

Shai Rahimipour

List of Publications by Year in descending order

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citations

279701

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315616

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docs citations

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times ranked

2307
citing authors

#	ARTICLE	IF	CITATIONS
1	Cu ²⁺ -Induced self-assembly and amyloid formation of a cyclic α -peptide: structure and function. <i>Physical Chemistry Chemical Physics</i> , 2022, 24, 6699-6715.	1.3	3
2	Glucose-Functionalized Liposomes for Reducing False Positives in Cancer Diagnosis. <i>ACS Nano</i> , 2021, 15, 1301-1309.	7.3	14
3	Polydopamine Nanoparticles Containing a Cisplatin Analog for Anticancer Treatment and Diagnostics. <i>ACS Applied Nano Materials</i> , 2021, 4, 14126-14135.	2.4	2
4	Gold Nanorod-Based Bio-Barcode Sensor Array for Enzymatic Detection in Biomedical Applications. <i>ACS Applied Nano Materials</i> , 2020, 3, 8414-8423.	2.4	7
5	Bifunctional Carbon Dots ⁺ Magnetic and Fluorescent Hybrid Nanoparticles for Diagnostic Applications. <i>Nanomaterials</i> , 2020, 10, 1384.	1.9	13
6	Biological Logic Gate Using Gold Nanoparticles and Fluorescence Lifetime Imaging Microscopy. <i>ACS Applied Nano Materials</i> , 2019, 2, 6527-6536.	2.4	26
7	Neuroprotective Effect of Nerve Growth Factor Loaded in Porous Silicon Nanostructures in an Alzheimer's Disease Model and Potential Delivery to the Brain. <i>Small</i> , 2019, 15, e1904203.	5.2	30
8	Porous Materials: Neuroprotective Effect of Nerve Growth Factor Loaded in Porous Silicon Nanostructures in an Alzheimer's Disease Model and Potential Delivery to the Brain (<i>Small</i> 45/2019). <i>Small</i> , 2019, 15, 1970245.	5.2	0
9	Photoactive chlorin e6 is a multifunctional modulator of amyloid- β aggregation and toxicity via specific interactions with its histidine residues. <i>Chemical Science</i> , 2019, 10, 208-217.	3.7	25
10	Computer-Aided Design and Synthesis of 1-[(3,4-Dihydroxybenzylidene)amino]phenyl]-5-oxopyrrolidine-3-carboxylic Acid as an Nrf2 Enhancer. <i>ChemPlusChem</i> , 2018, 83, 320-333.	5.3	9
11	Computer-Aided Design and Synthesis of 1-[(3,4-Dihydroxybenzylidene)amino]phenyl]-5-oxopyrrolidine-3-carboxylic Acid as an Nrf2 Enhancer. <i>ChemPlusChem</i> , 2018, 83, 318-318.	1.3	2
12	Inhibition of tau-derived hexapeptide aggregation and toxicity by a self-assembled cyclic α -peptide conformational inhibitor. <i>Chemical Communications</i> , 2018, 54, 5980-5983.	2.2	33
13	Distinct Effects of O-GlcNAcylation and Phosphorylation of a Tau-Derived Amyloid Peptide on Aggregation of the Native Peptide. <i>Chemistry - A European Journal</i> , 2018, 24, 14039-14043.	1.7	7
14	Specific Binding of Cu(II) Ions to Amyloid-Beta Peptides Bound to Aggregation-Inhibiting Molecules or SDS Micelles Creates Complexes that Generate Radical Oxygen Species. <i>Journal of Alzheimer's Disease</i> , 2016, 54, 971-982.	1.2	34
15	Sonochemically-Produced Metal-Containing Polydopamine Nanoparticles and Their Antibacterial and Antibiofilm Activity. <i>Langmuir</i> , 2016, 32, 5201-5212.	1.6	52
16	Self-Assembled Cyclic α -Peptides as Generic Conformational Inhibitors of the β -Synuclein Aggregation and Toxicity: In Vitro and Mechanistic Studies. <i>Chemistry - A European Journal</i> , 2016, 22, 14236-14246.	1.7	34
17	Selective Inhibition of Aggregation and Toxicity of a Tau-Derived Peptide using Its Glycosylated Analogues. <i>Chemistry - A European Journal</i> , 2016, 22, 5945-5952.	1.7	37
18	A versatile water-soluble chelating and radical scavenging platform. <i>Chemical Communications</i> , 2016, 52, 2350-2353.	2.2	13

