## Charles M Deber

## List of Publications by Year in descending order

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papers citations h-index

165

docs citations

h-index g-index

165
7974
times ranked citing authors

82

#	Article	IF	CITATIONS
1	Detergent binding explains anomalous SDS-PAGE migration of membrane proteins. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 1760-1765.	7.1	659
2	Non-random Distribution of Amino Acids in the Transmembrane Segments of Human Type I Single Span Membrane Proteins. Journal of Molecular Biology, 1993, 229, 602-608.	4.2	361
3	Roles of Hydrophobicity and Charge Distribution of Cationic Antimicrobial Peptides in Peptide-Membrane Interactions. Journal of Biological Chemistry, 2012, 287, 7738-7745.	3.4	317
4	Proline residues in transmembrane helixes: structural or dynamic role?. Biochemistry, 1991, 30, 8919-8923.	2.5	263
5	Basis for Selectivity of Cationic Antimicrobial Peptides for Bacterial Versus Mammalian Membranes. Journal of Biological Chemistry, 2005, 280, 33960-33967.	3.4	244
6	Why cyclic peptides? Complementary approaches to conformations. Accounts of Chemical Research, 1976, 9, 106-113.	15.6	205
7	Cationic Hydrophobic Peptides with Antimicrobial Activity. Antimicrobial Agents and Chemotherapy, 2002, 46, 3585-3590.	3.2	194
8	The structure of "unstructured―regions in peptides and proteins: Role of the polyproline II helix in protein folding and recognition*. Biopolymers, 2005, 80, 179-185.	2.4	181
9	A measure of helical propensity for amino acids in membrane environments. Nature Structural and Molecular Biology, 1994, 1, 368-373.	8.2	171
10	TM Finder: A prediction program for transmembrane protein segments using a combination of hydrophobicity and nonpolar phase helicity scales. Protein Science, 2001, 10, 212-219.	7.6	128
11	Anionic Phospholipids Modulate Peptide Insertion into Membranes. Biochemistry, 1997, 36, 5476-5482.	2.5	115
12	Transmembrane Aromatic Amino Acid Distribution in P-glycoprotein. Journal of Molecular Biology, 1994, 235, 554-564.	4.2	114
13	Peptides in membranes: Helicity and hydrophobicity. Biopolymers, 1995, 37, 295-318.	2.4	108
14	Guidelines for membrane protein engineering derived from de novo designed model peptides., 1998, 47, 41-62.		105
15	The Affinity of GXXXG Motifs in Transmembrane Helix-Helix Interactions Is Modulated by Long-range Communication. Journal of Biological Chemistry, 2004, 279, 16591-16597.	3.4	103
16	Central nervous system myelin: structure, function, and pathology. Clinical Biochemistry, 1991, 24, 113-134.	1.9	101
17	SDS Micelles as a Membrane-Mimetic Environment for Transmembrane Segments. Biochemistry, 2009, 48, 12096-12103.	2.5	101
18	Cyclic peptides. I. Cyclo(tri-L-prolyl) and derivatives. Synthesis and molecular conformation from nuclear magnetic resonance. Journal of the American Chemical Society, 1971, 93, 4893-4897.	13.7	95

