

Harald Kolmar

List of Publications by Year in descending order

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202
papers

6,226
citations

66343

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106344

65
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all docs

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docs citations

217
times ranked

5851
citing authors

#	ARTICLE	IF	CITATIONS
1	Dimerisation of the Glycophorin A Transmembrane Segment in Membranes Probed with the ToxR Transcription Activator. <i>Journal of Molecular Biology</i> , 1996, 263, 525-530.	4.2	237
2	Decorating microbes: surface display of proteins on <i>Escherichia coli</i> . <i>Trends in Biotechnology</i> , 2011, 29, 79-86.	9.3	198
3	Camelid and shark single domain antibodies: structural features and therapeutic potential. <i>Current Opinion in Structural Biology</i> , 2017, 45, 10-16.	5.7	165
4	The <i>vsr</i> gene product of <i>E. coli</i> K-12 is a strand- and sequence-specific DNA mismatch endonuclease. <i>Nature</i> , 1991, 353, 776-778.	27.8	142
5	The DegP and DegQ periplasmic endoproteases of <i>Escherichia coli</i> : specificity for cleavage sites and substrate conformation. <i>Journal of Bacteriology</i> , 1996, 178, 5925-5929.	2.2	130
6	Bi-specific Aptamers Mediating Tumor Cell Lysis. <i>Journal of Biological Chemistry</i> , 2011, 286, 21896-21905.	3.4	124
7	The crystal structure of SdsA1, an alkylsulfatase from <i>Pseudomonas aeruginosa</i> , defines a third class of sulfatases. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 7631-7636.	7.1	115
8	Ultra-high-throughput screening based on cell-surface display and fluorescence-activated cell sorting for the identification of novel biocatalysts. <i>Current Opinion in Biotechnology</i> , 2004, 15, 323-329.	6.6	114
9	Triazole Bridge Disulfide Bond Replacement by Ruthenium Catalyzed Formation of 1,5-Disubstituted 1,2,3-Triazoles. <i>Angewandte Chemie - International Edition</i> , 2011, 50, 5207-5211.	13.8	112
10	Biological diversity and therapeutic potential of natural and engineered cystine knot miniproteins. <i>Current Opinion in Pharmacology</i> , 2009, 9, 608-614.	3.5	108
11	Structural insights and biomedical potential of IgNAR scaffolds from sharks. <i>MAbs</i> , 2015, 7, 15-25.	5.2	102
12	Therapeutic antibody engineering by high efficiency cell screening. <i>FEBS Letters</i> , 2014, 588, 278-287.	2.8	95
13	Engineered Cystine Knot Miniproteins as Potent Inhibitors of Human Mast Cell Tryptase β^2 . <i>Journal of Molecular Biology</i> , 2010, 395, 167-175.	4.2	92
14	The cystine knot of a squash-type protease inhibitor as a structural scaffold for <i>Escherichia coli</i> cell surface display of conformationally constrained peptides. <i>Protein Engineering, Design and Selection</i> , 1999, 12, 797-806.	2.1	87
15	Alternative binding proteins: Biological activity and therapeutic potential of cystine knot miniproteins. <i>FEBS Journal</i> , 2008, 275, 2684-2690.	4.7	87
16	Single-Cell High-Throughput Screening To Identify Enantioselective Hydrolytic Enzymes. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 5085-5088.	13.8	81
17	The potential of cystine-knot microproteins as novel pharmacophoric scaffolds in oral peptide drug delivery. <i>Journal of Drug Targeting</i> , 2006, 14, 137-146.	4.4	79
18	A generic system for the <i>Escherichia coli</i> cell-surface display of lipolytic enzymes. <i>FEBS Letters</i> , 2005, 579, 1177-1182.	2.8	76

