

Albert Breier

List of Publications by Year in descending order

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120
papers

1,822
citations

331642

21
h-index

330122

37
g-index

120
all docs

120
docs citations

120
times ranked

2067
citing authors

#	ARTICLE	IF	CITATIONS
1	Different mechanisms of drug resistance to hypomethylating agents in the treatment of myelodysplastic syndromes and acute myeloid leukemia. <i>Drug Resistance Updates</i> , 2022, 61, 100805.	14.4	17
2	The Roles of microRNAs in Cancer Multidrug Resistance. <i>Cancers</i> , 2022, 14, 1090.	3.7	22
3	Sulphoraphane Affinity-Based Chromatography for the Purification of Myrosinase from <i>Lepidium sativum</i> Seeds. <i>Biomolecules</i> , 2022, 12, 406.	4.0	5
4	Changes in Apoptotic Pathways in MOLM-13 Cell Lines after Induction of Resistance to Hypomethylating Agents. <i>International Journal of Molecular Sciences</i> , 2021, 22, 2076.	4.1	5
5	Optimisation of Recombinant Myrosinase Production in <i>Pichia pastoris</i> . <i>International Journal of Molecular Sciences</i> , 2021, 22, 3677.	4.1	11
6	Insight into Bortezomib Focusing on Its Efficacy against P-gp-Positive MDR Leukemia Cells. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5504.	4.1	2
7	Development of Multidrug Resistance in Acute Myeloid Leukemia Is Associated with Alterations of the LPHN1/GAL-9/TIM-3 Signaling Pathway. <i>Cancers</i> , 2021, 13, 3629.	3.7	9
8	Cell Death Effects Induced by Sulforaphane and Allyl Isothiocyanate on P-Glycoprotein Positive and Negative Variants in L1210 Cells. <i>Molecules</i> , 2020, 25, 2093.	3.8	7
9	Development of Resistance to Endoplasmic Reticulum Stress-Inducing Agents in Mouse Leukemic L1210 Cells. <i>Molecules</i> , 2020, 25, 2517.	3.8	6
10	Overexpression of GRP78/BiP in P-Glycoprotein-Positive L1210 Cells is Responsible for Altered Response of Cells to Tunicamycin as a Stressor of the Endoplasmic Reticulum. <i>Cells</i> , 2020, 9, 890.	4.1	9
11	Screening of Phenanthroquinolizidine Alkaloid Derivatives for Inducing Cell Death of L1210 Leukemia Cells with Negative and Positive P-glycoprotein Expression. <i>Molecules</i> , 2019, 24, 2127.	3.8	5
12	Overexpression of the ABCB1 drug transporter in acute myeloid leukemia cells is associated with downregulation of latrophilin-1. <i>General Physiology and Biophysics</i> , 2018, 37, 353-357.	0.9	5
13	Interplay between P-Glycoprotein Expression and Resistance to Endoplasmic Reticulum Stressors. <i>Molecules</i> , 2018, 23, 337.	3.8	32
14	Triorganotin Derivatives Induce Cell Death Effects on L1210 Leukemia Cells at Submicromolar Concentrations Independently of P-glycoprotein Expression. <i>Molecules</i> , 2018, 23, 1053.	3.8	8
15	Detection of the Mitochondrial Membrane Potential by the Cationic Dye JC-1 in L1210 Cells with Massive Overexpression of the Plasma Membrane ABCB1 Drug Transporter. <i>International Journal of Molecular Sciences</i> , 2018, 19, 1985.	4.1	100
16	L1210 Cells Overexpressing ABCB1 Drug Transporters Are Resistant to Inhibitors of the N- and O-glycosylation of Proteins. <i>Molecules</i> , 2017, 22, 1104.	3.8	6
17	The expression of P-gp in leukemia cells is associated with cross-resistance to protein N-glycosylation inhibitor tunicamycin. <i>General Physiology and Biophysics</i> , 2016, 35, 497-510.	0.9	12
18	The expression of P-glycoprotein in leukemia cells is associated with the upregulated expression of nestin, a class 6 filament protein. <i>Leukemia Research</i> , 2016, 48, 32-39.	0.8	8