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19	Missense mutations in transmembrane domains of proteins: Phenotypic propensity of polar residues for human disease. Proteins: Structure, Function and Bioinformatics, 2004, 54, 648-656.	2.6	94
20	Glycine and $\hat{l}^2$ -branched residues support and modulate peptide helicity in membrane environments. FEBS Letters, 1992, 311, 217-220.	2.8	92
21	Aromatic and Cationâ^'Ï€ Interactions Enhance Helixâ^'Helix Association in a Membrane Environment. Biochemistry, 2007, 46, 9208-9214.	2.5	92
22	Cyclic peptides. 17. Metal and amino acid complexes of cyclo(Pro-Gly)4 and analogs studied by nuclear magnetic resonance and circular dichroism. Journal of the American Chemical Society, 1977, 99, 4788-4798.	13.7	91
23	Interhelical hydrogen bonds in the CFTR membrane domain. Nature Structural Biology, 2001, 8, 597-601.	9.7	90
24	Method to generate highly stable D-amino acid analogs of bioactive helical peptides using a mirror image of the entire PDB. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 1505-1510.	7.1	89
25	Retention of Native-like Oligomerization States in Transmembrane Segment Peptides:  Application to the Escherichia coli Aspartate Receptor. Biochemistry, 2001, 40, 11106-11113.	2.5	88
26	Helix Induction in Antimicrobial Peptides by Alginate in Biofilms. Journal of Biological Chemistry, 2004, 279, 38749-38754.	3.4	88
27	Polar residue tagging of transmembrane peptides. Biopolymers, 2003, 71, 675-685.	2.4	86
28	Uncoupling Hydrophobicity and Helicity in Transmembrane Segments. Journal of Biological Chemistry, 1998, 273, 23645-23648.	3.4	85
29	Packing of Coat Protein Amphipathic and Transmembrane Helices in Filamentous Bacteriophage M13: Role of Small Residues in Protein Oligomeriza tion. Journal of Molecular Biology, 1995, 252, 6-14.	4.2	81
30	Membrane interactions of designed cationic antimicrobial peptides: The two thresholds. Biopolymers, 2008, 89, 360-371.	2.4	78
31	Bromo-A23187: A nonfluorescent calcium ionophore for use with fluorescent probes. Analytical Biochemistry, 1985, 146, 349-352.	2.4	77
32	Membrane protein misassembly in disease. Biochimica Et Biophysica Acta - Biomembranes, 2012, 1818, 1115-1122.	2.6	75
33	Disease-associated mutations in conserved residues of ALK-1 kinase domain. European Journal of Human Genetics, 2003, 11, 279-287.	2.8	<b>7</b> 3
34	Sequence Context Strongly Modulates Association of Polar Residues in Transmembrane Helices. Journal of Molecular Biology, 2003, 331, 255-262.	4.2	72
35	Peptides in membranes: Lipid-induced secondary structure of substance P. Biopolymers, 1987, 26, \$109-\$121.	2.4	70
36	Conformations of proline residues in membrane environments. Biopolymers, 1990, 29, 149-157.	2.4	68

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37	Threshold hydrophobicity dictates helical conformations of peptides in membrane environments., 1998, 39, 465-470.		68
38	Cyclic peptides. V. Proton and carbon-13 nuclear magnetic resonance determination of the preferred .beta. conformation for proline-containing cyclic hexapeptides. Journal of the American Chemical Society, 1973, 95, 258-260.	13.7	65
39	Transmembrane domain mediated self-assembly of major coat protein subunits from Ff bacteriophage11Edited by G. von Heijne. Journal of Molecular Biology, 2002, 315, 63-72.	4.2	64
40	Misfolding of the Cystic Fibrosis Transmembrane Conductance Regulator and Disease. Biochemistry, 2008, 47, 1465-1473.	2.5	62
41	Amino acid composition of the membrane and aqueous domains of integral membrane proteins. Archives of Biochemistry and Biophysics, 1986, 251, 68-76.	3.0	60
42	Therapeutic design of peptide modulators of protein-protein interactions in membranes. Biochimica Et Biophysica Acta - Biomembranes, 2017, 1859, 577-585.	2.6	57
43	Folding proteins into membranes. Nature Structural and Molecular Biology, 1996, 3, 815-818.	8.2	56
44	Conformational Aspects of Polypeptides. VII. Reversal of The Helical Sense of Poly-L-Aspartate Esters. Journal of the American Chemical Society, 1962, 84, 3773-3774.	13.7	55
45	Cyclic peptides. VIII. Carbon-13 and proton nuclear magnetic resonance evidence for slow cis'-trans' rotation in a cyclic tetrapeptide. Journal of the American Chemical Society, 1974, 96, 4015-4017.	13.7	54
46	Polar Residues in Membrane Domains of Proteins: Molecular Basis for Helixâ^'Helix Association in a Mutant CFTR Transmembrane Segmentâ€. Biochemistry, 2002, 41, 3647-3653.	2.5	53
47	Efficiency of detergents at maintaining membrane protein structures in their biologically relevant forms. Biochimica Et Biophysica Acta - Biomembranes, 2012, 1818, 1351-1358.	2.6	51
48	Acrylamide concentration determines the direction and magnitude of helical membrane protein gel shifts. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 15668-15673.	7.1	50
49	Polar mutations in membrane proteins as a biophysical basis for disease. Biopolymers, 2002, 66, 350-358.	2.4	49
50	Transfer of peptide hormones from aqueous to membrane phases. Biopolymers, 1985, 24, 105-116.	2.4	48
51	Oligomerization of a Peptide Derived from the Transmembrane Region of the Sodium Pump $\hat{I}^3$ Subunit: Effect of the Pathological Mutation G41R. Journal of Molecular Biology, 2002, 322, 583-590.	4.2	48
52	Cyclic peptides. Amino acid-cyclic peptides complexes. Journal of the American Chemical Society, 1974, 96, 7566-7568.	13.7	47
53	Optimizing synthesis and expression of transmembrane peptides and proteins. Methods, 2007, 41, 370-380.	3.8	46
54	The Structure and Function of Central Nervous System Myelin. Critical Reviews in Clinical Laboratory Sciences, 1993, 30, 29-64.	6.1	44

