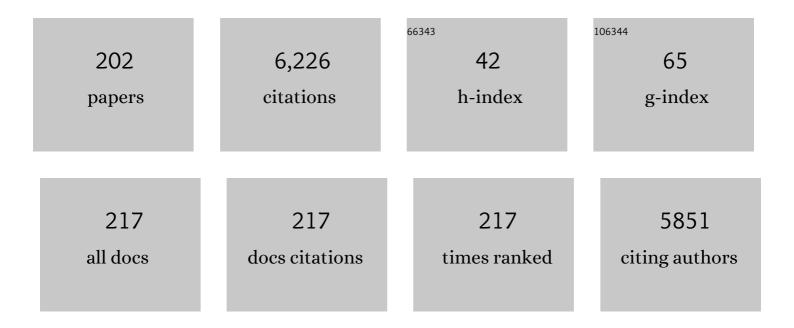
Harald Kolmar

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Beyond bispecificity: Controlled Fab arm exchange for the generation of antibodies with multiple specificities. MAbs, 2022, 14, 2018960.	5.2	17
2	Generation of a host cell line containing a <scp>MAR</scp> â€rich landing pad for siteâ€specific integration and expression of transgenes. Biotechnology Progress, 2022, 38, e3254.	2.6	3
3	Methoxy-Monobenzoylmethane Protects Skin from UV-Induced Damages in a Randomized, Placebo Controlled, Double-Blinded Human In Vivo Study and Prevents Signs of Inflammation While Improving the Skin Barrier. Dermatology and Therapy, 2022, 12, 435-449.	3.0	1
4	A Generic Strategy to Generate Bifunctional Two-in-One Antibodies by Chicken Immunization. Frontiers in Immunology, 2022, 13, 888838.	4.8	5
5	Humanization of Chicken-Derived Antibodies by Yeast Surface Display. Methods in Molecular Biology, 2022, 2491, 335-360.	0.9	2
6	Antibody Library Screening Using Yeast Biopanning and Fluorescence-Activated Cell Sorting. Methods in Molecular Biology, 2022, 2491, 177-193.	0.9	0
7	Streamlining the Transition From Yeast Surface Display of Antibody Fragment Immune Libraries to the Production as IgG Format in Mammalian Cells. Frontiers in Bioengineering and Biotechnology, 2022, 10, .	4.1	9
8	Engineering of ultraID, a compact and hyperactive enzyme for proximity-dependent biotinylation in living cells. Communications Biology, 2022, 5, .	4.4	31
9	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. Journal of Immunology, 2021, 206, 225-236.	0.8	32
10	From cell line development to the formulated drug product: The art of manufacturing therapeutic monoclonal antibodies. International Journal of Pharmaceutics, 2021, 594, 120164.	5.2	24
11	Humanization of Chickenâ€Derived scFv Using Yeast Surface Display and NGS Data Mining. Biotechnology Journal, 2021, 16, e2000231.	3.5	12
12	Isolation of Common Light Chain Antibodies from Immunized Chickens Using Yeast Biopanning and Fluorescenceâ€Activated Cell Sorting. Biotechnology Journal, 2021, 16, e2000240.	3.5	16
13	Review: High temperature short time treatment of cell culture media and feed solutions to mitigate adventitious viral contamination in the biopharmaceutical industry. Biotechnology Progress, 2021, 37, e3117.	2.6	2
14	Use of 5â€Thioâ€Lâ€Fucose to modulate binding affinity of therapeutic proteins. Biotechnology and Bioengineering, 2021, 118, 1818-1831.	3.3	6
15	Enhancing the Pharmacokinetics and Antitumor Activity of an Î \pm -Amanitin-Based Small-Molecule Drug Conjugate via Conjugation with an Fc Domain. Journal of Medicinal Chemistry, 2021, 64, 4117-4129.	6.4	20
16	Functional paper-based materials for diagnostics. ChemTexts, 2021, 7, 14.	1.9	23
17	Characterization of soy protein hydrolysates and influence of its iron content on monoclonal antibody production by a murine hybridoma cell line. Biotechnology Progress, 2021, 37, e3147.	2.6	2
18	Inside Front Cover Image, Volume 118, Number 5, May 2021. Biotechnology and Bioengineering, 2021, 118, ii.	3.3	0

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19	Carbohydrate binding module-fused antibodies improve the performance of cellulose-based lateral flow immunoassays. Scientific Reports, 2021, 11, 7880.	3.3	12