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19	A decrease in cellular microRNA-27a content is involved in azacytidine-induced P-glycoprotein expression in SKM-1 cells. <i>Toxicology in Vitro</i> , 2016, 36, 81-88.	2.4	6
20	Acute myeloid leukemia cells MOLM-13 and SKM-1 established for resistance by azacytidine are crossresistant to P-glycoprotein substrates. <i>Toxicology in Vitro</i> , 2015, 29, 1405-1415.	2.4	16
21	Selection of resistant acute myeloid leukemia SKM-1 and MOLM-13 cells by vincristine-, mitoxantrone- and lenalidomide-induced upregulation of P-glycoprotein activity and downregulation of CD33 cell surface exposure. <i>European Journal of Pharmaceutical Sciences</i> , 2015, 77, 29-39.	4.0	17
22	Molecular changes induced by repeated restraint stress in the heart: the effect of oxytocin receptor antagonist atosiban. <i>Canadian Journal of Physiology and Pharmacology</i> , 2015, 93, 827-834.	1.4	6
23	Reduced UDP-glucose Levels Are Associated with P-glycoprotein Over-expression in L1210 Cells and Limit Glucosylceramide Synthase Activity. <i>Anticancer Research</i> , 2015, 35, 2627-34.	1.1	5
24	Lenalidomide treatment induced the normalization of marker protein levels in blood plasma of patients with 5q-myelodysplastic syndrome. <i>General Physiology and Biophysics</i> , 2015, 34, 399-406.	0.9	3
25	Vincristine-induced expression of P-glycoprotein in MOLM-13 and SKM-1 acute myeloid leukemia cell lines is associated with coexpression of nestin transcript. <i>General Physiology and Biophysics</i> , 2014, 33, 425-431.	0.9	9
26	Effect of 9-cis retinoic acid and all-trans retinoic acid in combination with verapamil on P-glycoprotein expression in L1210 cells. <i>Neoplasma</i> , 2014, 62, 553-565.	1.6	15
27	Lectin detection of cell surface saccharides remodeling induced by development of P-glycoprotein mediated multidrug resistance phenotype in L1210 leukemia cells. <i>Acta Chimica Slovaca</i> , 2014, 7, 52-56.	0.8	2
28	New Insight into P-Glycoprotein as a Drug Target. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2013, 13, 159-170.	1.7	135
29	New insight into p-glycoprotein as a drug target. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2013, 13, 159-70.	1.7	65
30	Detection of Glycomic Alterations Induced by Overexpression of P-Glycoprotein on the Surfaces of L1210 Cells Using Sialic Acid Binding Lectins. <i>International Journal of Molecular Sciences</i> , 2012, 13, 15177-15192.	4.1	11
31	Potential of Anticancer Drugs: Effects of Pentoxifylline on Neoplastic Cells. <i>International Journal of Molecular Sciences</i> , 2012, 13, 369-382.	4.1	26
32	P-glycoprotein depresses cisplatin sensitivity in L1210 cells by inhibiting cisplatin-induced caspase-3 activation. <i>Toxicology in Vitro</i> , 2012, 26, 435-444.	2.4	55
33	New Insight into P-Glycoprotein as a Drug Target. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2012, 13, 159-170.	1.7	10
34	β_2 integrin as regulator of metastatic potential. <i>Acta Pharmacologica Sinica</i> , 2011, 32, 279-279.	6.1	0
35	Tunicamycin Depresses P-Glycoprotein Glycosylation Without an Effect on Its Membrane Localization and Drug Efflux Activity in L1210 Cells. <i>International Journal of Molecular Sciences</i> , 2011, 12, 7772-7784.	4.1	33
36	Effect of thapsigargin on P-glycoprotein-negative and P-glycoprotein-positive L1210 mouse leukaemia cells. <i>General Physiology and Biophysics</i> , 2010, 29, 396-401.	0.9	8