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19	Shark Attack: High affinity binding proteins derived from shark vNAR domains by stepwise in vitro affinity maturation. <i>Journal of Biotechnology</i> , 2014, 191, 236-245.	3.8	74
20	Display of Passenger Proteins on the Surface of Escherichia coli K-12 by the Enterohemorrhagic E. coli Intimin EaeA. <i>Journal of Bacteriology</i> , 2001, 183, 7273-7284.	2.2	71
21	Arranged Sevenfold: Structural Insights into the C-Terminal Oligomerization Domain of Human C4b-Binding Protein. <i>Journal of Molecular Biology</i> , 2013, 425, 1302-1317.	4.2	69
22	Inhibition of platelet aggregation by grafting RGD and KGD sequences on the structural scaffold of small disulfide-rich proteins. <i>Platelets</i> , 2006, 17, 153-157.	2.3	67
23	Knottin cyclization: impact on structure and dynamics. <i>BMC Structural Biology</i> , 2008, 8, 54.	2.3	66
24	Balancing Selectivity and Efficacy of Bispecific Epidermal Growth Factor Receptor (EGFR) $\tilde{\text{A}}$ - c-MET Antibodies and Antibody-Drug Conjugates. <i>Journal of Biological Chemistry</i> , 2016, 291, 25106-25119.	3.4	66
25	Trypsin inhibition by macrocyclic and open-chain variants of the squash inhibitor MCoTI-II. <i>Biological Chemistry</i> , 2005, 386, 1301-6.	2.5	65
26	Crystal structure of the electron transfer complex rubredoxin $\tilde{\text{A}}$ -rubredoxin reductase of <i>Pseudomonas aeruginosa</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 12276-12281.	7.1	65
27	A generic approach to engineer antibody pH-switches using combinatorial histidine scanning libraries and yeast display. <i>MAbs</i> , 2015, 7, 138-151.	5.2	64
28	Single-domain antibodies for biomedical applications. <i>Immunopharmacology and Immunotoxicology</i> , 2016, 38, 21-28.	2.4	64
29	Braces for the Peptide Backbone: Insights into Structure $\tilde{\text{A}}$ Activity Relationships of Protease Inhibitor Mimics with Locked Amide Conformations. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 3708-3712.	13.8	62
30	Contribution of the intramolecular disulfide bridge to the folding stability of REIv, the variable domain of a human immunoglobulin $\tilde{\text{I}}$ light chain. <i>Folding & Design</i> , 1996, 1, 431-440.	4.5	61
31	Sequence Requirements of the GPNG $\tilde{\text{I}}$ -Turn of the Ecballium elaterium Trypsin Inhibitor II Explored by Combinatorial Library Screening. <i>Journal of Biological Chemistry</i> , 1999, 274, 21037-21043.	3.4	61
32	Beyond antibody engineering: directed evolution of alternative binding scaffolds and enzymes using yeast surface display. <i>Microbial Cell Factories</i> , 2018, 17, 32.	4.0	58
33	Grafting of thrombopoietin-mimetic peptides into cystine knot miniproteins yields high-affinity thrombopoietin antagonists and agonists. <i>FEBS Journal</i> , 2007, 274, 86-95.	4.7	57
34	Epitope mapping and affinity purification of monospecific antibodies by Escherichia coli cell surface display of gene-derived random peptide libraries. <i>Journal of Immunological Methods</i> , 2001, 257, 163-173.	1.4	56
35	Combinatorial Optimization of Cystine-Knot Peptides towards High-Affinity Inhibitors of Human Matriptase-1. <i>PLoS ONE</i> , 2013, 8, e76956.	2.5	55
36	Evaluation and improvement of the properties of the novel cystine-knot microprotein McoEeTI for oral administration. <i>International Journal of Pharmaceutics</i> , 2007, 332, 72-79.	5.2	52