20	Comparison of Membrane Depth Determination Techniques for Active Ingredient Skin Penetration Studies Using Microdialysis. Skin Pharmacology and Physiology, 2021, 34, 203-213.	2.5	5
21	Design of a Trispecific Checkpoint Inhibitor and Natural Killer Cell Engager Based on a 2 + 1 Common Light Chain Antibody Architecture. Frontiers in Immunology, 2021, 12, 669496.	4.8	23
22	Recombinant Antibody Production Using a Dual-Promoter Single Plasmid System. Antibodies, 2021, 10, 18.	2.5	6
23	Treating Bladder Cancer: Engineering of Current and Next Generation Antibody-, Fusion Protein-, mRNA-, Cell- and Viral-Based Therapeutics. Frontiers in Oncology, 2021, 11, 672262.	2.8	11
24	Toward Fabrication of Bioactive Papers: Covalent Immobilization of Peptides and Proteins. Biomacromolecules, 2021, 22, 2954-2962.	5.4	7
25	Generation and Biological Evaluation of Fc Antigen Binding Fragment-Drug Conjugates as a Novel Antibody-Based Format for Targeted Drug Delivery. Bioconjugate Chemistry, 2021, 32, 1699-1710.	3.6	10
26	Synthetic Integrin-Targeting Dextran-Fc Hybrids Efficiently Inhibit Tumor Proliferation In Vitro. Frontiers in Chemistry, 2021, 9, 693097.	3.6	2
27	S-Sulfocysteine – Investigation of cellular uptake in CHO cells. Journal of Biotechnology, 2021, 335, 27-38.	3.8	1
28	Protease-Activation of Fc-Masked Therapeutic Antibodies to Alleviate Off-Tumor Cytotoxicity. Frontiers in Immunology, 2021, 12, 715719.	4.8	7
29	Effect of Conjugation Site and Technique on the Stability and Pharmacokinetics of Antibody-Drug Conjugates. Journal of Pharmaceutical Sciences, 2021, 110, 3776-3785.	3.3	13
30	EGFR binding Fc domain-drug conjugates: stable and highly potent cytotoxic molecules mediate selective cell killing. Biological Chemistry, 2021, .	2.5	0
31	Multivalent dextran hybrids for efficient cytosolic delivery of biomolecular cargoes. Journal of Peptide Science, 2021, 27, e3298.	1.4	5
32	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. Frontiers in Immunology, 2021, 12, 742418.	4.8	11
33	Grabbing the Bull by Both Horns: Bovine Ultralong CDR-H3 Paratopes Enable Engineering of â€~Almost Natural' Common Light Chain Bispecific Antibodies Suitable For Effector Cell Redirection. Frontiers in Immunology, 2021, 12, 801368.	4.8	11
34	Native llama Nanobody Library Panning Performed by Phage and Yeast Display Provides Binders Suitable for C-Reactive Protein Detection. Biosensors, 2021, 11, 496.	4.7	10
35	The metal-binding properties of the long chaplin from Streptomyces mobaraensis: A bioinformatic and biochemical approach. Journal of Inorganic Biochemistry, 2020, 202, 110878.	3.5	2
36	Specific Targeting of Lymphoma Cells Using Semisynthetic Anti-Idiotype Shark Antibodies. Frontiers in Immunology, 2020, 11, 560244.	4.8	7

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37	Engineering therapeutic antibodies for patient safety: tackling the immunogenicity problem. Protein Engineering, Design and Selection, 2020, 33, .	2.1	16
38	Expeditious Generation of Biparatopic Common Light Chain Antibodies via Chicken Immunization and Yeast Display Screening. Frontiers in Immunology, 2020, 11, 606878.	4.8	17
39	Sustainable Peptide Synthesis Enabled by a Transient Protecting Group. Angewandte Chemie, 2020, 132, 13084-13090.	2.0	2
40	Intein mediated high throughput screening for bispecific antibodies. MAbs, 2020, 12, 1731938.	5.2	23