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37	Why the xanthine derivatives are used to study of P-glycoprotein-mediated multidrug resistance in L1210/VCR line cells. <i>General Physiology and Biophysics</i> , 2010, 29, 215-221.	0.9	0
38	Acute treatment with polyphenol quercetin improves postischemic recovery of isolated perfused rat hearts after global ischemia. <i>Canadian Journal of Physiology and Pharmacology</i> , 2010, 88, 465-471.	1.4	36
39	The presence of P-glycoprotein in L1210 cells directly induces down-regulation of cell surface saccharide targets of concanavalin A. <i>Anticancer Research</i> , 2010, 30, 3661-8.	1.1	21
40	Effect of quercetin on kinetic properties of renal Na,K-ATPase in normotensive and hypertensive rats. <i>Journal of Physiology and Pharmacology</i> , 2010, 61, 593-8.	1.1	11
41	Vincristine-Induced Overexpression of P-Glycoprotein in L1210 Cells Is Associated with Remodeling of Cell Surface Saccharides. <i>Journal of Proteome Research</i> , 2009, 8, 513-520.	3.7	20
42	Multidrug resistant P-glycoprotein positive L1210/VCR cells are also cross-resistant to cisplatin via a mechanism distinct from P-glycoprotein-mediated drug efflux activity. <i>General Physiology and Biophysics</i> , 2009, 28, 391-403.	0.9	13
43	Membrane transport and apoptosis-related proteins in radiation-associated acute myeloid leukemia following the Chernobyl accident. <i>General Physiology and Biophysics</i> , 2009, 28, 63-69.	0.9	0
44	Does any relationship exist between P-glycoprotein-mediated multidrug resistance and intracellular calcium homeostasis. <i>General Physiology and Biophysics</i> , 2009, 28 Spec No Focus, F89-95.	0.9	25
45	H ₂ S and HS ⁻ donor NaHS releases nitric oxide from nitrosothiols, metal nitrosyl complex, brain homogenate and murine L1210 leukaemia cells. <i>Pflugers Archiv European Journal of Physiology</i> , 2008, 457, 271-279.	2.8	77
46	Combined treatment of P-gp-positive L1210/VCR cells by verapamil and all-trans retinoic acid induces down-regulation of P-glycoprotein expression and transport activity. <i>Toxicology in Vitro</i> , 2008, 22, 96-105.	2.4	19
47	Overexpression of P-glycoprotein in L1210/VCR cells is associated with changes in several endoplasmic reticulum proteins that may be partially responsible for the lack of thapsigargin sensitivity. <i>General Physiology and Biophysics</i> , 2008, 27, 211-21.	0.9	10
48	Modulation of expression of Na ⁺ /Ca ²⁺ -exchanger in heart of rat and mouse under stress. <i>Acta Physiologica</i> , 2007, 190, 127-136.	3.8	15
49	Inhibitory effect of DIDS, NPPB, and phloretin on intracellular chloride channels. <i>Pflugers Archiv European Journal of Physiology</i> , 2007, 455, 349-357.	2.8	38
50	L1210 cells cultivated under the selection pressure of doxorubicin or vincristine express common mechanisms of multidrug resistance based on the overexpression of P-glycoprotein. <i>Toxicology in Vitro</i> , 2006, 20, 1560-1568.	2.4	14
51	LY294,002, a specific inhibitor of PI3K/Akt kinase pathway, antagonizes P-glycoprotein-mediated multidrug resistance. <i>European Journal of Pharmaceutical Sciences</i> , 2006, 29, 426-434.	4.0	75
52	Structural differences between sensitive and resistant L1210 cells. <i>General Physiology and Biophysics</i> , 2006, 25, 427-38.	0.9	4
53	Inhibition of (Na ⁺)/K ⁺)-ATPase by Cibacron Blue 3G-A and its analogues. <i>General Physiology and Biophysics</i> , 2006, 25, 439-53.	0.9	4
54	P-Glycoprotein - Implications of Metabolism of Neoplastic Cells and Cancer Therapy. <i>Current Cancer Drug Targets</i> , 2005, 5, 457-468.	1.6	105

