Andreas Matouschek

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 771-782.	4.2	855
2	Mapping the transition state and pathway of protein folding by protein engineering. Nature, 1989, 340, 122-126.	27.8	715
3	Transient folding intermediates characterized by protein engineering. Nature, 1990, 346, 440-445.	27.8	501
4	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 783-804.	4.2	421
5	An unstructured initiation site is required for efficient proteasome-mediated degradation. Nature Structural and Molecular Biology, 2004, 11, 830-837.	8.2	404
6	ATP-Dependent Proteases Degrade Their Substrates by Processively Unraveling Them from the Degradation Signal. Molecular Cell, 2001, 7, 627-637.	9.7	380
7	Aggregated and Monomeric α-Synuclein Bind to the S6′ Proteasomal Protein and Inhibit Proteasomal Function. Journal of Biological Chemistry, 2003, 278, 11753-11759.	3.4	364
8	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 805-818.	4.2	269
9	Targeting proteins for degradation. Nature Chemical Biology, 2009, 5, 815-822.	8.0	260
10	Inefficient degradation of truncated polyglutamine proteins by the proteasome. EMBO Journal, 2004, 23, 4307-4318.	7.8	258
11	Detection and characterization of a folding intermediate in barnase by NMR. Nature, 1990, 346, 488-490.	27.8	241
12	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 819-835.	4.2	222
13	Regulated protein turnover: snapshots of the proteasome in action. Nature Reviews Molecular Cell Biology, 2014, 15, 122-133.	37.0	212
14	Cyclophilin catalyzes protein folding in yeast mitochondria Proceedings of the National Academy of Sciences of the United States of America, 1995, 92, 6319-6323.	7.1	206
15	Application of physical organic chemistry to engineered mutants of proteins: Hammond postulate behavior in the transition state of protein folding Proceedings of the National Academy of Sciences of the United States of America, 1993, 90, 7814-7818.	7.1	199
16	Intrinsically Disordered Segments Affect Protein Half-Life in the Cell and during Evolution. Cell Reports, 2014, 8, 1832-1844.	6.4	192
17	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 847-859.	4.2	169
18	Lack of a Robust Unfoldase Activity Confers a Unique Level of Substrate Specificity to the Universal AAA Protease EtsH. Molecular Cell. 2003. 11, 659-669.	9.7	163

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#	Article	IF	CITATIONS
19	Protein unfolding by mitochondria. EMBO Reports, 2000, 1, 404-410.	4.5	160
20	Protein unfolding — an important process in vivo?. Current Opinion in Structural Biology, 2003, 13, 98-109.	5.7	153
21	Defining the geometry of the two-component proteasome degron. Nature Chemical Biology, 2011, 7, 161-167.	8.0	149
22	Active unfolding of precursor proteins during mitochondrial protein import. EMBO Journal, 1997, 16, 6727-6736.	7.8	140
23	Movement of the position of the transition state in protein folding. Biochemistry, 1995, 34, 13656-13662.	2.5	133
24	Protein unfolding in the cell. Trends in Biochemical Sciences, 2004, 29, 593-600.	7.5	125
25	Protein unfolding by the mitochondrial membrane potential. Nature Structural Biology, 2002, 9, 301-307.	9.7	119
26	Mitochondria unfold precursor proteins by unraveling them from their N-termini. Nature Structural Biology, 1999, 6, 1132-1138.	9.7	110
27	The dimensions of the protein import channels in the outer and inner mitochondrial membranes. Proceedings of the National Academy of Sciences of the United States of America, 1999, 96, 13086-13090.	7.1	109
28	Controlling a Single Protein in a Nanopore through Electrostatic Traps. Journal of the American Chemical Society, 2008, 130, 4081-4088.	13.7	109
29	Disordered Proteinaceous Machines. Chemical Reviews, 2014, 114, 6806-6843.	47.7	109
30	Sequence composition of disordered regions fine-tunes protein half-life. Nature Structural and Molecular Biology, 2015, 22, 214-221.	8.2	109
31	Substrate selection by the proteasome during degradation of protein complexes. Nature Chemical Biology, 2009, 5, 29-36.	8.0	108
32	A conserved processing mechanism regulates the activity of transcription factors Cubitus interruptus and NF-κB. Nature Structural and Molecular Biology, 2005, 12, 1045-1053.	8.2	106
33	Paradigms of protein degradation by the proteasome. Current Opinion in Structural Biology, 2014, 24, 156-164.	5.7	102
34	The proteasome 19S cap and its ubiquitin receptors provide a versatile recognition platform for substrates. Nature Communications, 2020, 11, 477.	12.8	101
35	The folding of an enzyme. Journal of Molecular Biology, 1992, 224, 837-845.	4.2	100
36	Recognition of Client Proteins by the Proteasome. Annual Review of Biophysics, 2017, 46, 149-173.	10.0	99