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55	Activity of novel non-amphipathic cationic antimicrobial peptides against Candida species. Journal of Antimicrobial Chemotherapy, 2006, 57, 899-907.	3.0	43
56	Positive Charge Patterning and Hydrophobicity of Membrane-Active Antimicrobial Peptides as Determinants of Activity, Toxicity, and Pharmacokinetic Stability. Journal of Medicinal Chemistry, 2019, 62, 6276-6286.	6.4	43
57	NMR investigation of the charge isomers of bovine myelin basic protein. Archives of Biochemistry and Biophysics, 1984, 233, 151-160.	3.0	42
58	Peptide Models of Membrane Protein Folding. Biochemistry, 2009, 48, 3036-3045.	2.5	42
59	Activity of a novel antimicrobial peptide against Pseudomonas aeruginosa biofilms. Scientific Reports, 2018, 8, 14728.	3.3	42
60	Influence of glycine residues on peptide conformation in membran environments. International Journal of Peptide and Protein Research, 1992, 40, 243-248.	0.1	40
61	Putting the β-breaks on membrane protein misfolding. Nature Structural Biology, 2002, 9, 318-319.	9.7	39
62	A Transmembrane Segment Mimic Derived from Escherichia coli Diacylglycerol Kinase Inhibits Protein Activity. Journal of Biological Chemistry, 2003, 278, 22056-22060.	3.4	38
63	Modulation of Na,K-ATPase by the Î <sup>3</sup> Subunit. Journal of Biological Chemistry, 2003, 278, 40437-40441.	3.4	38
64	Potential-sensitive membrane association of a fluorescent dye. FEBS Letters, 1987, 224, 337-342.	2.8	37
65	A lipid vesicle system for probing voltage-dependent peptide-lipid interactions: Application to alamethicin channel formation. Biopolymers, 1989, 28, 267-272.	2.4	36
66	Minimum energy conformations of proline-containing helices. Biopolymers, 1992, 32, 399-406.	2.4	36
67	Hydrophobic Helical Hairpins:  Design and Packing Interactions in Membrane Environments. Biochemistry, 2004, 43, 14361-14369.	2.5	36
68	Transmembrane Domain of Myelin Protein Zero Can Form Dimers:  Possible Implications for Myelin Construction. Biochemistry, 2007, 46, 12164-12173.	2.5	36
69	Correction factors for membrane protein molecular weight readouts on sodium dodecyl sulfate–polyacrylamide gel electrophoresis. Analytical Biochemistry, 2013, 434, 67-72.	2.4	34
70	Structure of the EmrEmultidrug transporter and its use for inhibitor peptide design. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E7932-E7941.	7.1	34
71	Peptide Mimics of the M13 Coat Protein Transmembrane Segment. Journal of Biological Chemistry, 2000, 275, 16155-16159.	3.4	32
72	Hydrophobic Interactions in Complexes of Antimicrobial Peptides with Bacterial Polysaccharides. Chemical Biology and Drug Design, 2007, 69, 405-412.	3.2	32