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37	Towards click bioconjugations on cube-octameric silsesquioxane scaffolds. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 2212.	2.8	49
38	Locked by Design: A Conformationally Constrained Transglutaminase Tag Enables Efficient Site-Specific Conjugation. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13420-13424.	13.8	49
39	Combinatorial tuning of peptidic drug candidates: high-affinity matriptase inhibitors through incremental structure-guided optimization. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 1848.	2.8	48
40	A Chemoenzymatic Approach to Protein Immobilization onto Crystalline Cellulose Nanoscaffolds. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 12618-12623.	13.8	48
41	Ultrahigh-Throughput Screening to Identify <i>E. coli</i> Cells Expressing Functionally Active Enzymes on their Surface. <i>ChemBioChem</i> , 2007, 8, 943-949.	2.6	47
42	Bioconjugation on cube-octameric silsesquioxanes. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 2224.	2.8	44
43	Chemical Synthesis, Backbone Cyclization and Oxidative Folding of Cystine-knot Peptides – Promising Scaffolds for Applications in Drug Design. <i>Molecules</i> , 2012, 17, 12533-12552.	3.8	43
44	Engineering bispecific antibodies with defined chain pairing. <i>New Biotechnology</i> , 2017, 39, 167-173.	4.4	43
45	Potent inhibitors of human matriptase-1 based on the scaffold of sunflower trypsin inhibitor. <i>Journal of Peptide Science</i> , 2014, 20, 415-420.	1.4	42
46	Natural and Engineered Cystine Knot Miniproteins for Diagnostic and Therapeutic Applications. <i>Current Pharmaceutical Design</i> , 2011, 17, 4329-4336.	1.9	41
47	In Vivo Enzyme Immobilization by Inclusion Body Display. <i>Applied and Environmental Microbiology</i> , 2010, 76, 5563-5569.	3.1	40
48	Synthesis and characterization of new generation open tubular silica capillaries for liquid chromatography. <i>Journal of Chromatography A</i> , 2012, 1265, 88-94.	3.7	40
49	Functional genomics of <i>Pseudomonas aeruginosa</i> to identify habitat-specific determinants of pathogenicity. <i>International Journal of Medical Microbiology</i> , 2007, 297, 615-623.	3.6	39
50	Structure of <i>Ecballium elaterium</i> trypsin inhibitor II (EETI-II): a rigid molecular scaffold. <i>Acta Crystallographica Section D: Biological Crystallography</i> , 2005, 61, 1255-1262.	2.5	38
51	Recent progress in transglutaminase-mediated assembly of antibody-drug conjugates. <i>Analytical Biochemistry</i> , 2020, 595, 113615.	2.4	38
52	Engineering IgG-Like Bispecific Antibodies – An Overview. <i>Antibodies</i> , 2018, 7, 28.	2.5	37
53	A fusion protein system for the recombinant production of short disulfide bond rich cystine knot peptides using barnase as a purification handle. <i>Protein Expression and Purification</i> , 2005, 39, 82-89.	1.3	36
54	Covalent Attachment of Enzymes to Paper Fibers for Paper-Based Analytical Devices. <i>Frontiers in Chemistry</i> , 2018, 6, 214.	3.6	35