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37	Effect of protein structure on mitochondrial import. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 15435-15440.	7.1	94
38	Rad23 escapes degradation because it lacks a proteasome initiation region. Nature Communications, 2011, 2, 192.	12.8	87
39	[6] Protein engineering in analysis of protein folding pathways and stability. Methods in Enzymology, 1991, 202, 82-112.	1.0	81
40	Hsp60-independent protein folding in the matrix of yeast mitochondria EMBO Journal, 1996, 15, 764-774.	7.8	81
41	Extrapolation to water of kinetic and equilibrium data for the unfolding of barnase in urea solutions. Protein Engineering, Design and Selection, 1994, 7, 1089-1095.	2.1	71
42	ATP-dependent Proteases Differ Substantially in Their Ability to Unfold Globular Proteins. Journal of Biological Chemistry, 2009, 284, 18674-18684.	3.4	69
43	Concurrent Translocation of Multiple Polypeptide Chains through the Proteasomal Degradation Channel. Journal of Biological Chemistry, 2002, 277, 34760-34765.	3.4	57
44	Conserved Sequence Preferences Contribute to Substrate Recognition by the Proteasome. Journal of Biological Chemistry, 2016, 291, 14526-14539.	3.4	56
45	Ubiquitinâ€like domains can target to the proteasome but proteolysis requires a disordered region. EMBO Journal, 2016, 35, 1522-1536.	7.8	52
46	Sequence- and Species-Dependence of Proteasomal Processivity. ACS Chemical Biology, 2012, 7, 1444-1453.	3.4	50
47	β-Synuclein Reduces Proteasomal Inhibition by α-Synuclein but Not γ-Synuclein. Journal of Biological Chemistry, 2005, 280, 7562-7569.	3.4	49
48	The Structure of Precursor Proteins during Import into Mitochondria. Journal of Biological Chemistry, 1999, 274, 12759-12764.	3.4	47
49	Top-Down 193-nm Ultraviolet Photodissociation Mass Spectrometry for Simultaneous Determination of Polyubiquitin Chain Length and Topology. Analytical Chemistry, 2015, 87, 1812-1820.	6.5	41
50	The Force Exerted by the Membrane Potential during Protein Import into the Mitochondrial Matrix. Biophysical Journal, 2004, 86, 3647-3652.	0.5	38
51	Protein targeting to ATP-dependent proteases. Current Opinion in Structural Biology, 2008, 18, 43-51.	5.7	36
52	A Rapid and Versatile Method for Generating Proteins with Defined Ubiquitin Chains. Biochemistry, 2016, 55, 1898-1908.	2.5	36
53	Mechanisms of substrate recognition by the 26S proteasome. Current Opinion in Structural Biology, 2021, 67, 161-169.	5.7	34
54	A Three-part Signal Governs Differential Processing of Gli1 and Gli3 Proteins by the Proteasome. Journal of Biological Chemistry, 2011, 286, 39051-39058.	3.4	33

