and Monomeric \hat{I}^1 -Synuclein Bind to the S6 \hat{I}^2 Proteasomal Protein and Inhibit Proteasomal Function. <i>Journal of Biological Chemistry</i> , 2003, 278, 11753-11759.	1.6	364

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55	Concurrent Translocation of Multiple Polypeptide Chains through the Proteasomal Degradation Channel. <i>Journal of Biological Chemistry</i> , 2002, 277, 34760-34765.	1.6	57
56	Protein unfolding by the mitochondrial membrane potential. <i>Nature Structural Biology</i> , 2002, 9, 301-307.	9.7	119
57	ATP-Dependent Proteases Degrade Their Substrates by Processively Unraveling Them from the Degradation Signal. <i>Molecular Cell</i> , 2001, 7, 627-637.	4.5	380
58	Barreling through the outer membrane. , 2001, 8, 284-286.		24
59	Recognizing misfolded proteins in the endoplasmic reticulum. , 2000, 7, 265-266.		5
60	Protein unfolding by mitochondria. <i>EMBO Reports</i> , 2000, 1, 404-410.	2.0	160
61	Effect of the protein import machinery at the mitochondrial surface on precursor stability. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2000, 97, 12991-12996.	3.3	21
62	The Structure of Precursor Proteins during Import into Mitochondria. <i>Journal of Biological Chemistry</i> , 1999, 274, 12759-12764.	1.6	47
63	The dimensions of the protein import channels in the outer and inner mitochondrial membranes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1999, 96, 13086-13090.	3.3	109
64	Mitochondria unfold precursor proteins by unraveling them from their N-termini. <i>Nature Structural Biology</i> , 1999, 6, 1132-1138.	9.7	110
65	Active unfolding of precursor proteins during mitochondrial protein import. <i>EMBO Journal</i> , 1997, 16, 6727-6736.	3.5	140
66	Hsp60-independent protein folding in the matrix of yeast mitochondria.. <i>EMBO Journal</i> , 1996, 15, 764-774.	3.5	81
67	Hsp60-independent protein folding in the matrix of yeast mitochondria. <i>EMBO Journal</i> , 1996, 15, 764-74.	3.5	25
68	Cyclophilin catalyzes protein folding in yeast mitochondria.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1995, 92, 6319-6323.	3.3	206
69	Movement of the position of the transition state in protein folding. <i>Biochemistry</i> , 1995, 34, 13656-13662.	1.2	133
70	Import and Folding of Proteins by Mitochondria. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 1995, 60, 609-617.	2.0	6
71	Extrapolation to water of kinetic and equilibrium data for the unfolding of barnase in urea solutions. <i>Protein Engineering, Design and Selection</i> , 1994, 7, 1089-1095.	1.0	71
72	Application of physical organic chemistry to engineered mutants of proteins: Hammond postulate behavior in the transition state of protein folding.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 7814-7818.	3.3	199

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73	Pathway of protein folding. Faraday Discussions, 1992, 93, 183.	1.6	13
74	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 783-804.	2.0	421
75	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 805-818.	2.0	269
76	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 819-835.	2.0	222
77	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 771-782.	2.0	855
78	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 837-845.	2.0	100
79	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 847-859.	2.0	169
80	Pathway and stability of protein folding. Philosophical Transactions of the Royal Society B: Biological Sciences, 1991, 332, 171-176.	1.8	22
81	Physical-organic molecular biology: pathway and stability of protein folding. Pure and Applied Chemistry, 1991, 63, 187-194.	0.9	7
82	[6] Protein engineering in analysis of protein folding pathways and stability. Methods in Enzymology, 1991, 202, 82-112.	0.4	81
83	Transient folding intermediates characterized by protein engineering. Nature, 1990, 346, 440-445.	13.7	501
84	Detection and characterization of a folding intermediate in barnase by NMR. Nature, 1990, 346, 488-490.	13.7	241
85	Mapping the transition state and pathway of protein folding by protein engineering. Nature, 1989, 340, 122-126.	13.7	715
86	ATP-Dependent Proteases: The Cell's Degradation Machines. , 0, , 239-260.		1