

Teruna J Siahaan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1076819/publications.pdf>

Version: 2024-02-01

154
papers

4,680
citations

109321

35
h-index

123424

61
g-index

160
all docs

160
docs citations

160
times ranked

5395
citing authors

#	ARTICLE	IF	CITATIONS
1	Inhibition of LFA-1/ICAM-1 and VLA-4/VCAM-1 as a therapeutic approach to inflammation and autoimmune diseases. <i>Medicinal Research Reviews</i> , 2002, 22, 146-167.	10.5	339
2	The Role of Thiols and Disulfides on Protein Stability. <i>Current Protein and Peptide Science</i> , 2009, 10, 614-625.	1.4	317
3	PLGA Nanoparticle-peptide Conjugate Effectively Targets Intercellular Cell-Adhesion Molecule-1. <i>Bioconjugate Chemistry</i> , 2008, 19, 145-152.	3.6	176
4	ICAM-1 targeting of doxorubicin-loaded PLGA nanoparticles to lung epithelial cells. <i>European Journal of Pharmaceutical Sciences</i> , 2009, 37, 141-150.	4.0	161
5	Solution stability of linear vs. cyclic RGD peptides. <i>Chemical Biology and Drug Design</i> , 1999, 53, 530-541.	1.1	160
6	Doxorubicin-loaded iron oxide nanoparticles for glioblastoma therapy: a combinational approach for enhanced delivery of nanoparticles. <i>Scientific Reports</i> , 2020, 10, 11292.	3.3	160
7	Targeting ICAM-1/LFA-1 interaction for controlling autoimmune diseases: designing peptide and small molecule inhibitors. <i>Peptides</i> , 2003, 24, 487-501.	2.4	126
8	Peptide-mediated targeted drug delivery. <i>Medicinal Research Reviews</i> , 2012, 32, 637-658.	10.5	122
9	Cell Adhesion Molecules for Targeted Drug Delivery. <i>Journal of Pharmaceutical Sciences</i> , 2006, 95, 1856-1872.	3.3	108
10	Pathways and progress in improving drug delivery through the intestinal mucosa and blood-brain barriers. <i>Therapeutic Delivery</i> , 2014, 5, 1143-1163.	2.2	99
11	Effect of restricted conformational flexibility on the permeation of model hexapeptides across Caco-2 cell monolayers. <i>Pharmaceutical Research</i> , 1997, 14, 169-175.	3.5	83
12	Controlling Ligand Surface Density Optimizes Nanoparticle Binding to ICAM-1. <i>Journal of Pharmaceutical Sciences</i> , 2011, 100, 1045-1056.	3.3	78
13	Protein PEGylation for cancer therapy: bench to bedside. <i>Journal of Cell Communication and Signaling</i> , 2019, 13, 319-330.	3.4	76
14	The effect of beta-turn structure on the passive diffusion of peptides across Caco-2 cell monolayers. <i>Pharmaceutical Research</i> , 1997, 14, 1332-1340.	3.5	75
15	VEGF-A stimulation of leukocyte adhesion to colonic microvascular endothelium: implications for inflammatory bowel disease. <i>American Journal of Physiology - Renal Physiology</i> , 2006, 290, G648-G654.	3.4	72
16	Vaccine-like Controlled-Release Delivery of an Immunomodulating Peptide To Treat Experimental Autoimmune Encephalomyelitis. <i>Molecular Pharmaceutics</i> , 2012, 9, 979-985.	4.6	65
17	Esterase-sensitive cyclic prodrugs of peptides: evaluation of an acyloxyalkoxy promoiety in a model hexapeptide. <i>Pharmaceutical Research</i> , 1996, 13, 1615-1623.	3.5	62
18	Molecular Structure of the Apical Junction Complex and Its Contribution to the Paracellular Barrier. <i>Journal of Pharmaceutical Sciences</i> , 1997, 86, 977-984.	3.3	55