41	FACS-Based Functional Protein Screening via Microfluidic Co-encapsulation of Yeast Secretor and Mammalian Reporter Cells. Scientific Reports, 2020, 10, 10182.	3.3	27
42	A Generic Procedure for the Isolation of pH- and Magnesium-Responsive Chicken scFvs for Downstream Purification of Human Antibodies. Frontiers in Bioengineering and Biotechnology, 2020, 8, 688.	4.1	10
43	Recent progress in transglutaminase-mediated assembly of antibody-drug conjugates. Analytical Biochemistry, 2020, 595, 113615.	2.4	38
44	Solvent-Containing Closure Material Can Be Used to Prevent Follicular Penetration of Caffeine and Fluorescein Sodium Salt on Porcine Ear Skin. Skin Pharmacology and Physiology, 2020, 33, 117-126.	2.5	4
45	A Bioorthogonal Click Chemistry Toolbox for Targeted Synthesis of Branched and Wellâ€Defined Protein–Protein Conjugates. Angewandte Chemie, 2020, 132, 12985-12993.	2.0	10
46	A Bioorthogonal Click Chemistry Toolbox for Targeted Synthesis of Branched and Wellâ€Đefined Protein–Protein Conjugates. Angewandte Chemie - International Edition, 2020, 59, 12885-12893.	13.8	28
47	Sustainable Peptide Synthesis Enabled by a Transient Protecting Group. Angewandte Chemie - International Edition, 2020, 59, 12984-12990.	13.8	28
48	Selection and Characterization of Anti-idiotypic Shark Antibody Domains. Methods in Molecular Biology, 2020, 2070, 191-209.	0.9	2
49	Rapid Generation of Chicken Immune Libraries for Yeast Surface Display. Methods in Molecular Biology, 2020, 2070, 289-302.	0.9	17
50	Isolation of Tailor-Made Antibody Fragments from Yeast-Displayed B-Cell Receptor Repertoires by Multiparameter Fluorescence-Activated Cell Sorting. Methods in Molecular Biology, 2020, 2070, 249-266.	0.9	0
51	Isolation of Anti-Hapten Antibodies by Fluorescence-Activated Cell Sorting of Yeast-Displayed B-Cell Receptor Gene Repertoires. Methods in Molecular Biology, 2020, 2070, 267-287.	0.9	0
52	Simplifying the Detection of Surface Presentation Levels in Yeast Surface Display by Intracellular tGFP Expression. Methods in Molecular Biology, 2020, 2070, 211-222.	0.9	3
53	Glutamine-walking: Creating reactive substrates for transglutaminase-mediated protein labeling. Methods in Enzymology, 2020, 644, 121-148.	1.0	3
54	Dissecting capture and twisting of aureolysin and pseudolysin: functional amino acids of the Dispase autolysis-inducing protein. Biochemical Journal, 2020, 477, 2595-2606.	3.7	1

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55	Dual Function pH Responsive Bispecific Antibodies for Tumor Targeting and Antigen Depletion in Plasma. Frontiers in Immunology, 2019, 10, 1892.	4.8	26
56	Ultrafast Singleâ€Scan 2Dâ€NMR Spectroscopic Detection of a PHIPâ€Hyperpolarized Protease Inhibitor. Chemistry - A European Journal, 2019, 25, 4025-4030.	3.3	30
57	Protein engineering comes of age. Biological Chemistry, 2019, 400, 255-256.	2.5	0
58	TRAILâ€Inspired Multivalent Dextran Conjugates Efficiently Induce Apoptosis upon DR5 Receptor Clustering. ChemBioChem, 2019, 20, 3006-3012.	2.6	16
59	Tailoring Activity and Selectivity of Microbial Transglutaminase. Methods in Molecular Biology, 2019, 2012, 151-169.	0.9	2
60	SpyLigase-Catalyzed Modification of Antibodies. Methods in Molecular Biology, 2019, 2012, 171-192.	0.9	2
61	Site-Specific Antibody–Drug Conjugation Using Microbial Transglutaminase. Methods in Molecular Biology, 2019, 2012, 135-149.	0.9	9