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73	Influence of hydrocarbon-stapling on membrane interactions of synthetic antimicrobial peptides. Bioorganic and Medicinal Chemistry, 2018, 26, 1189-1196.	3.0	32
74	Anti-Infectives Restore ORKAMBI® Rescue of F508del-CFTR Function in Human Bronchial Epithelial Cells Infected with Clinical Strains of P. aeruginosa. Biomolecules, 2020, 10, 334.	4.0	32
75	Combining hydrophobicity and helicity: a novel approach to membrane protein structure prediction. Bioorganic and Medicinal Chemistry, 1999, 7, 1-7.	3.0	31
76	Hydrophobicity and helicity of membrane-interactive peptides containing peptoid residues. Biopolymers, 2002, 65, 254-262.	2.4	31
77	Evidence for Assembly of Small Multidrug Resistance Proteins by a "Two-faced―Transmembrane Helix. Journal of Biological Chemistry, 2006, 281, 15546-15553.	3.4	31
78	Expression and Purification of Two Hydrophobic Double-Spanning Membrane Proteins Derived from the Cystic Fibrosis Transmembrane Conductance Regulator. Protein Expression and Purification, 2002, 25, 81-86.	1.3	30
79	Non-Native Interhelical Hydrogen Bonds in the Cystic Fibrosis Transmembrane Conductance Regulator Domain Modulated by Polar Mutations. Biochemistry, 2004, 43, 8077-8083.	2.5	30
80	A novel method for monitoring the cytosolic delivery of peptide cargo. Journal of Controlled Release, 2009, 137, 2-7.	9.9	30
81	Interhelical Packing in Detergent Micelles. Journal of Biological Chemistry, 2002, 277, 6067-6072.	3.4	29
82	Blockade of G Protein-Coupled Receptors and the Dopamine Transporter by a Transmembrane Domain Peptide: Novel Strategy for Functional Inhibition of Membrane Proteins in Vivo. Journal of Pharmacology and Experimental Therapeutics, 2003, 307, 481-489.	2.5	29
83	Inside-out Signaling Promotes Dynamic Changes in the Carcinoembryonic Antigen-related Cellular Adhesion Molecule 1 (CEACAM1) Oligomeric State to Control Its Cell Adhesion Properties. Journal of Biological Chemistry, 2013, 288, 29654-29669.	3.4	29
84	Lipid Solvation Effects Contribute to the Affinity of Gly-xxx-Gly Motif-Mediated Helixâ^'Helix Interactionsâ€. Biochemistry, 2006, 45, 8507-8515.	2.5	28
85	Peptides as transmembrane segments: Decrypting the determinants for helix–helix interactions in membrane proteins. Biopolymers, 2007, 88, 217-232.	2.4	28
86	Congenital Heart Block Maternal Sera Autoantibodies Target an Extracellular Epitope on the $\hat{l}\pm 1G$ T-Type Calcium Channel in Human Fetal Hearts. PLoS ONE, 2013, 8, e72668.	2.5	28
87	Conformations of neurotensin in solution and in membrane environments studied by 2â€D NMR spectroscopy. International Journal of Peptide and Protein Research, 1991, 37, 528-535.	0.1	27
88	The Assembly Motif of a Bacterial Small Multidrug Resistance Protein. Journal of Biological Chemistry, 2009, 284, 9870-9875.	3.4	26
89	Protein Structure in Membrane Domains. Annual Review of Biophysics, 2012, 41, 135-155.	10.0	26
90	Efflux by Small Multidrug Resistance Proteins Is Inhibited by Membrane-interactive Helix-stapled Peptides. Journal of Biological Chemistry, 2015, 290, 1752-1759.	3.4	26