#	ARTICLE	IF	CITATIONS
19	Improving the selectivity of HAV-peptides in modulating E-cadherin-E-cadherin interactions in the intercellular junction of MDCK cell monolayers. <i>Pharmaceutical Research</i> , 2001, 18, 446-453.	3.5	55
20	Linear and cyclic LFA-1 and ICAM-1 peptides inhibit T cell adhesion and function. <i>Peptides</i> , 2000, 21, 1161-1167.	2.4	52
21	Synthesis of a Novel Esterase-Sensitive Cyclic Prodrug System for Peptides That Utilizes a α -Trimethyl Lock-Facilitated Lactonization Reaction. <i>Journal of Organic Chemistry</i> , 1997, 62, 1363-1367.	3.2	51
22	Increasing paracellular porosity by E-cadherin peptides: discovery of bulge and groove regions in the EC1-domain of E-cadherin. <i>Pharmaceutical Research</i> , 2002, 19, 1170-1179.	3.5	51
23	Modulation of cellular adhesion in bovine brain microvessel endothelial cells by a decapeptide. <i>Brain Research</i> , 1997, 747, 103-113.	2.2	50
24	Synthesis of a Novel Esterase-Sensitive Cyclic Prodrug of a Hexapeptide Using an (Acyloxy)alkoxy Promoiety. <i>Journal of Organic Chemistry</i> , 1997, 62, 1356-1362.	3.2	48
25	The effect of conformation on membrane permeability of an acyloxyalkoxy-linked cyclic prodrug of a model hexapeptide. <i>Pharmaceutical Research</i> , 1996, 13, 1657-1662.	3.5	45
26	Adhesion of pancreatic beta cells to biopolymer films. <i>Biopolymers</i> , 2009, 91, 676-685.	2.4	44
27	Enhancement of Drug Absorption through the Blood-Brain Barrier and Inhibition of Intercellular Tight Junction Resealing by E-Cadherin Peptides. <i>Molecular Pharmaceutics</i> , 2011, 8, 239-249.	4.6	44
28	Acyloxyalkoxy-based cyclic prodrugs of opioid peptides: evaluation of the chemical and enzymatic stability as well as their transport properties across Caco-2 cell monolayers. <i>Pharmaceutical Research</i> , 1999, 16, 24-29.	3.5	43
29	Modulation of Blood-Brain Barrier Permeability in Mice Using Synthetic E-Cadherin Peptide. <i>Molecular Pharmaceutics</i> , 2014, 11, 974-981.	4.6	42
30	PEPTIDES DERIVED FROM ICAM-1 AND LFA-1 MODULATE T CELL ADHESION AND IMMUNE FUNCTION IN A MIXED LYMPHOCYTE CULTURE1. <i>Transplantation</i> , 1999, 68, 685-692.	1.0	41
31	Structure and Function of the Intercellular Junctions: Barrier of Paracellular Drug Delivery. <i>Current Pharmaceutical Design</i> , 2006, 12, 2813-2824.	1.9	40
32	Antigen-Specific Suppression of Experimental Autoimmune Encephalomyelitis by a Novel Bifunctional Peptide Inhibitor. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2007, 322, 879-886.	2.5	40
33	Modulation of Intercellular Junctions by Cyclic-ADT Peptides as a Method to Reversibly Increase Blood-Brain Barrier Permeability. <i>Journal of Pharmaceutical Sciences</i> , 2015, 104, 1065-1075.	3.3	39
34	Calcium Condensed LABL-TAT Complexes Effectively Target Gene Delivery to ICAM-1 Expressing Cells. <i>Molecular Pharmaceutics</i> , 2011, 8, 788-798.	4.6	38
35	Separation and Analysis of Peptides and Proteins. <i>Analytical Chemistry</i> , 1997, 69, 29-58.	6.5	36
36	Binding and internalization of an LFA-1-derived cyclic peptide by ICAM receptors on activated lymphocyte: a potential ligand for drug targeting to ICAM-1-expressing cells. , 2001, 18, 329-335.		36