62	Biochemical study of sortase E2 fromStreptomycesÂmobaraensisand determination of transglutaminase crossâ€linking sites. FEBS Letters, 2019, 593, 1944-1956.	2.8	2
63	Efficient Siteâ€Specific Antibody–Drug Conjugation by Engineering a Natureâ€Derived Recognition Tag for Microbial Transglutaminase. ChemBioChem, 2019, 20, 2411-2419.	2.6	18
64	Dextramabs: A Novel Format of Antibodyâ€Đrug Conjugates Featuring a Multivalent Polysaccharide Scaffold. ChemistryOpen, 2019, 8, 354-357.	1.9	19
65	Facile generation of antibody heavy and light chain diversities for yeast surface display by Golden Gate Cloning. Biological Chemistry, 2019, 400, 383-393.	2.5	24
66	Impact of Acetylated and Non-Acetylated Fucose Analogues on IgG Glycosylation. Antibodies, 2019, 8, 9.	2.5	14
67	Yeast Surface Display in Combination with Fluorescenceâ€activated Cell Sorting Enables the Rapid Isolation of Antibody Fragments Derived from Immunized Chickens. Biotechnology Journal, 2019, 14, 1800466.	3.5	30
68	Lightâ€Controlled Chemoenzymatic Immobilization of Proteins towards Engineering of Bioactive Papers. Chemistry - A European Journal, 2019, 25, 1746-1751.	3.3	13
69	A tightly regulated and adjustable CRISPR-dCas9 based AND gate in yeast. Nucleic Acids Research, 2019, 47, 509-520.	14.5	22
70	Microbial transglutaminase for biotechnological and biomedical engineering. Biological Chemistry, 2019, 400, 257-274.	2.5	27
71	Site-Specific Conjugation of Thiol-Reactive Cytotoxic Agents to Nonnative Cysteines of Engineered Monoclonal Antibodies. Methods in Molecular Biology, 2019, 2033, 1-14.	0.9	5
72	A novel one-step approach for the construction of yeast surface display Fab antibody libraries. Microbial Cell Factories, 2018, 17, 3.	4.0	31

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73	Shark attack: Haiantikörper für Biomedizin und Biotechnologie. BioSpektrum, 2018, 24, 142-145.	0.0	Ο
74	Isolation of pH-Sensitive Antibody Fragments by Fluorescence-Activated Cell Sorting and Yeast Surface Display. Methods in Molecular Biology, 2018, 1685, 311-331.	0.9	10
75	Generation of Semi-Synthetic Shark IgNAR Single-Domain Antibody Libraries. Methods in Molecular Biology, 2018, 1701, 147-167.	0.9	15
76	Structure of a glutamine donor mimicking inhibitory peptide shaped by the catalytic cleft of microbial transglutaminase. FEBS Journal, 2018, 285, 4684-4694.	4.7	11
77	A Streamlined Approach for the Construction of Large Yeast Surface Display Fab Antibody Libraries. Methods in Molecular Biology, 2018, 1827, 145-161.	0.9	13
78	Construction of Histidine-Enriched Shark IgNAR Variable Domain Antibody Libraries for the Isolation of pH-Sensitive vNAR Fragments. Methods in Molecular Biology, 2018, 1827, 109-127.	0.9	3
79	Destructive twisting of neutral metalloproteases: the catalysis mechanism of the Dispase autolysisâ€inducing protein fromStreptomyces mobaraensisDSM40487. FEBS Journal, 2018, 285, 4246-4264.	4.7	7
80	Selection of Antibodies with Tailored Properties by Application of High-Throughput Multiparameter Fluorescence-Activated Cell Sorting of Yeast-Displayed Immune Libraries. Molecular Biotechnology, 2018, 60, 727-735.	2.4	10
81	Engineering IgG-Like Bispecific Antibodies—An Overview. Antibodies, 2018, 7, 28.	2.5	37