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91	A minimal helical-hairpin motif provides molecular-level insights into misfolding and pharmacological rescue of CFTR. Communications Biology, 2018, 1, 154.	4.4	25
92	The position of the Gly-xxx-Gly motif in transmembrane segments modulates dimer affinityThis paper is one of a selection of papers published in this Special Issue, entitled CSBMCB — Membrane Proteins in Health and Disease Biochemistry and Cell Biology, 2006, 84, 1006-1012.	2.0	23
93	Self-association of the Transmembrane Domain of an Anthrax Toxin Receptor. Journal of Molecular Biology, 2006, 360, 145-156.	4.2	23
94	Destabilization of the Transmembrane Domain Induces Misfolding in a Phenotypic Mutant of Cystic Fibrosis Transmembrane Conductance Regulator. Journal of Biological Chemistry, 2005, 280, 4968-4974.	3.4	22
95	Cystic fibrosis transmembrane conductance regulator: expression and helicity of a double membrane-spanning segment. FEBS Letters, 1998, 431, 29-33.	2.8	21
96	Role of the Extracellular Loop in the Folding of a CFTR Transmembrane Helical Hairpin. Biochemistry, 2007, 46, 7099-7106.	2.5	20
97	Binding of human normal and multiple sclerosis-derived myelin basic protein to phospholipid vesicles: Effects on membrane head group and bilayer regions. Archives of Biochemistry and Biophysics, 1986, 245, 455-463.	3.0	19
98	Conformation of proline residues in bacteriorhodopsin. Biochemical and Biophysical Research Communications, 1990, 172, 862-869.	2.1	19
99	Functional Rescue of DeltaF508-CFTR by Peptides Designed to Mimic Sorting Motifs. Chemistry and Biology, 2009, 16, 520-530.	6.0	19
100	Positions of Polar Amino Acids Alter Interactions between Transmembrane Segments and Detergents. Biochemistry, 2011, 50, 3928-3935.	2.5	19
101	Drug Efflux by a Small Multidrug Resistance Protein Is Inhibited by a Transmembrane Peptide. Antimicrobial Agents and Chemotherapy, 2012, 56, 3911-3916.	3.2	19
102	Peptide-Based Efflux Pump Inhibitors of the Small Multidrug Resistance Protein from Pseudomonas aeruginosa. Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	19
103	Modulation of the Oligomerization of Myelin Proteolipid Protein by Transmembrane Helix Interaction Motifs. Biochemistry, 2010, 49, 6896-6902.	2.5	18
104	Complexation of Zn(II) to a native sequence tripeptide of human serum albumin studied by 13C nuclear magnetic resonance. Canadian Journal of Chemistry, 1980, 58, 757-766.	1.1	17
105	Aqueous solubility and membrane interactions of hydrophobic peptides with peptoid tags. Biopolymers, 2004, 76, 110-118.	2.4	17
106	CFTR transmembrane segments are impaired in their conformational adaptability by a pathogenic loop mutation and dynamically stabilized by Lumacaftor. Journal of Biological Chemistry, 2020, 295, 1985-1991.	3.4	17
107	Isosteric metal complexes of ionophore A23187. FEBS Letters, 1979, 105, 360-364.	2.8	16
108	Manipulation of peptide conformations by fine-tuning of the environment and/or the primary sequence. Biopolymers, 1995, 35, 667-675.	2.4	16