#	ARTICLE	IF	CITATIONS
37	Mechanism of binding and internalization of ICAM-1-derived cyclic peptides by LFA-1 on the surface of T cells: a potential method for targeted drug delivery. <i>Pharmaceutical Research</i> , 2003, 20, 1523-1532.	3.5	35
38	Codelivery of antigen and an immune cell adhesion inhibitor is necessary for efficacy of soluble antigen arrays in experimental autoimmune encephalomyelitis. <i>Molecular Therapy - Methods and Clinical Development</i> , 2014, 1, 14008.	4.1	35
39	Modulation of Melphalan Resistance in Glioma Cells with a Peripheral Benzodiazepine Receptor Ligand [†] Melphalan Conjugate. <i>Journal of Medicinal Chemistry</i> , 1997, 40, 1726-1730.	6.4	32
40	Immune modulating peptides for the treatment and suppression of multiple sclerosis. <i>Clinical Immunology</i> , 2012, 144, 127-138.	3.2	30
41	Single-step grafting of aminoxy-peptides to hyaluronan: A simple approach to multifunctional therapeutics for experimental autoimmune encephalomyelitis. <i>Journal of Controlled Release</i> , 2013, 168, 334-340.	9.9	30
42	Comparison of Linear and Cyclic His-Ala-Val Peptides in Modulating the Blood-Brain Barrier Permeability: Impact on Delivery of Molecules to the Brain. <i>Journal of Pharmaceutical Sciences</i> , 2016, 105, 797-807.	3.3	30
43	Suppression of Type 1 Diabetes in NOD Mice by Bifunctional Peptide Inhibitor: Modulation of the Immunological Synapse Formation. <i>Chemical Biology and Drug Design</i> , 2007, 70, 227-236.	3.2	29
44	Inhibition of the adherence of T-lymphocytes to epithelial cells by a cyclic peptide derived from inserted domain of lymphocyte function-associated antigen-1. <i>Inflammation</i> , 2001, 25, 203-214.	3.8	28
45	Suppression of EAE and prevention of blood-brain barrier breakdown after vaccination with novel bifunctional peptide inhibitor. <i>Neuropharmacology</i> , 2012, 62, 1874-1881.	4.1	28
46	Antigen-specific blocking of CD4-specific immunological synapse formation using BPI and current therapies for autoimmune diseases. <i>Medicinal Research Reviews</i> , 2012, 32, 727-764.	10.5	28
47	Endotoxaemia-augmented murine venous thrombosis is dependent on TLR-4 and ICAM-1, and potentiated by neutropenia. <i>Thrombosis and Haemostasis</i> , 2017, 117, 339-348.	3.4	28
48	Prophylactic and therapeutic suppression of experimental autoimmune encephalomyelitis by a novel bifunctional peptide inhibitor. <i>Clinical Immunology</i> , 2008, 129, 69-79.	3.2	26
49	Structure, Size, and Solubility of Antigen Arrays Determines Efficacy in Experimental Autoimmune Encephalomyelitis. <i>AAPS Journal</i> , 2014, 16, 1185-1193.	4.4	26
50	Co-Delivery of Autoantigen and B7 Pathway Modulators Suppresses Experimental Autoimmune Encephalomyelitis. <i>AAPS Journal</i> , 2014, 16, 1204-1213.	4.4	26
51	Improving Brain Delivery of Biomolecules via BBB Modulation in Mouse and Rat: Detection using MRI, NIRF, and Mass Spectrometry. <i>Nanotheranostics</i> , 2017, 1, 217-231.	5.2	26
52	A modified coumarinic acid-based cyclic prodrug of an opioid peptide: its enzymatic and chemical stability and cell permeation characteristics. <i>Pharmaceutical Research</i> , 2002, 19, 794-801.	3.5	25
53	Immune response to controlled release of immunomodulating peptides in a murine experimental autoimmune encephalomyelitis (EAE) model. <i>Journal of Controlled Release</i> , 2010, 141, 145-152.	9.9	25
54	Solution Structure of a Cyclic RGD Peptide That Inhibits Platelet Aggregation. <i>Journal of Biomolecular Structure and Dynamics</i> , 1996, 14, 1-11.	3.5	24