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55	Host cell protein quantification by fourier transform mid infrared spectroscopy (FT-MIR). <i>Biotechnology and Bioengineering</i> , 2013, 110, 252-259.	3.3	34
56	Effective PHIP Labeling of Bioactive Peptides Boosts the Intensity of the NMR Signal. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 12941-12945.	13.8	34
57	Semi-synthetic vNAR libraries screened against therapeutic antibodies primarily deliver anti-idiotypic binders. <i>Scientific Reports</i> , 2017, 7, 9676.	3.3	34
58	Generation of human bispecific common light chain antibodies by combining animal immunization and yeast display. <i>Protein Engineering, Design and Selection</i> , 2017, 30, 291-301.	2.1	33
59	Intimin-Mediated Export of Passenger Proteins Requires Maintenance of a Translocation-Competent Conformation. <i>Journal of Bacteriology</i> , 2005, 187, 522-533.	2.2	32
60	Head-to-Tail Cyclized Cysteine-Knot Peptides by a Combined Recombinant and Chemical Route of Synthesis. <i>ChemBioChem</i> , 2008, 9, 33-37.	2.6	32
61	Cysteine-knot peptides targeting cancer-relevant human cytotoxic T lymphocyte-associated antigen 4 (CTLA4). <i>Journal of Peptide Science</i> , 2015, 21, 651-660.	1.4	32
62	At-line mid infrared spectroscopy for monitoring downstream processing unit operations. <i>Process Biochemistry</i> , 2015, 50, 997-1005.	3.7	32
63	Affinity Maturation of B7-H6 Translates into Enhanced NK Cell-Mediated Tumor Cell Lysis and Improved Proinflammatory Cytokine Release of Bispecific Immunoligands via Nkp30 Engagement. <i>Journal of Immunology</i> , 2021, 206, 225-236.	0.8	32
64	Sunflower Trypsin Inhibitor 1 Derivatives as Molecular Scaffolds for the Development of Novel Peptidic Radiopharmaceuticals. <i>Molecular Imaging and Biology</i> , 2010, 12, 377-385.	2.6	31
65	Autotransporters with GDSL Passenger Domains: Molecular Physiology and Biotechnological Applications. <i>ChemBioChem</i> , 2011, 12, 1476-1485.	2.6	31
66	REAL-Select: Full-Length Antibody Display and Library Screening by Surface Capture on Yeast Cells. <i>PLoS ONE</i> , 2014, 9, e114887.	2.5	31
67	Spontaneous Isopeptide Bond Formation as a Powerful Tool for Engineering Site-Specific Antibody-Drug Conjugates. <i>Scientific Reports</i> , 2016, 6, 39291.	3.3	31
68	Isolation of a pH-Sensitive IgNAR Variable Domain from a Yeast-Displayed, Histidine-Doped Master Library. <i>Marine Biotechnology</i> , 2016, 18, 161-167.	2.4	31
69	A novel one-step approach for the construction of yeast surface display Fab antibody libraries. <i>Microbial Cell Factories</i> , 2018, 17, 3.	4.0	31
70	Engineering of ultraID, a compact and hyperactive enzyme for proximity-dependent biotinylation in living cells. <i>Communications Biology</i> , 2022, 5, .	4.4	31
71	Functional Cell-Surface Display of a Lipase-Specific Chaperone. <i>ChemBioChem</i> , 2007, 8, 55-60.	2.6	30
72	Ultrafast Single-Scan 2D-NMR Spectroscopic Detection of a PHIP-Hyperpolarized Protease Inhibitor. <i>Chemistry - A European Journal</i> , 2019, 25, 4025-4030.	3.3	30

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73	Yeast Surface Display in Combination with Fluorescence-activated Cell Sorting Enables the Rapid Isolation of Antibody Fragments Derived from Immunized Chickens. <i>Biotechnology Journal</i> , 2019, 14, 1800466.	3.5	30
74	PHIP-label: parahydrogen-induced polarization in propargylglycine-containing synthetic oligopeptides. <i>Chemical Communications</i> , 2013, 49, 7839.	4.1	29
75	Engineered cystine-knot miniproteins for diagnostic applications. <i>Expert Review of Molecular Diagnostics</i> , 2010, 10, 361-368.	3.1	28
76	Directed Evolution of a Bond-Forming Enzyme: Ultrahigh-Throughput Screening of Microbial Transglutaminase Using Yeast Surface Display. <i>Chemistry - A European Journal</i> , 2018, 24, 15195-15200.	3.3	28
77	Generation of Potent Anti-HER1/2 Immunotoxins by Protein Ligation Using Split Inteins. <i>ACS Chemical Biology</i> , 2018, 13, 2058-2066.	3.4	28
78	A Bioorthogonal Click Chemistry Toolbox for Targeted Synthesis of Branched and Well-Defined Protein-Protein Conjugates. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 12885-12893.	13.8	28
79	Sustainable Peptide Synthesis Enabled by a Transient Protecting Group. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 12984-12990.	13.8	28
80	Self-Assembled Hybrid Aptamer-Fc Conjugates for Targeted Delivery: A Modular Chemoenzymatic Approach. <i>ACS Chemical Biology</i> , 2015, 10, 2158-2165.	3.4	27
81	A simplified procedure for antibody engineering by yeast surface display: Coupling display levels and target binding by ribosomal skipping. <i>Biotechnology Journal</i> , 2017, 12, 1600454.	3.5	27
82	Microbial transglutaminase for biotechnological and biomedical engineering. <i>Biological Chemistry</i> , 2019, 400, 257-274.	2.5	27
83	FACS-Based Functional Protein Screening via Microfluidic Co-encapsulation of Yeast Secretor and Mammalian Reporter Cells. <i>Scientific Reports</i> , 2020, 10, 10182.	3.3	27
84	Oxidative Folding of Peptides with Cystine-Knot Architectures: Kinetic Studies and Optimization of Folding Conditions. <i>ChemBioChem</i> , 2013, 14, 137-146.	2.6	26
85	Preparation and kinetic performance assessment of thick film 10 μ m open tubular silica capillaries in normal phase high pressure liquid chromatography. <i>Journal of Chromatography A</i> , 2013, 1315, 127-134.	3.7	26
86	Engineering a Constrained Peptidic Scaffold towards Potent and Selective Furin Inhibitors. <i>ChemBioChem</i> , 2015, 16, 2441-2444.	2.6	26
87	Dual Function pH Responsive Bispecific Antibodies for Tumor Targeting and Antigen Depletion in Plasma. <i>Frontiers in Immunology</i> , 2019, 10, 1892.	4.8	26
88	Between two worlds: a comparative study on in vitro and in silico inhibition of trypsin and matriptase by redox-stable SFTI-1 variants at near physiological pH. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 7753.	2.8	25
89	Performance evaluation of thick film open tubular silica capillary by reversed phase liquid chromatography. <i>Journal of Chromatography A</i> , 2013, 1283, 110-115.	3.7	25
90	An Apoptosis-Inducing Peptidic Heptad That Efficiently Clusters Death Receptor-5. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 5085-5089.	13.8	25

