Snezna Rogelj

List of Publications by Year in descending order

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		201385	133063
59	5,366	27	59
papers	citations	h-index	g-index
59	59	59	5512
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	A toxicological study on photo-degradation products of environmental ibuprofen: Ecological and human health implications. Ecotoxicology and Environmental Safety, 2020, 188, 109892.	2.9	29
2	Synergistic action of substituted indole derivatives and clinically used antibiotics against drug-resistant bacteria. Future Microbiology, 2020, 15, 579-590.	1.0	5
3	Microtubuleâ€Targeting 7â€Deazahypoxanthines Derived from Marine Alkaloid Rigidins: Exploration of the N3 and N9 Positions and Interaction with Multidrugâ€Resistance Proteins. ChemMedChem, 2019, 14, 322-333.	1.6	5
4	Photo-physical properties of substituted 2,3-distyryl indoles: Spectroscopic, computational and biological insights. Journal of Photochemistry and Photobiology A: Chemistry, 2019, 376, 73-79.	2.0	1
5	Photoactivated 2,3-distyrylindoles kill multi-drug resistant bacteria. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 1879-1886.	1.0	13
6	Spectroscopic Study of a Photoactive Antibacterial Agent: 2,3-Distyrylindole. Journal of Physical Chemistry A, 2018, 122, 937-945.	1.1	1
7	Desymmetrization of Cyclopropenes via the Potassium-Templated Diastereoselective 7- <i>exo</i> - <i>trig</i> Cycloaddition of Tethered Amino Alcohols toward Enantiopure Cyclopropane-Fused Oxazepanones with Antimycobacterial Activity. Journal of Organic Chemistry, 2018. 83. 5650-5664.	1.7	8
8	Abiotic degradation and environmental toxicity of ibuprofen: Roles of mineral particles and solar radiation. Water Research, 2018, 131, 22-32.	5. 3	45
9	Fate, Transformation, and Toxicological Impacts of Pharmaceutical and Personal Care Products in Surface Waters. Environmental Health Insights, 2018, 12, 117863021879583.	0.6	18
10	Metal-Templated Assembly of Cyclopropane-Fused Diazepanones and Diazecanones via <i>exo</i> - <i>trig</i> Nucleophilic Cyclization of Cyclopropenes with Tethered Carbamates. Journal of Organic Chemistry, 2018, 83, 13743-13753.	1.7	8
11	Design, synthesis, and evaluation of cystargolide-based \hat{l}^2 -lactones as potent proteasome inhibitors. European Journal of Medicinal Chemistry, 2018, 157, 962-977.	2.6	9
12	A total synthesis of (â^')-hortonone C. Tetrahedron, 2017, 73, 359-364.	1.0	4
13	5,10b-Ethanophenanthridine amaryllidaceae alkaloids inspire the discovery of novel bicyclic ring systems with activity against drug resistant cancer cells. European Journal of Medicinal Chemistry, 2016, 120, 313-328.	2.6	16
14	Novel Microtubule-Targeting 7-Deazahypoxanthines Derived from Marine Alkaloid Rigidins with Potent in Vitro and in Vivo Anticancer Activities. Journal of Medicinal Chemistry, 2016, 59, 480-485.	2.9	17
15	New method for the synthesis of ammonium salts of O,O'-alkylenedithiophosphoric acid and octathiotetraphosphetane. Crystal structure features's of diethylammonium salt of O,O'-propylenedithiophosphoric acid. Phosphorus, Sulfur and Silicon and the Related Elements, 2016, 191, 405-410.	0.8	11
16	Synthetic and Biological Studies of Sesquiterpene Polygodial: Activity of 9â€Epipolygodial against Drugâ€Resistant Cancer Cells. ChemMedChem, 2015, 10, 2014-2026.	1.6	22
17	Expanding our Understanding of Sequence-Function Relationships of Type II Polyketide Biosynthetic Gene Clusters: Bioinformatics-Guided Identification of Frankiamicin A from Frankia sp. EAN1pec. PLoS ONE, 2015, 10, e0121505.	1.1	25
18	Activity of 2-Aryl-2-(3-indolyl)acetohydroxamates against Drug-Resistant Cancer Cells. Journal of Medicinal Chemistry, 2015, 58, 2206-2220.	2.9	46