#	ARTICLE	IF	CITATIONS
55	Modulation of the Cellular Junction Protein E-Cadherin in Bovine Brain Microvessel Endothelial Cells by Cadherin Peptides. <i>Drug Delivery</i> , 1997, 4, 187-193.	5.7	24
56	Inhibition of E-Cadherin-Mediated Homotypic Adhesion of Caco-2 Cells: A Novel Evaluation Assay for Peptide Activities in Modulating Cell-Cell Adhesion. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006, 317, 309-316.	2.5	24
57	Synergistic inhibitory activity of $\hat{1}\pm$ - and $\hat{1}^2$ -LFA-1 peptides on LFA-1/ICAM-1 interaction. <i>Peptides</i> , 2001, 22, 1955-1962.	2.4	23
58	ICAM-1 Peptide Inhibitors of T-cell Adhesion bind to the allosteric site of LFA-1. An NMR Characterization. <i>Chemical Biology and Drug Design</i> , 2007, 70, 347-353.	3.2	23
59	Antigen-Specific Suppression of Experimental Autoimmune Encephalomyelitis by a Novel Bifunctional Peptide Inhibitor: Structure Optimization and Pharmacokinetics. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010, 332, 1136-1145.	2.5	23
60	Inhibition of ICAM-1/LFA-1-mediated heterotypic T-cell adhesion to epithelial cells: design of ICAM-1 cyclic peptides. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 1399-1402.	2.2	22
61	Nanoparticles Targeting Dendritic Cell Surface Molecules Effectively Block T Cell Conjugation and Shift Response. <i>ACS Nano</i> , 2011, 5, 1693-1702.	14.6	22
62	Influence of particle size, an elongated particle geometry, and adjuvants on dendritic cell activation. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2015, 94, 542-549.	4.3	21
63	Rapid Determination of Substrate Specificity of Clostridium histolyticum $\hat{1}^2$ -Collagenase Using an Immobilized Peptide Library. <i>Journal of Biological Chemistry</i> , 2002, 277, 8366-8371.	3.4	20
64	$\hat{1}\pm$ -L-Integrin I domain cyclic peptide antagonist selectively inhibits T cell adhesion to pancreatic islet microvascular endothelium. <i>American Journal of Physiology - Renal Physiology</i> , 2005, 288, G67-G73.	3.4	20
65	Characterization of Binding Properties of ICAM-1 Peptides to LFA-1: Inhibitors of T-cell Adhesion. <i>Chemical Biology and Drug Design</i> , 2006, 68, 20-28.	3.2	20
66	Mechanism of Internalization of an ICAM-1-Derived Peptide by Human Leukemic Cell Line HL-60: Influence of Physicochemical Properties on Targeted Drug Delivery. <i>Molecular Pharmaceutics</i> , 2007, 4, 749-758.	4.6	20
67	Evaluation of the physical stability of the EC5 domain of E-cadherin: Effects of pH, temperature, ionic strength, and disulfide bonds. <i>Journal of Pharmaceutical Sciences</i> , 2009, 98, 63-73.	3.3	20
68	Autoimmune therapies targeting costimulation and emerging trends in multivalent therapeutics. <i>Therapeutic Delivery</i> , 2011, 2, 873-889.	2.2	20
69	Structural and ICAM-1-Docking Properties of a Cyclic Peptide from the I-domain of LFA-1: An inhibitor of ICAM-1/LFA-1-mediated T-cell adhesion. <i>Journal of Biomolecular Structure and Dynamics</i> , 2002, 19, 789-799.	3.5	19
70	Inhibition of homotypic adhesion of T-cells: secondary structure of an ICAM-1-derived cyclic peptide. <i>Chemical Biology and Drug Design</i> , 1997, 49, 517-526.	1.1	19
71	Disulfide Bond Formation Promotes the cis- and trans-Dimerization of the E-cadherin-derived First Repeat. <i>Journal of Biological Chemistry</i> , 2002, 277, 16002-16010.	3.4	18
72	Synthesis and chemical stability of a disulfide bond in a model cyclic pentapeptide: Cyclo(1,4)-Cys-Gly-Phe-Cys-Gly-OH. <i>Journal of Pharmaceutical Sciences</i> , 2006, 95, 2222-2234.	3.3	18

