

R Nagaraj

List of Publications by Year in descending order

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50
papers

2,431
citations

201674

27
h-index

206112

48
g-index

52
all docs

52
docs citations

52
times ranked

1861
citing authors

#	ARTICLE	IF	CITATIONS
1	Is Levinthal's question answered after a revisit?. <i>Journal of Biomolecular Structure and Dynamics</i> , 2013, 31, 961-962.	3.5	0
2	Is Protein Folding Still a Challenge?. <i>Journal of Biomolecular Structure and Dynamics</i> , 2011, 28, 639-640.	3.5	5
3	Cyclic Homooligomers of Furanoid Sugar Amino Acids. <i>Journal of Organic Chemistry</i> , 2003, 68, 6257-6263.	3.2	35
4	Host-defense Antimicrobial Peptides: Importance of Structure for Activity. <i>Current Pharmaceutical Design</i> , 2002, 8, 727-742.	1.9	96
5	Tigerinins: Novel Antimicrobial Peptides from the Indian Frog <i>Rana tigerina</i> . <i>Journal of Biological Chemistry</i> , 2001, 276, 2701-2707.	3.4	99
6	Synthesis and conformational studies of peptidomimetics containing a carbocyclic 1,3-diacid. <i>Tetrahedron</i> , 2001, 57, 9169-9175.	1.9	11
7	Synthesis and structural studies of oligomers of 6-amino-2,5-anhydro-6-deoxy-d-mannonic acid. <i>Tetrahedron Letters</i> , 2000, 41, 8167-8171.	1.4	31
8	Antibacterial and Hemolytic Activities of Single Tryptophan Analogs of Indolicidin. <i>Biochemical and Biophysical Research Communications</i> , 2000, 274, 714-716.	2.1	62
9	Synthesis and Conformational Studies of Peptidomimetics Containing Furanoid Sugar Amino Acids and a Sugar Diacid. <i>Journal of Organic Chemistry</i> , 2000, 65, 6441-6457.	3.2	92
10	Biological activities of C-terminal 15-residue synthetic fragment of melittin: design of an analog with improved antibacterial activity. <i>FEBS Letters</i> , 1999, 448, 62-66.	2.8	79
11	Interaction of antimicrobial peptides with biological and model membranes: structural and charge requirements for activity. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1999, 1462, 29-54.	2.6	295
12	Addition and omission analogs of the 13-residue antibacterial and hemolytic peptide PKLLKTFLSKWIG: structural preferences, model membrane binding and biological activities. <i>Chemical Biology and Drug Design</i> , 1999, 53, 47-55.	1.1	11
13	Interaction of indolicidin, a 13-residue peptide rich in tryptophan and proline and its analogues with model membranes. <i>Journal of Biosciences</i> , 1998, 23, 9-13.	1.1	21
14	Folded Conformation in Peptides Containing Furanoid Sugar Amino Acids. <i>Journal of the American Chemical Society</i> , 1998, 120, 12962-12963.	13.7	84
15	Demonstration of endo-cis-(2S,3R)-Bicyclo[2.2.1]hept-5-en-2,3-dicarbonyl Unit as a Reverse-Turn Scaffold and Nucleator of Two-Stranded Parallel β -Sheets: A Design, Synthesis, Crystal Structure, and Self-Assembling Properties of Norborneno Peptide Analogues. <i>Journal of the American Chemical Society</i> , 1998, 120, 8448-8460.	13.7	49
16	Identification of the region that plays an important role in determining antibacterial activity of bovine seminalplasmin. <i>FEBS Letters</i> , 1997, 400, 289-292.	2.8	11
17	Manual solid-phase syntheses of peptides on resins with high loading capacity requiring small volumes of solvents. <i>Journal of Chemical Sciences</i> , 1997, 109, 319-323.	1.5	2
18	Requirements for antibacterial and hemolytic activities in the bovine neutrophil derived 13-residue peptide indolicidin. <i>FEBS Letters</i> , 1996, 395, 48-52.	2.8	110