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91	Facile generation of antibody heavy and light chain diversities for yeast surface display by Golden Gate Cloning. <i>Biological Chemistry</i> , 2019, 400, 383-393.	2.5	24
92	From cell line development to the formulated drug product: The art of manufacturing therapeutic monoclonal antibodies. <i>International Journal of Pharmaceutics</i> , 2021, 594, 120164.	5.2	24
93	From pico to nano: biofunctionalization of cube-octameric silsesquioxanes by peptides and miniproteins. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 6287.	2.8	23
94	Feasibility study of semi-selective protein precipitation with salt-tolerant copolymers for industrial purification of therapeutic antibodies. <i>Biotechnology and Bioengineering</i> , 2013, 110, 2915-2927.	3.3	23
95	Intein mediated high throughput screening for bispecific antibodies. <i>MAbs</i> , 2020, 12, 1731938.	5.2	23
96	Functional paper-based materials for diagnostics. <i>ChemTexts</i> , 2021, 7, 14.	1.9	23
97	Design of a Trispecific Checkpoint Inhibitor and Natural Killer Cell Engager Based on a 2 + 1 Common Light Chain Antibody Architecture. <i>Frontiers in Immunology</i> , 2021, 12, 669496.	4.8	23
98	Fmoc-Assisted Synthesis of a 29-Residue Cystine-Knot Trypsin Inhibitor Containing a Guaninyl Amino Acid at the P1-Position. <i>European Journal of Organic Chemistry</i> , 2004, 2004, 4931-4935.	2.4	22
99	DNA Libraries for the Construction of Phage Libraries: Statistical and Structural Requirements and Synthetic Methods. <i>Molecules</i> , 2011, 16, 1625-1641.	3.8	22
100	A tightly regulated and adjustable CRISPR-dCas9 based AND gate in yeast. <i>Nucleic Acids Research</i> , 2019, 47, 509-520.	14.5	22
101	ToxR co-operative interactions are not modulated by environmental conditions or periplasmic domain conformation. <i>Molecular Microbiology</i> , 1999, 31, 305-317.	2.5	20
102	Enhancing the Pharmacokinetics and Antitumor Activity of an Î±-Amanitin-Based Small-Molecule Drug Conjugate via Conjugation with an Fc Domain. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 4117-4129.	6.4	20
103	Characterisation of the barrier caused by lumenally secreted gastro-intestinal proteolytic enzymes for two novel cystine-knot microproteins. <i>Amino Acids</i> , 2008, 35, 195-200.	2.7	19
104	Application of copper(i) catalyzed azide-alkyne [3+2] cycloaddition to the synthesis of template-assembled multivalent peptide conjugates. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 4177.	2.8	19
105	Dextramabs: A Novel Format of Antibody-Drug Conjugates Featuring a Multivalent Polysaccharide Scaffold. <i>ChemistryOpen</i> , 2019, 8, 354-357.	1.9	19
106	Structure of the Dispase Autolysis-inducing Protein from <i>Streptomyces mobaraensis</i> and Glutamine Cross-linking Sites for Transglutaminase. <i>Journal of Biological Chemistry</i> , 2016, 291, 20417-20426.	3.4	18
107	Efficient Site-Specific Antibody-Drug Conjugation by Engineering a Nature-Derived Recognition Tag for Microbial Transglutaminase. <i>ChemBioChem</i> , 2019, 20, 2411-2419.	2.6	18
108	Immunoglobulin Mutant Library Genetically Screened for Folding Stability Exploiting Bacterial Signal Transduction. <i>Journal of Molecular Biology</i> , 1995, 251, 471-476.	4.2	17