82	Directed Evolution of a Bondâ€Forming Enzyme: Ultrahighâ€Throughput Screening of Microbial Transglutaminase Using Yeast Surface Display. Chemistry - A European Journal, 2018, 24, 15195-15200.	3.3	28
83	Covalent Attachment of Enzymes to Paper Fibers for Paper-Based Analytical Devices. Frontiers in Chemistry, 2018, 6, 214.	3.6	35
84	Beyond antibody engineering: directed evolution of alternative binding scaffolds and enzymes using yeast surface display. Microbial Cell Factories, 2018, 17, 32.	4.0	58
85	Generation of Potent Anti-HER1/2 Immunotoxins by Protein Ligation Using Split Inteins. ACS Chemical Biology, 2018, 13, 2058-2066.	3.4	28
86	Highlight issue: protein design. Biological Chemistry, 2017, 398, 1-2.	2.5	2
87	Generation of human bispecific common light chain antibodies by combining animal immunization and yeast display. Protein Engineering, Design and Selection, 2017, 30, 291-301.	2.1	33
88	Engineering bispecific antibodies with defined chain pairing. New Biotechnology, 2017, 39, 167-173.	4.4	43
89	Semi-synthetic vNAR libraries screened against therapeutic antibodies primarily deliver anti-idiotypic binders. Scientific Reports, 2017, 7, 9676.	3.3	34
90	Camelid and shark single domain antibodies: structural features and therapeutic potential. Current Opinion in Structural Biology, 2017, 45, 10-16.	5.7	165

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91	A simplified procedure for antibody engineering by yeast surface display: Coupling display levels and target binding by ribosomal skipping. Biotechnology Journal, 2017, 12, 1600454.	3.5	27
92	An Apoptosisâ€Inducing Peptidic Heptad That Efficiently Clusters Death Receptorâ€5. Angewandte Chemie - International Edition, 2016, 55, 5085-5089.	13.8	25
93	Nanoskalige, biologisch abbaubare organischâ€anorganische Hybride für effiziente Zellaufnahme und Wirkstofftransport. Angewandte Chemie, 2016, 128, 15063-15068.	2.0	0
94	Spontaneous Isopeptide Bond Formation as a Powerful Tool for Engineering Site-Specific Antibody-Drug Conjugates. Scientific Reports, 2016, 6, 39291.	3.3	31
95	PROLink—Single Step Circularization and Purification Procedure for the Generation of an Improved Variant of Human Growth Hormone. Bioconjugate Chemistry, 2016, 27, 1341-1347.	3.6	11
96	Structure of the Dispase Autolysis-inducing Protein from Streptomyces mobaraensis and Glutamine Cross-linking Sites for Transglutaminase. Journal of Biological Chemistry, 2016, 291, 20417-20426.	3.4	18
97	Balancing Selectivity and Efficacy of Bispecific Epidermal Growth Factor Receptor (EGFR) × c-MET Antibodies and Antibody-Drug Conjugates. Journal of Biological Chemistry, 2016, 291, 25106-25119.	3.4	66
98	Nanoscale Biodegradable Organic–Inorganic Hybrids for Efficient Cell Penetration and Drug Delivery. Angewandte Chemie - International Edition, 2016, 55, 14842-14846.	13.8	16
99	Ein Apoptoseâ€induzierendes Heptamer, das effizient den Todesrezeptorâ€5 bündelt. Angewandte Chemie, 2016, 128, 5169-5173.	2.0	2
100	Coupled reactions on bioparticles: Stereoselective reduction with cofactor regeneration on PhaC inclusion bodies. Biotechnology Journal, 2016, 11, 890-898.	3.5	7
101	Isolation of a pH-Sensitive IgNAR Variable Domain from a Yeast-Displayed, Histidine-Doped Master Library. Marine Biotechnology, 2016, 18, 161-167.	2.4	31
102	Single-domain antibodies for biomedical applications. Immunopharmacology and Immunotoxicology, 2016, 38, 21-28.	2.4	64