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19	Jonquailine, a new pretazettine-type alkaloid isolated from Narcissus jonquilla quail, with activity against drug-resistant cancer. Fìtoterapìâ, 2015, 102, 41-48.	1.1	23
20	Total synthesis and absolute stereochemistry of the proteasome inhibitors cystargolides A and B. Organic and Biomolecular Chemistry, 2015, 13, 10127-10130.	1.5	7
21	Wittig derivatization of sesquiterpenoid polygodial leads to cytostatic agents with activity against drug resistant cancer cells and capable of pyrrolylation of primary amines. European Journal of Medicinal Chemistry, 2015, 103, 226-237.	2.6	16
22	Sol-Generating Chemical Vapor into Liquid (SG-CViL) deposition $\hat{a} \in \text{``a facile method for encapsulation of diverse cell types in silica matrices. Journal of Materials Chemistry B, 2015, 3, 1032-1041.}$	2.9	1
23	Lipophilic prodrug conjugates allow facile and rapid synthesis of high-loading capacity liposomes without the need for post-assembly purification. Journal of Liposome Research, 2015, 25, 232-260.	1.5	7
24	Synthetic and Biological Studies of Tubulin Targeting C2â€Substituted 7â€Deazahypoxanthines Derived from Marine Alkaloid Rigidins. ChemMedChem, 2014, 9, 1428-1435.	1.6	29
25	Synthesis and Some Properties of Transition Metal Complexes Based on the Octathiophophetane Ammonium Salts. Heteroatom Chemistry, 2014, 25, 434-441.	0.4	1
26	Synthesis and biological evaluation of unnatural derivatives of narciclasine: 7-aza-nornarciclasine and its N-oxide. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 4236-4238.	1.0	23
27	A catalytic approach to the MH-031 lactone: application to the synthesis of geralcin analogs. Tetrahedron Letters, 2014, 55, 6266-6268.	0.7	10
28	Exploring Natural Product Chemistry and Biology with Multicomponent Reactions. 5. Discovery of a Novel Tubulin-Targeting Scaffold Derived from the Rigidin Family of Marine Alkaloids. Journal of Medicinal Chemistry, 2013, 56, 6886-6900.	2.9	45
29	Extracellular protein disulfide isomerase regulates ligand-binding activity of $\hat{l}\pm M\hat{l}^22$ integrin and neutrophil recruitment during vascular inflammation. Blood, 2013, 121, 3789-3800.	0.6	111
30	Green Chemistry. Reaction of Elemental Phosphorus (P ₄) and Elemental Sulfur with Protonodonor Reagents: New Methods for the Synthesis of Ammonium Salts of ⟨i>S,⟨i>S,∂i>â€∂ialkyltetrathiophosphoric Acids and Octathiotetraphosphetane. Heteroatom Chemistry, 2013, 24, 163-167.	0.4	8
31	Antiproliferative activity of 2,3-disubstituted indoles toward apoptosis-resistant cancers cells. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 3277-3282.	1.0	9
32	Bulbispermine: A Crinineâ€Type Amaryllidaceae Alkaloid Exhibiting Cytostatic Activity toward Apoptosisâ€Resistant Glioma Cells. ChemMedChem, 2012, 7, 815-822.	1.6	33
33	Structural Simplification of Bioactive Natural Products with Multicomponent Synthesis. 3. Fused Uracil-Containing Heterocycles as Novel Topoisomerase-Targeting Agents. Journal of Medicinal Chemistry, 2011, 54, 2012-2021.	2.9	73
34	One-Pot Multicomponent Synthesis of Diversely Substituted 2-Aminopyrroles. A Short General Synthesis of Rigidins A, B, C, and D. Organic Letters, 2011, 13, 1118-1121.	2.4	73
35	Anticancer Properties of an Important Drug Lead Podophyllotoxin Can Be Efficiently Mimicked by Diverse Heterocyclic Scaffolds Accessible via One-Step Synthesis. Journal of Medicinal Chemistry, 2011, 54, 4234-4246.	2.9	60
36	Multicomponent synthesis of 2,3-dihydrochromeno [4,3-d] pyrazolo [3,4-b] pyridine-1,6-diones: a novel heterocyclic scaffold with antibacterial activity. Tetrahedron Letters, 2011, 52, 6643-6645.	0.7	91