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55	Expression of P-glycoprotein in L1210 cells is linked with rise in sensitivity to Ca ²⁺ . <i>Biochemical and Biophysical Research Communications</i> , 2005, 335, 777-784.	2.1	18
56	Prolongation of pentoxifylline aliphatic side chain positively affects the reversal of P-glycoprotein-mediated multidrug resistance in L1210/VCR line cells. <i>General Physiology and Biophysics</i> , 2005, 24, 461-6.	0.9	1
57	Reversal of P-glycoprotein mediated vincristine resistance of L1210/VCR cells by analogues of pentoxifylline. <i>European Journal of Pharmaceutical Sciences</i> , 2004, 21, 283-293.	4.0	26
58	Functional fluo-3/AM assay on P-glycoprotein transport activity in L1210/VCR cells by confocal microscopy. <i>General Physiology and Biophysics</i> , 2004, 23, 357-66.	0.9	13
59	P-glycoprotein-mediated multidrug resistance phenotype of L1210/VCR cells is associated with decreases of oligo- and/or polysaccharide contents. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2003, 1639, 213-224.	3.8	16
60	Hypoxia increases cell death in multidrug-resistant leukemia cells. Differences in viability and ultrastructure between sensitive and multidrug-resistant L1210 mouse leukemic cells under hypoxia. <i>General Physiology and Biophysics</i> , 2003, 22, 265-73.	0.9	2
61	Proteins released from liver after ischaemia induced an elevation of heart resistance against ischaemia-reperfusion injury: 1. Beneficial effect of protein fraction isolated from perfusate after ischaemia and reperfusion of liver. <i>General Physiology and Biophysics</i> , 2003, 22, 567-77.	0.9	4
62	Pentoxifylline influences drug transport activity of P-glycoprotein and decreases mdrl gene expression in multidrug resistant mouse leukemic L1210/VCR cells. <i>General Physiology and Biophysics</i> , 2002, 21, 103-9.	0.9	5
63	Carbonyl group of aliphatic side chain of pentoxifylline does not play role for P-glycoprotein antagonizing effect of pentoxifylline. <i>General Physiology and Biophysics</i> , 2002, 21, 471-8.	0.9	1
64	SB203580, a specific inhibitor of p38-MAPK pathway, is a new reversal agent of P-glycoprotein-mediated multidrug resistance. <i>European Journal of Pharmaceutical Sciences</i> , 2001, 14, 29-36.	4.0	122
65	Drug transporters and their role in multidrug resistance of neoplastic cells. <i>General Physiology and Biophysics</i> , 2001, 20, 215-37.	0.9	25
66	Reversal effect of specific inhibitors of extracellular-signal regulated protein kinase pathway on P-glycoprotein mediated vincristine resistance of L1210 cells. <i>General Physiology and Biophysics</i> , 2001, 20, 439-44.	0.9	23
67	Cytotoxic activity of several unrelated drugs on L1210 mouse leukemic cell sublines with P-glycoprotein (PGP) mediated multidrug resistance (MDR) phenotype. A QSAR study. <i>Neoplasma</i> , 2000, 47, 100-6.	1.6	11
68	Glutathione S-transferase does not play a role in multidrug resistance of L1210/VCR cell line. <i>Physiological Research</i> , 2000, 49, 447-53.	0.9	11
69	"Lysine is the Lord", thought some scientists in regard to the group interacting with fluorescein isothiocyanate in ATP-binding sites of P-type ATPases but, is it not cysteine?. <i>General Physiology and Biophysics</i> , 2000, 19, 253-63.	0.9	10
70	Differential expression of regulatory proteins in L1210/VCR cells with multidrug resistance mediated by P-glycoprotein. <i>General Physiology and Biophysics</i> , 1999, 18, 45-56.	0.9	4
71	Interaction of lactate dehydrogenase with anthraquinone dyes: characterization of ligands for dye-ligand chromatography. <i>Biomedical Applications</i> , 1998, 715, 273-281.	1.7	18
72	Ca(2+)-induced inhibition of sodium pump: noncompetitive inhibition in respect of magnesium and sodium cations. <i>General Physiology and Biophysics</i> , 1998, 17, 179-88.	0.9	7