#	ARTICLE	IF	CITATIONS
73	The importance of structural factors on the rate and the extent of N,O-acyl migration in cyclic and linear peptides. <i>Pharmaceutical Research</i> , 1995, 12, 323-328.	3.5	17
74	Suppression of MOG- and PLP-induced experimental autoimmune encephalomyelitis using a novel multivalent bifunctional peptide inhibitor. <i>Journal of Neuroimmunology</i> , 2013, 263, 20-27.	2.3	17
75	Brain Delivery of Drug and MRI Contrast Agent: Detection and Quantitative Determination of Brain Deposition of CPT-Glu Using LC-MS/MS and Gd-DTPA Using Magnetic Resonance Imaging. <i>Molecular Pharmaceutics</i> , 2016, 13, 379-390.	4.6	17
76	Probing the interaction between cHAVc3 peptide and the EC1 domain of E-cadherin using NMR and molecular dynamics simulations. <i>Journal of Biomolecular Structure and Dynamics</i> , 2017, 35, 92-104.	3.5	17
77	Derivatives of Melphalan Designed to Enhance Drug Accumulation in Cancer Cells. <i>Journal of Drug Targeting</i> , 1997, 4, 359-370.	4.4	16
78	Synthesis of a novel cyclic prodrug of RGD peptidomimetic to improve its cell membrane permeation. <i>Bioorganic Chemistry</i> , 2002, 30, 285-301.	4.1	16
79	Effects of Amino Acid Chirality and the Chemical Linker on the Cell Permeation Characteristics of Cyclic Prodrugs of Opioid Peptides. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 1261-1270.	6.4	16
80	Hyaluronic Acid Graft Polymers Displaying Peptide Antigen Modulate Dendritic Cell Response in Vitro. <i>Molecular Pharmaceutics</i> , 2014, 11, 367-373.	4.6	16
81	Rapid Identification of Fluorochrome Modification Sites in Proteins by LC ESI-Q-TOF Mass Spectrometry. <i>Bioconjugate Chemistry</i> , 2011, 22, 1330-1336.	3.6	15
82	Methotrexate disposition, anti-folate activity and efficacy in the collagen-induced arthritis mouse model. <i>European Journal of Pharmacology</i> , 2019, 853, 264-274.	3.5	15
83	Design, structure and biological activity of β -turn peptides of CD2 protein for inhibition of T-cell adhesion. <i>FEBS Journal</i> , 2004, 271, 2873-2886.	0.2	14
84	Sequence Recognition of α -LFA-1-derived Peptides by ICAM-1 Cell Receptors: Inhibitors of T-cell Adhesion. <i>Chemical Biology and Drug Design</i> , 2007, 70, 237-246.	3.2	14
85	cIBR Effectively Targets Nanoparticles to LFA-1 on Acute Lymphoblastic T Cells. <i>Molecular Pharmaceutics</i> , 2010, 7, 146-155.	4.6	14
86	Improving In Vivo Brain Delivery of Monoclonal Antibody Using Novel Cyclic Peptides. <i>Pharmaceutics</i> , 2019, 11, 568.	4.5	14
87	Orf239342 from the mushroom <i>Agaricus bisporus</i> is a mannose binding protein. <i>Biochemical and Biophysical Research Communications</i> , 2019, 515, 99-103.	2.1	14
88	Noninvasive Brain Delivery and Efficacy of BDNF to Stimulate Neuroregeneration and Suppression of Disease Relapse in EAE Mice. <i>Molecular Pharmaceutics</i> , 2020, 17, 404-416.	4.6	14
89	Non-invasive Brain Delivery and Efficacy of BDNF in APP/PS1 Transgenic Mice. <i>Medical Research Archives</i> , 2020, 8, .	0.2	14
90	Steric hindrance is a key factor in the coupling reaction of (acyloxy) alkyl-halides with phenols to make a new promoiety for prodrugs. <i>Tetrahedron Letters</i> , 2002, 43, 577-579.	1.4	13