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19	New Perspectives in Reducing Amyloid Aggregation and Toxicity. Israel Journal of Chemistry, 2015, 55, 735-748.	1.0	2
20	Resolution of Two Sub-Populations of Conformers and Their Individual Dynamics by Time Resolved Ensemble Level FRET Measurements. PLoS ONE, 2015, 10, e0143732.	1.1	14
21	Antibacterial and Antibiofilm Surfaces through Polydopamine-Assisted Immobilization of Lysostaphin as an Antibacterial Enzyme. Langmuir, 2015, 31, 1064-1073.	1.6	89
22	Tubular cell phenotype in HIV-associated nephropathy: Role of phospholipid lysophosphatidic acid. Experimental and Molecular Pathology, 2015, 99, 109-115.	0.9	5
23	Hypericin and its Derivatives Act as Radiosensitizing Agents That Can Inhibit Tumor Initiating Cell Viability. Clinical Cancer Drugs, 2015, 2, 119-127.	0.3	2
24	Structure-based study of antiamyloidogenic cyclic d,l- α -peptides. Tetrahedron, 2014, 70, 7639-7644.	1.0	14
25	Multifunctional Cyclic d,l- α -Peptide Architectures Stimulate Non-Insulin Dependent Glucose Uptake in Skeletal Muscle Cells and Protect Them Against Oxidative Stress. Journal of Medicinal Chemistry, 2013, 56, 6709-6718.	2.9	14
26	Sonochemically produced polydopamine nanocapsules with selective antimicrobial activity. Chemical Communications, 2013, 49, 5721.	2.2	69
27	In Vitro and Mechanistic Studies of an Antiamyloidogenic Self-Assembled Cyclic d,l- α -Peptide Architecture. Journal of the American Chemical Society, 2013, 135, 3474-3484.	6.6	95
28	Effective Targeting of A β to Macrophages by Sonochemically Prepared Surface-Modified Protein Microspheres. Biomacromolecules, 2013, 14, 110-116.	2.6	16
29	Surface-modified protein nanospheres as potential antiviral agents. Chemical Communications, 2012, 48, 8359.	2.2	16
30	Non-leaching antimicrobial surfaces through polydopamine bio-inspired coating of quaternary ammonium salts or an ultrashort antimicrobial lipopeptide. Journal of Materials Chemistry, 2012, 22, 2026-2032.	6.7	112
31	A facile one-pot sonochemical synthesis of surface-coated mannosyl protein microspheres for detection and killing of bacteria. Chemical Communications, 2011, 47, 12277.	2.2	18
32	Surface-Modified Protein Microspheres Capture Amyloid β and Inhibit its Aggregation and Toxicity. Chemistry - A European Journal, 2011, 17, 11171-11177.	1.7	36
33	Inside Cover: Surface-Modified Protein Microspheres Capture Amyloid β and Inhibit its Aggregation and Toxicity (Chem. Eur. J. 40/2011). Chemistry - A European Journal, 2011, 17, 11074-11074.	1.7	0
34	Towards the Efficiency of Pharmacologically Active Quinoid Compounds: Electron Transfer and Formation of Reactive Oxygen Species. Applied Magnetic Resonance, 2010, 37, 629-648.	0.6	21
35	One-Step Preparation of Multifunctional Chitosan Microspheres by a Simple Sonochemical Method. Chemistry - A European Journal, 2010, 16, 562-567.	1.7	43
36	Antibacterial cyclic d,l- α -glycopeptides. Chemical Communications, 2009, , 3693.	2.2	60