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55	A masked initiation region in retinoblastoma protein regulates its proteasomal degradation. Nature Communications, 2020, 11, 2019.	12.8	33
56	To degrade or release: ubiquitin-chain remodeling. Trends in Cell Biology, 2007, 17, 419-421.	7.9	29
57	Finding a protein's Achilles heel. Nature Structural and Molecular Biology, 2003, 10, 674-676.	8.2	27
58	Proteasomal Degradation from Internal Sites Favors Partial Proteolysis <i>via</i> Remote Domain Stabilization. ACS Chemical Biology, 2011, 6, 1087-1095.	3.4	27
59	Substrate selection by the proteasome through initiation regions. Protein Science, 2019, 28, 1222-1232.	7.6	26
60	An Inducible System for Rapid Degradation of Specific Cellular Proteins Using Proteasome Adaptors. PLoS ONE, 2016, 11, e0152679.	2.5	25
61	Hsp60-independent protein folding in the matrix of yeast mitochondria. EMBO Journal, 1996, 15, 764-74.	7.8	25
62	Barreling through the outer membrane. , 2001, 8, 284-286.		24
63	Pathway and stability of protein folding. Philosophical Transactions of the Royal Society B: Biological Sciences, 1991, 332, 171-176.	4.0	22
64	An assay for 26S proteasome activity based on fluorescence anisotropy measurements of dye-labeled protein substrates. Analytical Biochemistry, 2016, 509, 50-59.	2.4	22
65	Effect of the protein import machinery at the mitochondrial surface on precursor stability. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 12991-12996.	7.1	21
66	Pathway of protein folding. Faraday Discussions, 1992, 93, 183.	3.2	13
67	Regulation of Proteasomal Degradation by Modulating Proteasomal Initiation Regions. ACS Chemical Biology, 2015, 10, 2537-2543.	3.4	13
68	Mouse Mammary Tumor Virus Signal Peptide Uses a Novel p97-Dependent and Derlin-Independent Retrotranslocation Mechanism To Escape Proteasomal Degradation. MBio, 2017, 8, .	4.1	12
69	Where to start and when to stop. Nature Structural and Molecular Biology, 2006, 13, 668-670.	8.2	10
70	Mechanical unfolding of spectrin reveals a super-exponential dependence of unfolding rate on force. Scientific Reports, 2019, 9, 11101.	3.3	9
71	Physical-organic molecular biology: pathway and stability of protein folding. Pure and Applied Chemistry, 1991, 63, 187-194.	1.9	7
72	How to pick a protein and pull at it. Nature Structural and Molecular Biology, 2008, 15, 1135-1136.	8.2	7

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73	Scalable In Vitro Proteasome Activity Assay. Methods in Molecular Biology, 2018, 1844, 321-341.	0.9	7
74	Import and Folding of Proteins by Mitochondria. Cold Spring Harbor Symposia on Quantitative Biology, 1995, 60, 609-617.	1.1	6
75	Recognizing misfolded proteins in the endoplasmic reticulum. , 2000, 7, 265-266.		5
76	An Ancient Portal to Proteolysis. Science, 2012, 337, 813-814.	12.6	5
77	Making It Easier to Regulate Protein Stability. Chemistry and Biology, 2010, 17, 917-918.	6.0	3
78	Pup grows up: in vitro characterization of the degradation of pupylated proteins. EMBO Journal, 2010, 29, 1163-1164.	7.8	3
79	Ramping up degradation for proliferation. Nature Cell Biology, 2016, 18, 141-142.	10.3	3
80	Use of Multiple Ion Fragmentation Methods to Identify Protein Cross-Links and Facilitate Comparison of Data Interpretation Algorithms. Journal of Proteome Research, 2020, 19, 2758-2771.	3.7	3
81	Chance, Destiny, and the Inner Workings of ClpXP. Cell, 2014, 158, 479-480.	28.9	2
82	Decoding without the cipher. Nature Chemical Biology, 2019, 15, 210-212.	8.0	2
83	Design principles that protect the proteasome from selfâ€destruction. Protein Science, 2022, 31, 556-567.	7.6	2
84	ATP-Dependent Proteases: The Cell's Degradation Machines. , 0, , 239-260.		1
85	1P041 SELECTING PROTEINS FOR DEGRADATION: THE INITIATION STEP(Proteins-functions, methodology,) Tj ET(Qq110.78	34314 rgBT /(
86	How ClpX Unfolds GFP in Stages by Pulling. Journal of Molecular Biology, 2011, 413, 1-3.	4.2	0