#	ARTICLE	IF	CITATIONS
91	Utilization of I-domain of LFA-1 to Target Drug and Marker Molecules to Leukocytes. <i>Theranostics</i> , 2011, 1, 277-289.	10.0	13
92	Methotrexate (MTX)â€œcIBR Conjugate for Targeting MTX to Leukocytes: Conjugate Stability and In Vivo Efficacy in Suppressing Rheumatoid Arthritis. <i>Journal of Pharmaceutical Sciences</i> , 2012, 101, 3275-3291.	3.3	13
93	Validation of Cadherin HAV6 Peptide in the Transient Modulation of the Blood-Brain Barrier for the Treatment of Brain Tumors. <i>Pharmaceutics</i> , 2019, 11, 481.	4.5	13
94	Conjugates of Cell Adhesion Peptides for Therapeutics and Diagnostics Against Cancer and Autoimmune Diseases. <i>Current Topics in Medicinal Chemistry</i> , 2018, 17, 3425-3443.	2.1	13
95	Synthesis of Cyclic Prodrugs of Aggrastat and Its Analogue with a Modified Phenylpropionic Acid Linker. <i>Organic Letters</i> , 2002, 4, 549-552.	4.6	12
96	Peptides and Drug Delivery. <i>Advances in Experimental Medicine and Biology</i> , 2017, 1030, 167-184.	1.6	12
97	Secondary Structure of the HAV Peptide Which Regulates Cadherin-Cadherin Interaction. <i>Journal of Biomolecular Structure and Dynamics</i> , 1995, 13, 447-455.	3.5	11
98	Conjugation with L-Glutamate for in vivo Brain Drug Delivery. <i>Journal of Drug Targeting</i> , 2001, 9, 23-37.	4.4	11
99	Effect of Modification of the Physicochemical Properties of ICAM-1-Derived Peptides on Internalization and Intracellular Distribution in the Human Leukemic Cell Line HL-60. <i>Molecular Pharmaceutics</i> , 2009, 6, 396-406.	4.6	11
100	The Role of Covalent Dimerization on the Physical and Chemical Stability of the EC1 Domain of Human E-Cadherin. <i>Journal of Pharmaceutical Sciences</i> , 2009, 98, 3562-3574.	3.3	11
101	Characterization of Multiple Stable Conformers of the EC5 Domain of E-cadherin and the Interaction of EC5 with E-cadherin Peptides. <i>Chemical Biology and Drug Design</i> , 2009, 73, 584-598.	3.2	11
102	Deamidation of model β -turn cyclic peptides in the solid state. <i>Journal of Pharmaceutical Sciences</i> , 2005, 94, 2616-2631.	3.3	10
103	Pulmonary Drug Delivery: <i>Pharmaceutical Chemistry and Aerosol Technology</i> . , 2005, , 341-361.		10
104	Structural Modifications of ICAM-1 Cyclic Peptides to Improve the Activity to Inhibit Heterotypic Adhesion of T cells. <i>Chemical Biology and Drug Design</i> , 2008, 72, 27-33.	3.2	10
105	Immune Tolerance Induction against Experimental Autoimmune Encephalomyelitis (EAE) Using A New PLP-B7AP Conjugate that Simultaneously Targets B7/CD28 Costimulatory Signal and TCR/MHC-II Signal. <i>Journal of Multiple Sclerosis</i> , 2014, 02, .	0.1	10
106	Comparison of the Solution Conformations of a Cell-Adhesive Peptide LBE and its Reverse Sequence EBL. <i>Journal of Biomolecular Structure and Dynamics</i> , 1999, 17, 429-444.	3.5	9
107	Oral Protein and Peptide Drug Delivery. , 2005, , 189-200.		9
108	A Peptide from the Beta-strand Region of CD2 Protein that Inhibits Cell Adhesion and Suppresses Arthritis in a Mouse Model. <i>Chemical Biology and Drug Design</i> , 2010, 76, 234-244.	3.2	9