103	Engineering a Constrained Peptidic Scaffold towards Potent and Selective Furin Inhibitors. ChemBioChem, 2015, 16, 2441-2444.	2.6	26
104	Locked by Design: A Conformationally Constrained Transglutaminase Tag Enables Efficient Site‧pecific Conjugation. Angewandte Chemie - International Edition, 2015, 54, 13420-13424.	13.8	49
105	Cystineâ€knot peptides targeting cancerâ€relevant human cytotoxic T lymphocyteâ€associated antigen 4 (CTLAâ€4). Journal of Peptide Science, 2015, 21, 651-660.	1.4	32
106	Aptamers Binding to c-Met Inhibiting Tumor Cell Migration. PLoS ONE, 2015, 10, e0142412.	2.5	16
107	Combination of inverse electron-demand Diels–Alder reaction with highly efficient oxime ligation expands the toolbox of site-selective peptide conjugations. Chemical Communications, 2015, 51, 11130-11133.	4.1	13
108	At-line mid infrared spectroscopy for monitoring downstream processing unit operations. Process Biochemistry, 2015, 50, 997-1005.	3.7	32

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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. Marine Biotechnology, 2015, 17, 386-392.	2.4	17
110	Bacterial Secretion Systems for Use in Biotechnology: Autotransporter-Based Cell Surface Display and Ultrahigh-Throughput Screening of Large Protein Libraries. Springer Protocols, 2015, , 87-103.	0.3	0
111	Structural insights and biomedical potential of IgNAR scaffolds from sharks. MAbs, 2015, 7, 15-25.	5.2	102
112	A generic approach to engineer antibody pH-switches using combinatorial histidine scanning libraries and yeast display. MAbs, 2015, 7, 138-151.	5.2	64
113	Self-Assembled Hybrid Aptamer-Fc Conjugates for Targeted Delivery: A Modular Chemoenzymatic Approach. ACS Chemical Biology, 2015, 10, 2158-2165.	3.4	27
114	REAL-Select: Full-Length Antibody Display and Library Screening by Surface Capture on Yeast Cells. PLoS ONE, 2014, 9, e114887.	2.5	31
115	Feasibility of polyelectrolyte-driven Fab fragment separation. Biotechnology Journal, 2014, 9, 698-701.	3.5	0
116	A general strategy for antibody library screening via conversion of transient target binding into permanent reporter deposition. Protein Engineering, Design and Selection, 2014, 27, 41-47.	2.1	4
117	Potent inhibitors of human matriptaseâ€1 based on the scaffold of sunflower trypsin inhibitor. Journal of Peptide Science, 2014, 20, 415-420.	1.4	42
118	Protein Production in Yarrowia lipolytica Via Fusion to the Secreted Lipase Lip2p. Molecular Biotechnology, 2014, 56, 79-90.	2.4	7
119	Required polymer lengths per precipitated protein molecule in protein-polymer interaction. Journal of Polymer Research, 2014, 21, 1.	2.4	2
120	Azobenzene switch with a long-lived cis-state to photocontrol the enzyme activity of a histone deacetylase-like amidohydrolase. Biological Chemistry, 2014, 395, 401-412.	2.5	12
121	A Chemoenzymatic Approach to Protein Immobilization onto Crystalline Cellulose Nanoscaffolds. Angewandte Chemie - International Edition, 2014, 53, 12618-12623.	13.8	48
122	Effective PHIP Labeling of Bioactive Peptides Boosts the Intensity of the NMR Signal. Angewandte Chemie - International Edition, 2014, 53, 12941-12945.	13.8	34
123	Therapeutic antibody engineering by high efficiency cell screening. FEBS Letters, 2014, 588, 278-287.	2.8	95