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109	Solubilization of Hydrophobic Peptides by Reversible Cysteine PEGylation. Biochemical and Biophysical Research Communications, 1998, 245, 618-621.	2.1	16
110	Peptide models for protein-mediated cation transport. Canadian Journal of Biochemistry, 1980, 58, 865-870.	1.4	15
111	Calcium transport by ionophorous peptides in dog and human lymphocytes detected by quin-2 fluorescence. Biochemical and Biophysical Research Communications, 1986, 134, 731-735.	2.1	15
112	Evidence for similar function of transmembrane segments in receptor and membrane-anchored proteins. Biopolymers, 1988, 27, 1171-1182.	2.4	15
113	Surface recognition elements of membrane protein oligomerization. Proteins: Structure, Function and Bioinformatics, 2008, 70, 786-793.	2.6	15
114	The N Terminus of the Qcr7 Protein of the Cytochromebc 1 Complex Is Not Essential for Import into Mitochondria in Saccharomyces cerevisiae but Is Essential for Assembly of the Complex. Journal of Biological Chemistry, 1997, 272, 17495-17501.	3.4	14
115	Membrane interactions of the hydrophobic segment of diacylglycerol kinase epsilon. Biochimica Et Biophysica Acta - Biomembranes, 2007, 1768, 2549-2558.	2.6	14
116	Membrane protein assembly patterns reflect selection for non-proliferative structures. FEBS Letters, 2007, 581, 1335-1341.	2.8	13
117	Distinctions between Hydrophobic Helices in Globular Proteins and Transmembrane Segments as Factors in Protein Sorting. Journal of Biological Chemistry, 2009, 284, 5395-5402.	3.4	13
118	Loop Sequence Dictates the Secondary Structure of a Human Membrane Protein Hairpin. Biochemistry, 2013, 52, 2419-2426.	2.5	13
119	D-glucose binding increases secondary structure of human erythrocyte monosaccharide transport protein. Biochemical and Biophysical Research Communications, 1987, 145, 1087-1091.	2.1	12
120	Conformations of cyclic peptide/calcium complexes in solution. Biopolymers, 1982, 21, 169-179.	2.4	11
121	Novel Hydrophobic Standards for Membrane Protein Molecular Weight Determinations via Sodium Dodecyl Sulfateâ°'Polyacrylamide Gel Electrophoresis. Biochemistry, 2010, 49, 10589-10591.	2.5	11
122	Deletion of a terminal residue disrupts oligomerization of a transmembrane α-helixThis paper is one of a selection of papers published in this special issue entitled "Canadian Society of Biochemistry, Molecular & amp; Cellular Biology 52nd Annual Meeting — Protein Folding: Principles and Diseases― and has undergone the Journal's usual peer review process Biochemistry and Cell Biology, 2010, 88,	2.0	11
123	339-345. Uncoupling Amphipathicity and Hydrophobicity: Role of Charge Clustering in Membrane Interactions of Cationic Antimicrobial Peptides. Biochemistry, 2021, 60, 2586-2592.	2.5	11
124	Microheterogeneity of bovine myelin basic protein studied by nuclear magnetic resonance spectroscopy. Biopolymers, 1983, 22, 377-380.	2.4	10
125	Beta-branched residues adjacent to GG4 motifs promote the efficient association of glycophorin a transmembrane helices. Biopolymers, 2011, 96, 340-347.	2.4	10
126	Modulation of Substrate Efflux in Bacterial Small Multidrug Resistance Proteins by Mutations at the Dimer Interface. Journal of Bacteriology, 2011, 193, 5929-5935.	2.2	10