#	ARTICLE	IF	CITATIONS
109	Controlling immune response and demyelination using highly potent bifunctional peptide inhibitors in the suppression of experimental autoimmune encephalomyelitis. <i>Clinical and Experimental Immunology</i> , 2013, 172, 23-36.	2.6	9
110	In Vivo Brain Delivery and Brain Deposition of Proteins with Various Sizes. <i>Molecular Pharmaceutics</i> , 2019, 16, 4878-4889.	4.6	9
111	Synthesis and stability study of a modified phenylpropionic acid linker-based esterase-sensitive prodrug. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2002, 12, 3439-3442.	2.2	8
112	Modulation of Cell Adhesion Molecules in Various Epithelial Cell Lines after Treatment with PP2A. <i>Molecular Pharmaceutics</i> , 2005, 2, 170-184.	4.6	8
113	Gram-Negative Pneumonia Alters Large-Vein Cell-Adhesion Molecule Profile and Potentiates Experimental Stasis Venous Thrombosis. <i>Journal of Vascular Research</i> , 2016, 53, 186-195.	1.4	8
114	Synthesis of a Bifunctional Peptide Inhibitor-IgG1 Fc Fusion That Suppresses Experimental Autoimmune Encephalomyelitis. <i>Bioconjugate Chemistry</i> , 2017, 28, 1867-1877.	3.6	8
115	A Peptide Derived from LFA-1 Protein that Modulates T-cell Adhesion Binds to Soluble ICAM-1 Protein. <i>Journal of Biomolecular Structure and Dynamics</i> , 2003, 20, 635-644.	3.5	7
116	Factors That Impact the Developability of Drug Candidates: An Overview. , 2005, , 1-14.		7
117	Liposomes as Drug Delivery Vehicles. , 2005, , 411-434.		7
118	The aqueous conformation of cyclo(1,6)Ac-Cys-Arg-Gly-Asp-Phe-Pen-NH ₂ . <i>International Journal of Peptide and Protein Research</i> , 1994, 44, 427-434.	0.1	7
119	Localized production of human E-cadherin-derived first repeat in <i>Escherichia coli</i> . <i>Protein Expression and Purification</i> , 2002, 26, 449-454.	1.3	6
120	N-cadherin involvement in the heterotypic adherence of malignant T-cells to epithelia. <i>Molecular and Cellular Biochemistry</i> , 2002, 233, 1-8.	3.1	6
121	Reductive Alkylation of Lipase: Experimental and Molecular Modeling Approaches. <i>Applied Biochemistry and Biotechnology</i> , 2004, 118, 011-020.	2.9	6
122	Effects of An E-cadherin-Derived Peptide on the Gene Expression of Caco-2 Cells. <i>Pharmaceutical Research</i> , 2004, 21, 2085-2094.	3.5	6
123	Prodrug Approaches to Drug Delivery. , 2005, , 125-165.		6
124	I-Domain-Antigen Conjugate (IDAC) for Delivering Antigenic Peptides to APC: Synthesis, Characterization, and in Vivo EAE Suppression. <i>Bioconjugate Chemistry</i> , 2012, 23, 509-517.	3.6	6
125	Improving the stability of the EC1 domain of E-cadherin by thiol alkylation of the cysteine residue. <i>International Journal of Pharmaceutics</i> , 2012, 431, 16-25.	5.2	6
126	Peptide Delivery. , 2013, , 1702-1710.		6