124	Shark Attack: High affinity binding proteins derived from shark vNAR domains by stepwise in vitro affinity maturation. Journal of Biotechnology, 2014, 191, 236-245.	3.8	74
125	Effektive Markierung von bioaktiven Peptiden mit PHIPâ€Markern zur Steigerung der Empfindlichkeit von NMR‧ignalen. Angewandte Chemie, 2014, 126, 13155-13159.	2.0	13
126	Fragmentation Follows Structure: Top-Down Mass Spectrometry Elucidates the Topology of Engineered Cystine-Knot Miniproteins. PLoS ONE, 2014, 9, e108626.	2.5	5

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127	Polyelectrolyte–protein interaction at low ionic strength: required chain flexibility depending on protein average charge. Colloid and Polymer Science, 2013, 291, 1759-1769.	2.1	5
128	Feasibility study of semiâ€selective protein precipitation with saltâ€tolerant copolymers for industrial purification of therapeutic antibodies. Biotechnology and Bioengineering, 2013, 110, 2915-2927.	3.3	23
129	PHIP-label: parahydrogen-induced polarization in propargylglycine-containing synthetic oligopeptides. Chemical Communications, 2013, 49, 7839.	4.1	29
130	Oxidative Folding of Peptides with Cystineâ€Knot Architectures: Kinetic Studies and Optimization of Folding Conditions. ChemBioChem, 2013, 14, 137-146.	2.6	26
131	Preparation and kinetic performance assessment of thick film 10–20μ4m open tubular silica capillaries in normal phase high pressure liquid chromatography. Journal of Chromatography A, 2013, 1315, 127-134.	3.7	26
132	Arranged Sevenfold: Structural Insights into the C-Terminal Oligomerization Domain of Human C4b-Binding Protein. Journal of Molecular Biology, 2013, 425, 1302-1317.	4.2	69
133	Performance evaluation of thick film open tubular silica capillary by reversed phase liquid chromatography. Journal of Chromatography A, 2013, 1283, 110-115.	3.7	25
134	Structural characterization of <i>Spinacia oleracea</i> trypsin inhibitor III (SOTI-III). Acta Crystallographica Section D: Biological Crystallography, 2013, 69, 114-120.	2.5	4
135	Combinatorial tuning of peptidic drug candidates: high-affinity matriptase inhibitors through incremental structure-guided optimization. Organic and Biomolecular Chemistry, 2013, 11, 1848.	2.8	48
136	Bioconjugation on cube-octameric silsesquioxanes. Organic and Biomolecular Chemistry, 2013, 11, 2224.	2.8	44
137	Cube-octameric silsesquioxane-mediated cargo peptide delivery into living cancer cells. Organic and Biomolecular Chemistry, 2013, 11, 2258-2265.	2.8	15
138	Host cell protein quantification by fourier transform mid infrared spectroscopy (FTâ€MIR). Biotechnology and Bioengineering, 2013, 110, 252-259.	3.3	34
139	Matrix effects during monitoring of antibody and host cell proteins using attenuated total reflection spectroscopy. Biotechnology Progress, 2013, 29, 265-274.	2.6	9
140	Customization of copolymers to optimize selectivity and yield in polymerâ€driven antibody purification processes. Biotechnology Progress, 2013, 29, 1484-1493.	2.6	2
141	Midâ€infrared spectroscopyâ€based antibody aggregate quantification in cell culture fluids. Biotechnology Journal, 2013, 8, 912-917.	3.5	15
142	DegP Protease. , 2013, , 2567-2571.		0
143	Combinatorial Optimization of Cystine-Knot Peptides towards High-Affinity Inhibitors of Human Matriptase-1. PLoS ONE, 2013, 8, e76956.	2.5	55
144	From pico to nano: biofunctionalization of cube-octameric silsesquioxanes by peptides and miniproteins. Organic and Biomolecular Chemistry, 2012, 10, 6287.	2.8	23