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109	The Shark Strikes Twice: Hypervariable Loop 2 of Shark IgNAR Antibody Variable Domains and Its Potential to Function as an Autonomous Paratope. <i>Marine Biotechnology</i> , 2015, 17, 386-392.	2.4	17
110	Expeditious Generation of Biparatopic Common Light Chain Antibodies via Chicken Immunization and Yeast Display Screening. <i>Frontiers in Immunology</i> , 2020, 11, 606878.	4.8	17
111	Rapid Generation of Chicken Immune Libraries for Yeast Surface Display. <i>Methods in Molecular Biology</i> , 2020, 2070, 289-302.	0.9	17
112	Beyond bispecificity: Controlled Fab arm exchange for the generation of antibodies with multiple specificities. <i>MAbs</i> , 2022, 14, 2018960.	5.2	17
113	Alternative binding proteins get mature: Rivalling antibodies. <i>FEBS Journal</i> , 2008, 275, 2667-2667.	4.7	16
114	Aptamers Binding to c-Met Inhibiting Tumor Cell Migration. <i>PLoS ONE</i> , 2015, 10, e0142412.	2.5	16
115	Nanoscale Biodegradable Organic-Inorganic Hybrids for Efficient Cell Penetration and Drug Delivery. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 14842-14846.	13.8	16
116	TRAIL-Inspired Multivalent Dextran Conjugates Efficiently Induce Apoptosis upon DR5 Receptor Clustering. <i>ChemBioChem</i> , 2019, 20, 3006-3012.	2.6	16
117	Engineering therapeutic antibodies for patient safety: tackling the immunogenicity problem. <i>Protein Engineering, Design and Selection</i> , 2020, 33, .	2.1	16
118	Isolation of Common Light Chain Antibodies from Immunized Chickens Using Yeast Biopanning and Fluorescence-Activated Cell Sorting. <i>Biotechnology Journal</i> , 2021, 16, e2000240.	3.5	16
119	General mutagenesis/gene expression procedure for the construction of variant immunoglobulin domains in <i>Escherichia coli</i> . <i>Journal of Molecular Biology</i> , 1992, 228, 359-365.	4.2	15
120	Barnase Fusion as a Tool to Determine the Crystal Structure of the Small Disulfide-rich Protein McoEeTI. <i>Journal of Molecular Biology</i> , 2006, 356, 1-8.	4.2	15
121	Cube-octameric silsesquioxane-mediated cargo peptide delivery into living cancer cells. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 2258-2265.	2.8	15
122	Mid-Infrared spectroscopy-based antibody aggregate quantification in cell culture fluids. <i>Biotechnology Journal</i> , 2013, 8, 912-917.	3.5	15
123	Generation of Semi-Synthetic Shark IgNAR Single-Domain Antibody Libraries. <i>Methods in Molecular Biology</i> , 2018, 1701, 147-167.	0.9	15
124	Impact of Acetylated and Non-Acetylated Fucose Analogues on IgG Glycosylation. <i>Antibodies</i> , 2019, 8, 9.	2.5	14
125	Effektive Markierung von bioaktiven Peptiden mit PHIP-Markern zur Steigerung der Empfindlichkeit von NMR-Signalen. <i>Angewandte Chemie</i> , 2014, 126, 13155-13159.	2.0	13
126	Combination of inverse electron-demand Diels-Alder reaction with highly efficient oxime ligation expands the toolbox of site-selective peptide conjugations. <i>Chemical Communications</i> , 2015, 51, 11130-11133.	4.1	13