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73	Direct interaction between verapamil and doxorubicin causes the lack of reversal effect of verapamil on P-glycoprotein mediated resistance to doxorubicin in vitro using L1210/VCR cells. <i>Neoplasma</i> , 1998, 45, 248-53.	1.6	5
74	Mechanisms that may be involved in calcium tolerance of the diabetic heart. <i>Molecular and Cellular Biochemistry</i> , 1997, 176, 191-198.	3.1	31
75	Mechanisms that may be involved in calcium tolerance of the diabetic heart. , 1997, , 191-198.		13
76	Diabetic cardiomyopathy in rats: biochemical mechanisms of increased tolerance to calcium overload. <i>Diabetes Research and Clinical Practice</i> , 1996, 31, S93-S103.	2.8	25
77	Is cysteine residue important in FITC-sensitive ATP-binding site of P-type ATPases? A commentary to the state of the art. <i>Molecular and Cellular Biochemistry</i> , 1996, 160-161, 89-93.	3.1	6
78	Prevention by 7-oxo-prostacyclin of the calcium paradox in rat heart: Role of the sarcolemmal (Na,K)-ATPase. <i>Molecular and Cellular Biochemistry</i> , 1996, 160-161, 257-263.	3.1	5
79	Is cysteine residue important in FITC-sensitive ATP-binding site of P-type ATPases? A commentary to the state of the art. , 1996, , 89-93.		0
80	Overcoming of P-glycoprotein mediated vincristine resistance of L1210/VCR mouse leukemic cells could be induced by pentoxifyline but not by theophylline and caffeine. <i>Neoplasma</i> , 1996, 43, 11-5.	1.6	5
81	The membrane effect of benfluron: modulation of the heart sarcolemmal (Na ⁺ , K ⁺)-ATPase and Mg(2 ⁺)-ATPase activities. <i>General Physiology and Biophysics</i> , 1996, 15, 71-5.	0.9	1
82	Competitive inhibition of (Na/K)-ATPase by furylethylenes with respect to potassium ions. <i>General Physiology and Biophysics</i> , 1996, 15, 291-307.	0.9	0
83	Adaptation of the heart to ischemia by preconditioning: Effects on energy equilibrium, properties of sarcolemmal ATPases and release of cardioprotective proteins. <i>Molecular and Cellular Biochemistry</i> , 1995, 147, 129-137.	3.1	6
84	Inhibition of (Na/K)-ATPase by electrophilic substances: Functional implications. <i>Molecular and Cellular Biochemistry</i> , 1995, 147, 187-192.	3.1	10
85	The effects of calcium and calcium channel blockers on sodium pump. <i>Pflugers Archiv European Journal of Physiology</i> , 1995, 429, 716-721.	2.8	13
86	Screening of Binding Properties of Con-A Immobilized on Bead Cellulose by Flow Microcalorimetry Using Invertase and Anti-Con-A Antibody as Reporting Systems. <i>Analytical Letters</i> , 1995, 28, 2585-2594.	1.8	3
87	Inhibition of (Na/K)-ATPase by electrophilic substances: Functional implications. , 1995, , 187-192.		1
88	Distribution of proteins in aqueous two-phase systems formed by dextran and polyethylene glycol. Influence of protein hydrophobicity. <i>General Physiology and Biophysics</i> , 1995, 14, 277-91.	0.9	1
89	Effect of phorbol myristate acetate (PMA) on P-glycoprotein mediated vincristine resistance of L1210 cells. <i>General Physiology and Biophysics</i> , 1995, 14, 171-5.	0.9	2
90	Estimation of the effective hydrophobicity of protiens: Re-evaluation of methods. <i>Biotechnology Letters</i> , 1994, 8, 915-920.	0.5	1