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127	Relative role(s) of leucine versus isoleucine in the folding of membrane proteins. Peptide Science, 2019, 111, e24075.	1.8	10
128	Differential Binding of L- vs. D-isomers of Cationic Antimicrobial Peptides to the Biofilm Exopolysaccharide Alginate. Protein and Peptide Letters, 2013, 20, 843-847.	0.9	10
129	The hydrophobicity threshold for peptide insertion into membranes. Current Topics in Membranes, 2002, 52, 465-479.	0.9	9
130	Positional dependence of non-native polar mutations on folding of CFTR helical hairpins. Biochimica Et Biophysica Acta - Biomembranes, 2008, 1778, 79-87.	2.6	9
131	Sequence Hydropathy Dominates Membrane Protein Response to Detergent Solubilization. Biochemistry, 2012, 51, 6228-6237.	2.5	9
132	Effects of a polar amino acid substitution on helix formation and aggregate size along the detergent-induced peptide folding pathway. Biochimica Et Biophysica Acta - Biomembranes, 2013, 1828, 373-381.	2.6	9
133	Peptide-Based Approach to Inhibition of the Multidrug Resistance Efflux Pump AcrB. Biochemistry, 2020, 59, 3973-3981.	2.5	9
134	Conjugation of Polyethylene Glycol via a Disulfide Bond Confers Water Solubility upon a Peptide Model of a Protein Transmembrane Segment. Analytical Biochemistry, 1999, 275, 224-230.	2.4	8
135	Helicity of hydrophobic peptides in polar vs. non-polar environments. Physical Chemistry Chemical Physics, 1999, 1, 1539.	2.8	8
136	Transmembrane segment peptides with double D-amino acid replacements: Helicity, hydrophobicity, and antimicrobial activity. Biopolymers, 2003, 71, 77-84.	2.4	8
137	Functional response of the small multidrug resistance protein EmrE to mutations in transmembrane helix 2. FEBS Letters, 2014, 588, 3720-3725.	2.8	7
138	Helix-Helix Interactions: Is the Medium the Message?. Structure, 2015, 23, 437-438.	3.3	7
139	Evidence that the translocon may function as a hydropathy partitioning filter. Biochimica Et Biophysica Acta - Biomembranes, 2010, 1798, 1995-1998.	2.6	5
140	Converting a Marginally Hydrophobic Soluble Protein into a Membrane Protein. Journal of Molecular Biology, 2011, 407, 171-179.	4.2	5
141	Structural basis for misfolding at a disease phenotypic position in CFTR: Comparison of TM3/4 helix-loop-helix constructs with TM4 peptides. Biochimica Et Biophysica Acta - Biomembranes, 2012, 1818, 49-54.	2.6	5
142	Hydrophobic Blocks Facilitate Lipid Compatibility and Translocon Recognition of Transmembrane Protein Sequences. Biochemistry, 2015, 54, 1465-1473.	2.5	5
143	Structural effects of extracellular loop mutations in CFTR helical hairpins. Biochimica Et Biophysica Acta - Biomembranes, 2018, 1860, 1092-1098.	2.6	5
144	[4] Use of ionophores for manipulating intracellular ion concentrations. Methods in Neurosciences, 1995, 27, 52-68.	0.5	4

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145	Terminal Residue Hydrophobicity Modulates Transmembrane Helix–Helix Interactions. Biochemistry, 2014, 53, 3747-3757.	2.5	4
146	Structural impact of proline mutations in the loop region of an ancestral membrane protein. Biopolymers, 2016, 106, 37-42.	2.4	4
147	Hydrophobic Clusters Raise the Threshold Hydrophilicity for Insertion of Transmembrane Sequences in Vivo. Biochemistry, 2016, 55, 5772-5779.	2.5	4
148	Modulating Transmembrane α-Helix Interactions through pH-Sensitive Boundary Residues. Biochemistry, 2016, 55, 4306-4315.	2.5	4
149	Heat treatment of thioredoxin fusions increases the purity of αâ€helical transmembrane protein constructs. Protein Science, 2021, 30, 1974-1982.	7.6	4
150	The modulation of bovine milk d-galactosyltransferase by various phosphatidylethanolamines. Carbohydrate Research, 1986, 149, 47-58.	2.3	3
151	Design of Transmembrane Peptides: Coping with Sticky Situations. Methods in Molecular Biology, 2013, 1063, 197-210.	0.9	3
152	Design and Characterization of a Membrane Protein Unfolding Platform in Lipid Bilayers. PLoS ONE, 2015, 10, e0120253.	2.5	3
153	Enhanced proteolytic resistance of cationic antimicrobial peptides through lysine side chain analogs and cyclization. Biochemical and Biophysical Research Communications, 2022, 612, 105-109.	2.1	3
154	Design, expression, and purification of de novo transmembrane "hairpin―peptides. Biopolymers, 2012, 98, 546-556.	2.4	2
155	Deuterated Digoxin. Analytical Letters, 1989, 22, 2783-2790.	1.8	1
156	3 Membrane protein folding in detergents. , 2011, , 23-46.		1
157	Protection or Destruction: The LL-37/HNP1 Cooperativity Switch. Biophysical Journal, 2020, 119, 2370-2371.	0.5	1
158	Defining the Defect in F508 del CFTR: A Soluble Problem?. Chemistry and Biology, 2008, 15, 3-4.	6.0	0
159	Mechanistic Insight into Peptide-Based Efflux Pump Inhibitors Against Pathogenic Bacteria. Biophysical Journal, 2021, 120, 74a.	0.5	0
160	Peptide-Based Inhibition of AcrB Efflux Activity. Biophysical Journal, 2021, 120, 70a.	0.5	0