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145	Synthesis and characterization of new generation open tubular silica capillaries for liquid chromatography. Journal of Chromatography A, 2012, 1265, 88-94.	3.7	40
146	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. Organic and Biomolecular Chemistry, 2012, 10, 7753.	2.8	25
147	A sensitive method for rapid detection of alkyl halides and dehalogenase activity using a multistep enzyme assay. AMB Express, 2012, 2, 51.	3.0	3
148	Chemical Synthesis, Backbone Cyclization and Oxidative Folding of Cystine-knot Peptides — Promising Scaffolds for Applications in Drug Design. Molecules, 2012, 17, 12533-12552.	3.8	43
149	Braces for the Peptide Backbone: Insights into Structure–Activity Relationships of Protease Inhibitor Mimics with Locked Amide Conformations. Angewandte Chemie - International Edition, 2012, 51, 3708-3712.	13.8	62
150	DNA Libraries for the Construction of Phage Libraries: Statistical and Structural Requirements and Synthetic Methods. Molecules, 2011, 16, 1625-1641.	3.8	22
151	Decorating microbes: surface display of proteins on Escherichia coli. Trends in Biotechnology, 2011, 29, 79-86.	9.3	198
152	"Triazole Bridge†Disulfideâ€Bond Replacement by Rutheniumâ€Catalyzed Formation of 1,5â€Disubstituted 1,2,3â€Triazoles. Angewandte Chemie - International Edition, 2011, 50, 5207-5211.	13.8	112
153	Autotransporters with GDSL Passenger Domains: Molecular Physiology and Biotechnological Applications. ChemBioChem, 2011, 12, 1476-1485.	2.6	31
154	Natural and Engineered Cystine Knot Miniproteins for Diagnostic and Therapeutic Applications. Current Pharmaceutical Design, 2011, 17, 4329-4336.	1.9	41
155	Bi-specific Aptamers Mediating Tumor Cell Lysis. Journal of Biological Chemistry, 2011, 286, 21896-21905.	3.4	124
156	Sunflower Trypsin Inhibitor 1 Derivatives as Molecular Scaffolds for the Development of Novel Peptidic Radiopharmaceuticals. Molecular Imaging and Biology, 2010, 12, 377-385.	2.6	31
157	In Vivo Enzyme Immobilization by Inclusion Body Display. Applied and Environmental Microbiology, 2010, 76, 5563-5569.	3.1	40
158	Engineered Cystine Knot Miniproteins as Potent Inhibitors of Human Mast Cell Tryptase β. Journal of Molecular Biology, 2010, 395, 167-175.	4.2	92
159	Towards click bioconjugations on cube-octameric silsesquioxane scaffolds. Organic and Biomolecular Chemistry, 2010, 8, 2212.	2.8	49
160	Engineered cystine-knot miniproteins for diagnostic applications. Expert Review of Molecular Diagnostics, 2010, 10, 361-368.	3.1	28
161	Biological diversity and therapeutic potential of natural and engineered cystine knot miniproteins. Current Opinion in Pharmacology, 2009, 9, 608-614.	3.5	108
162	Application of copper(i) catalyzed azide–alkyne [3+2] cycloaddition to the synthesis of template-assembled multivalent peptide conjugates. Organic and Biomolecular Chemistry, 2009, 7, 4177.	2.8	19

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163	Microbodiesâ"¢. Methods in Molecular Biology, 2009, 535, 361-372.	0.9	2
164	Characterisation of the barrier caused by luminally secreted gastro-intestinal proteolytic enzymes for two novel cystine-knot microproteins. Amino Acids, 2008, 35, 195-200.	2.7	19
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