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19	Design, Synthesis, and Ion-Transport Properties of a Novel Family of Cyclic, Adamantane-Containing Cystine Peptides. <i>Angewandte Chemie International Edition in English</i> , 1996, 35, 1105-1107.	4.4	35
20	Structural and charge requirements for antimicrobial and hemolytic activity in the peptide PKLLETFLSKWIG, corresponding to the hydrophobic region of the antimicrobial protein bovine seminalplasmin. <i>International Journal of Peptide and Protein Research</i> , 1995, 46, 166-173.	0.1	18
21	Design of 16-residue peptides possessing antimicrobial and hemolytic activities or only antimicrobial activity from an inactive peptide. <i>International Journal of Peptide and Protein Research</i> , 1995, 46, 480-486.	0.1	24
22	Cell-lytic and antibacterial peptides that act by perturbing the barrier function of membranes: facets of their conformational features, structure-function correlations and membrane-perturbing abilities. <i>BBA - Biomembranes</i> , 1994, 1197, 109-131.	8.0	229
23	Studies on the synthesis of the toxins, pardaxin, Î±-toxin and their analogues by solid-phase methods. <i>Journal of Chemical Sciences</i> , 1994, 106, 1109-1121.	1.5	1
24	Identification of a second membrane-active 13-residue peptide segment in the antimicrobial protein, bovine seminalplasmin. <i>FEBS Letters</i> , 1993, 328, 239-242.	2.8	10
25	Interaction of the 47-residue antibacterial peptide seminalplasmin and its 13-residue fragment which has antibacterial and hemolytic activities with model membranes. <i>Biochemistry</i> , 1993, 32, 3124-3130.	2.5	20
26	Change of glutamic acid to lysine in a 13-residue antibacterial and hemolytic peptide results in enhanced antibacterial activity without increase in hemolytic activity. <i>Antimicrobial Agents and Chemotherapy</i> , 1992, 36, 2468-2472.	3.2	29
27	The antibacterial peptide seminal plasmin alters permeability of the inner membrane of <i>E. coli</i> . <i>FEBS Letters</i> , 1992, 303, 265-268.	2.8	22
28	Mass spectra of t-butylloxycarbonyl (BOC)-protected peptides. <i>Organic Mass Spectrometry</i> , 1990, 25, 97-100.	1.3	0
29	A synthetic 13-residue peptide corresponding to the hydrophobic region of bovine seminalplasmin has antibacterial activity and also causes lysis of red blood cells. <i>Journal of Biological Chemistry</i> , 1990, 265, 10438-10442.	3.4	38
30	A synthetic 13-residue peptide corresponding to the hydrophobic region of bovine seminalplasmin has antibacterial activity and also causes lysis of red blood cells. <i>Journal of Biological Chemistry</i> , 1990, 265, 10438-42.	3.4	32
31	Perturbation of the lipid bilayer of model membranes by synthetic signal peptides. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 1987, 903, 465-472.	2.6	19
32	Membrane channel-forming polypeptides. Aqueous phase aggregation and membrane-modifying activity of synthetic fluorescent alamethicin fragments. <i>Journal of Biological Chemistry</i> , 1982, 257, 2170-2176.	3.4	30
33	Conformations of synthetic alamethicin fragments. Evidence for 310 helical folding from 270-MHz hydrogen-1 nuclear magnetic resonance and circular dichroism studies. <i>Biochemistry</i> , 1981, 20, 2828-2835.	2.5	70
34	Determination of beta-turn conformation by laser Raman spectroscopy. <i>Biophysical Journal</i> , 1981, 36, 509-517.	0.5	40
35	Alamethicin and synthetic peptide fragments as uncouplers of mitochondrial oxidative phosphorylation. Effect of chain length and change. <i>Biochemical and Biophysical Research Communications</i> , 1981, 98, 548-555.	2.1	33
36	Hydrophobic channels in crystals of an Î±-aminoisobutyric acid pentapeptide. <i>Biochemical and Biophysical Research Communications</i> , 1981, 103, 898-904.	2.1	30

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37	Solution phase synthesis of alamethicin I. Tetrahedron, 1981, 37, 1263-1270.	1.9	61
38	Racemization at proline residues during peptide bond formation. Tetrahedron, 1981, 37, 2001-2005.	1.9	21
39	H N.M.R. STUDIES OF PROTECTED α -AMINOISOBUTYRIC ACID CONTAINING PEPTIDES. International Journal of Peptide and Protein Research, 1981, 18, 208-213.	0.1	12
40	The crystal structure of benzyloxycarbonyl-(α -aminoisobutyryl) ₂ -L-alanyl methyl ester. Acta Crystallographica Section B: Structural Crystallography and Crystal Chemistry, 1980, 36, 107-110.	0.4	26
41	Cation translocating effects of alamethicin and its synthetic fragments in lipid membranes. FEBS Letters, 1980, 121, 365-368.	2.8	25
42	Infrared studies on the conformation of synthetic alamethicin fragments and model peptides containing α -aminoisobutyric acid. Biochemistry, 1980, 19, 425-431.	2.5	116
43	ROTATIONAL ISOMERISM ABOUT THE C _α -CO BOND IN PROLINE DERIVATIVES. International Journal of Peptide and Protein Research, 1980, 16, 291-298.	0.1	19
44	Crystal and molecular structure of benzyloxycarbonyl- α -aminoisobutyryl-L-prolyl methylamide: The observation of an X ₂ -Pro ₃ Type III β -Turn. Biopolymers, 1979, 18, 1635-1646.	2.4	70
45	Infrared spectroscopy as a probe for the development of secondary structure in the amino-terminal segment of alamethicin. FEBS Letters, 1979, 100, 244-248.	2.8	31
46	Enkephalin analogs. Introduction of stereochemical constraints, metal binding sites and fluorescent groups. FEBS Letters, 1979, 106, 271-274.	2.8	13
47	Fluorescent hydrophobic peptide fragments of emerimicin. Models for the study of peptide aggregation and interactions with lipids and proteins. Biochemical and Biophysical Research Communications, 1979, 89, 1041-1049.	2.1	16
48	Stereochemically constrained linear peptides. Conformations of peptides containing α -aminoisobutyric acid. Journal of the American Chemical Society, 1979, 101, 16-20.	13.7	137
49	A stereochemically-constrained enkephalin analog. FEBS Letters, 1978, 96, 273-276.	2.8	17
50	The crystal and molecular structure of the amino terminal tetrapeptide of alamethicin. A novel 310 helical conformation. Biochemical and Biophysical Research Communications, 1977, 79, 292-298.	2.1	94