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91	2,4,6-trinitrobenzenesulfonic acid modification of the carboxyl-terminal region (C-domain) of calreticulin. <i>Molecular and Cellular Biochemistry</i> , 1994, 130, 19-28.	3.1	5
92	Inhibition of (Na/K)-ATPase by NFE induces an increase in mechanical activity of perfused guinea-pig heart. <i>General Physiology and Biophysics</i> , 1994, 13, 433-41.	0.9	1
93	Effect of pentoxifylline on P-glycoprotein mediated vincristine resistance of L1210 mouse leukemic cell line. <i>Neoplasma</i> , 1994, 41, 297-303.	1.6	4
94	Time dependence of [3H]-vincristine accumulation by L1210 mouse leukemic cells. Effect of P-glycoprotein overexpression. <i>General Physiology and Biophysics</i> , 1994, 13, 287-98.	0.9	5
95	Characterization of morphological and histochemical changes induced by overexpression of P-glycoprotein in mouse leukemic cell line L1210. <i>Neoplasma</i> , 1994, 41, 83-8.	1.6	7
96	Reversal effects of several Ca(2+)-entry blockers, neuroleptics and local anaesthetics on P-glycoprotein-mediated vincristine resistance of L1210/VCR mouse leukaemic cell line. <i>Drugs Under Experimental and Clinical Research</i> , 1994, 20, 13-8.	0.3	4
97	Purification of Glycerol Kinase by "Dye-Ligand" Chromatography and Hydrophobic Interaction Chromatography on Bead-Cellulose Derivatives. <i>Collection of Czechoslovak Chemical Communications</i> , 1993, 58, 445-451.	1.0	7
98	Overcoming of vincristine resistance in L1210/VCR cells by several corticosteroids. Collateral sensitivity of resistant cells. <i>Neoplasma</i> , 1993, 40, 21-5.	1.6	5
99	Adaptation of mouse leukemia cells L1210 to vincristine. Evidence for expression of P-glycoprotein. <i>Neoplasma</i> , 1992, 39, 73-7.	1.6	22
100	Application of adsorption kinetics for estimation of dissociation constants. <i>Journal of Proteomics</i> , 1991, 22, 185-193.	2.4	4
101	Trypsin entrapped within liposomes. Partition of a low-molecular-mass substrate as the main factor in kinetic control of hydrolysis. <i>Collection of Czechoslovak Chemical Communications</i> , 1991, 56, 712-717.	1.0	2
102	Hydrophobic partitioning of proteins in a two-phase aqueous system of poly(oxyethylene)-dextran alternatively derivatized by 2-hydroxy-3-phenoxypropyl group. <i>Collection of Czechoslovak Chemical Communications</i> , 1991, 56, 1270-1278.	1.0	4
103	Increased activity of sarcolemmal (Na+K+)-ATPase is involved in the late cardioprotective action of 7-oxo-prostacyclin. <i>Cardioscience</i> , 1991, 2, 105-8.	0.5	17
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109	Principles of selectivity of sodium and potassium binding sites of the Na ⁺ /K ⁺ -ATPase. A corollary hypothesis. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1988, 946, 129-134.	2.6	2
110	Application of a time-concentration model of adsorption for determination of the nature of adsorbent-adsorbate interaction. <i>Colloid and Polymer Science</i> , 1987, 265, 933-937.	2.1	7
111	Size-exclusion effect of a substrate upon kinetics of trypsin immobilized on porous bead cellulose. 2. Influence of hydrodynamic diameter of substrate. <i>Enzyme and Microbial Technology</i> , 1987, 9, 44-46.	3.2	7
112	Interaction of ATP with the active sites of ATPases in heart sarcolemma. Role of the hydroxylic group in position two on the ribose moiety. <i>General Physiology and Biophysics</i> , 1987, 6, 193-6.	0.9	1
113	Some properties of the active site and cation binding site of the heart sarcolemmal (Na ⁺ + K ⁺)-ATPase. <i>Biomedica Biochimica Acta</i> , 1987, 46, S553-6.	0.1	3
114	Size-exclusion effect of a substrate upon kinetics of trypsin immobilized on porous bead cellulose. 1. Influence of distribution coefficient of a substrate. <i>Enzyme and Microbial Technology</i> , 1986, 8, 109-114.	3.2	16
115	Quantitative criterion for evaluation of hydrophobic sorbents. <i>Biomedical Applications</i> , 1986, 376, 95-101.	1.7	5
116	Effect of the concentration of 5,5'-dithiobis(2-nitrobenzoic acid) on parameters of the kinetics of its chemisorption on thiol derivatives of cellulose. <i>Collection of Czechoslovak Chemical Communications</i> , 1986, 51, 545-552.	1.0	6
117	Selective chemisorbents. Part 3: Selective binding of thiols to benzaldehyde derivatives of cellulose. <i>Reactive Polymers, Ion Exchangers, Sorbents</i> , 1984, 2, 189-196.	0.0	4
118	Simple estimation of carrier binding capacity using sorption kinetics curve-fitting. <i>Journal of Proteomics</i> , 1984, 9, 267-275.	2.4	10
119	Aldehydic derivatives of bead cellulose?relationships between matrix structure and function in immobilization of enzymes catalyzing hydrolysis of high molecular substrates. <i>Biotechnology and Bioengineering</i> , 1982, 24, 2573-2582.	3.3	24
120	Different Mechanisms of Drug Resistance in Myelodysplastic Syndromes and Acute Myeloid Leukemia. , O, , .		1