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37	Unprecedented C-2 arylation of indole with diazonium salts: Syntheses of 2,3-disubstituted indoles and their antimicrobial activity. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 4720-4723.	1.0	41
38	Antibacterial action of a novel functionalized chitosan-arginine against Gram-negative bacteria. Acta Biomaterialia, 2010, 6, 2562-2571.	4.1	203
39	Structural Simplification of Bioactive Natural Products with Multicomponent Synthesis. 2. Antiproliferative and Antitubulin Activities of Pyrano[3,2- $\langle i\rangle c\langle j\rangle$] pyridones and Pyrano[3,2- $\langle i\rangle c\langle j\rangle$] quinolones. Journal of Medicinal Chemistry, 2008, 51, 2561-2570.	2.9	160
40	Antiproliferative and apoptosis inducing properties of pyrano[3,2-c]pyridones accessible by a one-step multicomponent synthesis. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 3872-3876.	1.0	64
41	Discovery and Investigation of Antiproliferative and Apoptosis-Inducing Properties of New Heterocyclic Podophyllotoxin Analogues Accessible by a One-Step Multicomponent Synthesis. Journal of Medicinal Chemistry, 2007, 50, 5183-5192.	2.9	135
42	Three-component synthesis and anticancer evaluation of polycyclic indenopyridines lead to the discovery of a novel indenoheterocycle with potent apoptosis inducing properties. Organic and Biomolecular Chemistry, 2007, 5, 3865.	1.5	101
43	Structural simplification of bioactive natural products with multicomponent synthesis: Dihydropyridopyrazole analogues of podophyllotoxin. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 1381-1385.	1.0	43
44	Design of a Highly Sensitive and Specific Nucleotide Sensor Based on Photon Upconverting Particles. Journal of the American Chemical Society, 2006, 128, 12410-12411.	6.6	235
45	Synthesis and Biological Evaluation of Aromatic Analogues of Conduritol F,l-chiro-Inositol, and Dihydroconduritol F Structurally Related to the Amaryllidaceae Anticancer Constituents. Journal of Organic Chemistry, 2006, 71, 5694-5707.	1.7	43
46	Sensitive, real-time PCR detects low-levels of contamination by Legionella pneumophila in commercial reagents. Molecular and Cellular Probes, 2006, 20, 147-153.	0.9	52
47	An immuno-PCR method for detecting Bacillus thuringiensis Cry1Ac toxin. Journal of Immunological Methods, 2006, 308, 109-115.	0.6	39
48	Rapid detection of Escherichia coli O157:H7 by immunomagnetic separation and real-time PCR. International Journal of Food Microbiology, 2005, 99, 47-57.	2.1	155
49	Single-Walled Carbon Nanotube Purification, Pelletization, and Surfactant-Assisted Dispersion:Â A Combined TEM and Resonant Micro-Raman Spectroscopy Study. Journal of Physical Chemistry B, 2005, 109, 4455-4463.	1.2	70
50	A highly sensitive immuno-PCR assay for detecting Group A Streptococcus. Journal of Immunological Methods, 2003, 279, 101-110.	0.6	49
51	Sulfhydryl Regulation of L-Selectin Shedding: Phenylarsine Oxide Promotes Activation-Independent L-Selectin Shedding from Leukocytes. Journal of Immunology, 2000, 164, 4120-4129.	0.4	104
52	Enzyme Destruction by a Protease Contaminant in Bacitracin. Biochemical and Biophysical Research Communications, 2000, 273, 829-832.	1.0	29
53	Enhanced aggregation of human neutrophils by MnCl2 or DTT differentiates the roles of L-selectin and Î ² 2-integrins. Journal of Leukocyte Biology, 1996, 60, 356-364.	1.5	11
54	Evidence for a third component in neutrophil aggregation: potential roles of O-linked glycoproteins as L-selectin counter-structures. Journal of Leukocyte Biology, 1995, 58, 510-518.	1.5	21

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55	Construction and expression of transforming gene resulting from fusion of basic fibroblast growth factor gene with signal peptide sequence. Methods in Enzymology, 1991, 198, 117-124.	0.4	2
56	Cyclic adenosine monophosphate-mediated induction of F9 teratocarcinoma differentiation in the absence of retinoic acid. Journal of Cellular Physiology, 1990, 143, 205-212.	2.0	9
57	Characterization of tumors produced by signal peptide-basic fibroblast growth factor-transformed cells. Journal of Cellular Biochemistry, 1989, 39, 13-23.	1.2	29
58	A human DNA segment with properties of the gene that predisposes to retinoblastoma and osteosarcoma. Nature, 1986, 323, 643-646.	13.7	2,853
59	The effect of retinoic acid on cyclic-AMP-binding proteins in mouse melanoma cells. FEBS Journal, 1984, 139, 351-357.	0.2	15