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37	Hypericin Derivatives: Substituent Effects on Radical-anion Formation. <i>Photochemistry and Photobiology</i> , 2007, 74, 149-156.	1.3	0
38	Generation of Free Radicals by Emodic Acid and its [d-Lys6]GnRH-conjugate. <i>Photochemistry and Photobiology</i> , 2007, 74, 226-236.	1.3	1
39	Novel cyclic azo-bridged analogs of gonadotropin-releasing hormone. <i>Journal of Peptide Science</i> , 2006, 12, 106-115.	0.8	6
40	?-Sulfonamido gonadotropin-releasing hormone analogs: synthesis and evaluation of several parent hormone properties. <i>Journal of Peptide Science</i> , 2005, 11, 45-52.	0.8	3
41	Novel Methyl Helianthrones as Photosensitizers: Synthesis and Biological Evaluation. <i>Photochemistry and Photobiology</i> , 2005, 81, 250.	1.3	4
42	Discovery of a Biologically Active Thiostrepton Fragment. <i>Journal of the American Chemical Society</i> , 2005, 127, 15042-15044.	6.6	66
43	Novel Methyl Helianthrones as Photosensitizers: Synthesis and Biological Evaluation. <i>Photochemistry and Photobiology</i> , 2005, 81, 250-258.	1.3	0
44	Novel Methyl Helianthrones as Photosensitizers: Synthesis and Biological Evaluation. <i>Photochemistry and Photobiology</i> , 2005, 81, 250-8.	1.3	0
45	Receptor-Mediated Targeting of a Photosensitizer by Its Conjugation to Gonadotropin-Releasing Hormone Analogues. <i>Journal of Medicinal Chemistry</i> , 2003, 46, 3965-3974.	2.9	55
46	Chemical and Photochemical Electron Transfer of New Helianthrone Derivatives: Aspects of Their Photodynamic Activity. <i>Journal of the American Chemical Society</i> , 2003, 125, 1376-1384.	6.6	38
47	The neuropeptides GnRH-II and GnRH-I are produced by human T cells and trigger laminin receptor gene expression, adhesion, chemotaxis and homing to specific organs. <i>Nature Medicine</i> , 2002, 8, 1421-1426.	15.2	109
48	The neuropeptides GnRH-II and GnRH-I are produced by human T cells and trigger laminin receptor gene expression, adhesion, chemotaxis and homing to specific organs. <i>Nature Medicine</i> , 2002, 8, 1421-1426.	15.2	24
49	Two forms of gonadotropin-releasing hormone (GnRH) are expressed in human breast tissue and overexpressed in breast cancer: a putative mechanism for the antiproliferative effect of GnRH by down-regulation of acidic ribosomal phosphoproteins P1 and P2. <i>Cancer Research</i> , 2002, 62, 1036-44.	0.4	58
50	Design, Synthesis, and Evaluation of a Long-Acting, Potent Analogue of Gonadotropin-Releasing Hormone. <i>Journal of Medicinal Chemistry</i> , 2001, 44, 3645-3652.	2.9	14
51	Hypericin Derivatives: Substituent Effects on Radical-anion Formation. <i>Photochemistry and Photobiology</i> , 2001, 74, 149.	1.3	10
52	Base catalyzed [2,3]-sigmatropic rearrangements of propargylic sulfonium and selenonium salts. <i>Tetrahedron Letters</i> , 2001, 42, 2911-2914.	0.7	11
53	Peptide Torsion Angle Measurements: Effects of Nondilute Spin Pairs on Carbon-Observed, Deuterium-Dephased PM5-REDOR. <i>Journal of Magnetic Resonance</i> , 2001, 148, 104-114.	1.2	9
54	Generation of Free Radicals by Emodic Acid and its [d-Lys6]GnRH-conjugate. <i>Photochemistry and Photobiology</i> , 2001, 74, 226.	1.3	24

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55	Two Isoforms of Gonadotropin-Releasing Hormone Are Coexpressed in Neuronal Cell Lines**This work was supported by the Israel Science Foundation, administered by the Israel Academy of Sciences and Humanities.. Endocrinology, 2001, 142, 830-837.	1.4	22
56	Design and Synthesis of Potent Hexapeptide and Heptapeptide Gonadotropin-Releasing Hormone Antagonists by Truncation of a Decapeptide Analogue Sequence. Journal of Medicinal Chemistry, 2000, 43, 2831-2836.	2.9	10
57	Structure-Activity Studies of Reduced-Size Gonadotropin-Releasing Hormone Agonists Derived from the Sequence of an Endothelin Antagonist. Journal of Medicinal Chemistry, 2000, 43, 2824-2830.	2.9	9
58	Solid-State NMR Determination of Peptide Torsion Angles: Applications of 2H-Dephased REDOR. Journal of the American Chemical Society, 2000, 122, 12263-12269.	6.6	29
59	The gonadotropin-releasing hormone family of neuropeptides in the brain of human, bovine and rat: identification of a third isoform. FEBS Letters, 1999, 463, 289-294.	1.3	59
60	Cytotoxic Peptides: Naphthoquinonyl Derivatives of Luteinizing Hormone-Releasing Hormone. International Journal of Peptide Research and Therapeutics, 1998, 5, 421-427.	0.1	0
61	Cytotoxic peptides: Naphthoquinonyl derivatives of luteinizing hormone-releasing hormone. International Journal of Peptide Research and Therapeutics, 1998, 5, 421-427.	0.1	5
62	Novel naphthoquinonyl derivatives: Potential structural components for the synthesis of cytotoxic peptides. International Journal of Peptide Research and Therapeutics, 1996, 3, 263-274.	0.1	18
63	On the Synthesis of Naphthoquinonyl Heterocyclic Amino Acids. Synthesis, 1996, 1996, 1468-1472.	1.2	28