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127	A Streamlined Approach for the Construction of Large Yeast Surface Display Fab Antibody Libraries. <i>Methods in Molecular Biology</i> , 2018, 1827, 145-161.	0.9	13
128	Light-Controlled Chemoenzymatic Immobilization of Proteins towards Engineering of Bioactive Papers. <i>Chemistry - A European Journal</i> , 2019, 25, 1746-1751.	3.3	13
129	Effect of Conjugation Site and Technique on the Stability and Pharmacokinetics of Antibody-Drug Conjugates. <i>Journal of Pharmaceutical Sciences</i> , 2021, 110, 3776-3785.	3.3	13
130	Azobenzene switch with a long-lived cis-state to photocontrol the enzyme activity of a histone deacetylase-like amidohydrolase. <i>Biological Chemistry</i> , 2014, 395, 401-412.	2.5	12
131	Humanization of Chicken-Derived scFv Using Yeast Surface Display and NGS Data Mining. <i>Biotechnology Journal</i> , 2021, 16, e2000231.	3.5	12
132	Carbohydrate binding module-fused antibodies improve the performance of cellulose-based lateral flow immunoassays. <i>Scientific Reports</i> , 2021, 11, 7880.	3.3	12
133	PROLink™ Single Step Circularization and Purification Procedure for the Generation of an Improved Variant of Human Growth Hormone. <i>Bioconjugate Chemistry</i> , 2016, 27, 1341-1347.	3.6	11
134	Structure of a glutamine donor mimicking inhibitory peptide shaped by the catalytic cleft of microbial transglutaminase. <i>FEBS Journal</i> , 2018, 285, 4684-4694.	4.7	11
135	Treating Bladder Cancer: Engineering of Current and Next Generation Antibody-, Fusion Protein-, mRNA-, Cell- and Viral-Based Therapeutics. <i>Frontiers in Oncology</i> , 2021, 11, 672262.	2.8	11
136	Milking the Cow: Cattle-Derived Chimeric Ultralong CDR-H3 Antibodies and Their Engineered CDR-H3-Only Knobbody Counterparts Targeting Epidermal Growth Factor Receptor Elicit Potent NK Cell-Mediated Cytotoxicity. <i>Frontiers in Immunology</i> , 2021, 12, 742418.	4.8	11
137	Grabbing the Bull by Both Horns: Bovine Ultralong CDR-H3 Paratopes Enable Engineering of "Almost Natural"™ Common Light Chain Bispecific Antibodies Suitable For Effector Cell Redirection. <i>Frontiers in Immunology</i> , 2021, 12, 801368.	4.8	11
138	Isolation of pH-Sensitive Antibody Fragments by Fluorescence-Activated Cell Sorting and Yeast Surface Display. <i>Methods in Molecular Biology</i> , 2018, 1685, 311-331.	0.9	10
139	Selection of Antibodies with Tailored Properties by Application of High-Throughput Multiparameter Fluorescence-Activated Cell Sorting of Yeast-Displayed Immune Libraries. <i>Molecular Biotechnology</i> , 2018, 60, 727-735.	2.4	10
140	A Generic Procedure for the Isolation of pH- and Magnesium-Responsive Chicken scFvs for Downstream Purification of Human Antibodies. <i>Frontiers in Bioengineering and Biotechnology</i> , 2020, 8, 688.	4.1	10
141	A Bioorthogonal Click Chemistry Toolbox for Targeted Synthesis of Branched and Well-Defined Protein-Protein Conjugates. <i>Angewandte Chemie</i> , 2020, 132, 12985-12993.	2.0	10
142	Generation and Biological Evaluation of Fc Antigen Binding Fragment-Drug Conjugates as a Novel Antibody-Based Format for Targeted Drug Delivery. <i>Bioconjugate Chemistry</i> , 2021, 32, 1699-1710.	3.6	10
143	Native Llama Nanobody Library Panning Performed by Phage and Yeast Display Provides Binders Suitable for C-Reactive Protein Detection. <i>Biosensors</i> , 2021, 11, 496.	4.7	10
144	Matrix effects during monitoring of antibody and host cell proteins using attenuated total reflection spectroscopy. <i>Biotechnology Progress</i> , 2013, 29, 265-274.	2.6	9

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