#	ARTICLE	IF	CITATIONS
127	Bifunctional Peptide Inhibitors Suppress Interleukin-6 Proliferation and Ameliorates Murine Collagen-Induced Arthritis. <i>Journal of Clinical & Cellular Immunology</i> , 2014, 05, .	1.5	6
128	Synthesis and comparison of physicochemical, transport, and antithrombic properties of a cyclic prodrug and the parent RGD peptidomimetic. <i>Tetrahedron</i> , 2003, 59, 2861-2869.	1.9	5
129	Ultrasound-Mediated Drug Delivery. , 2005, , 245-278.		4
130	Physiological, Biochemical, and Chemical Barriers to Oral Drug Delivery. , 2005, , 15-27.		4
131	Vaccinelike and Prophylactic Treatments of EAE with Novel I-Domain Antigen Conjugates (IDAC): Targeting Multiple Antigenic Peptides to APC. <i>Molecular Pharmaceutics</i> , 2013, 10, 297-306.	4.6	4
132	¹ H, ¹³ C and ¹⁵ N backbone assignment of the EC-1 domain of human E-cadherin. <i>Biomolecular NMR Assignments</i> , 2015, 9, 31-35.	0.8	4
133	A Tribute to Ronald T. Borchardt "Teacher, Mentor, Scientist, Colleague, Leader, Friend, and Family Man. <i>Journal of Pharmaceutical Sciences</i> , 2016, 105, 370-385.	3.3	4
134	Expression, purification, and structural study of the EC4 domain of E-cadherin. <i>Protein Expression and Purification</i> , 2004, 33, 72-79.	1.3	3
135	Efflux Transporters in Drug Excretion. , 2005, , 381-410.		3
136	Pathways for Drug Delivery to the Central Nervous System. , 2005, , 29-56.		3
137	Receptor-Mediated Drug Delivery. , 2005, , 167-187.		3
138	Solution structure of a novel T _H 1 cell adhesion inhibitor derived from the fragment of ICAM-1 receptor: Cyclo(1,8)-Cys-Pro-Arg-Gly-Gly-Ser-Val-Cys. <i>Biopolymers</i> , 2009, 91, 633-641.	2.4	3
139	Conformational analysis of cyclo(2,9)-QCRSVEGSCG-OH from the C-terminal loop of human growth hormone. <i>Chemical Biology and Drug Design</i> , 1997, 49, 15-22.	1.1	3
140	Gene Therapy and Gene Delivery. , 2005, , 305-319.		2
141	Methods of Delivering Molecules Through the Blood-Brain Barrier for Brain Diagnostics and Therapeutics. <i>Neuromethods</i> , 2019, , 9-43.	0.3	2
142	Enhancing Intestinal Absorption of a Model Macromolecule via the Paracellular Pathway using E-Cadherin Peptides. <i>Journal of Pharmaceutical Sciences</i> , 2021, 110, 2139-2148.	3.3	2
143	Antibody-Directed Drug Delivery. , 2005, , 363-379.		1
144	Regulatory and Intellectual Property Issues in Drug Delivery Research. , 2005, , 435-442.		1

#	ARTICLE	IF	CITATIONS
145	Physicochemical Properties, Formulation, and Drug Delivery. , 2005, , 57-71.		1
146	Parenteral Formulation for Peptides, Proteins, and Monoclonal Antibodies Drugs: A Commercial Development Overview. , 2005, , 321-339.		0
147	Metabolic Activation and Drug Targeting. , 2005, , 201-244.		0
148	Polycationic Peptides and Proteins in Drug Delivery: Focus on Nonclassical Transport. , 2005, , 279-304.		0
149	Targeted Bioavailability: A Fresh Look at Pharmacokinetic and Pharmacodynamic Issues in Drug Delivery. , 2005, , 73-82.		0
150	Presystemic and First-Pass Metabolism. , 2005, , 83-101.		0
151	Cell Culture Models for Drug Transport Studies. , 2005, , 103-124.		0
152	ICAM-1 Peptide Inhibitors of T-cell Adhesion bind to the allosteric site of LFA-1. An NMR Characterization. Chemical Biology and Drug Design, 2007, .	3.2	0
153	Cadherin peptide-induced enhancement of blood brain barrier (BBB) permeability. FASEB Journal, 2013, 27, 668.3.	0.5	0
154	Abstract 191: Endothelial Dysfunction Potentiates Deep Venous Thrombosis in a Mouse Model of Sepsis. Arteriosclerosis, Thrombosis, and Vascular Biology, 2014, 34, .	2.4	0