

Ravindra K Rawal

List of Publications by Year in descending order

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97
papers

3,531
citations

182225

30
h-index

169272

56
g-index

107
all docs

107
docs citations

107
times ranked

4945
citing authors

#	ARTICLE	IF	CITATIONS
1	RdRp (RNA-dependent RNA polymerase): A key target providing anti-virals for the management of various viral diseases. <i>Journal of Molecular Structure</i> , 2022, 1250, 131756.	1.8	16
2	Identifying novel putative ERK1/2 inhibitors via hybrid scaffold hopping â€“FBDD approach. <i>Journal of Biomolecular Structure and Dynamics</i> , 2021, , 1-16.	2.0	8
3	Vitamin D and immuno-pathology of COVID-19: many interactions but uncertain therapeutic benefits. <i>Expert Review of Anti-Infective Therapy</i> , 2021, 19, 1245-1258.	2.0	8
4	Synthetic and Medicinal Perspective of Fused-Thiazoles as Anticancer Agents. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2021, 21, 1379-1402.	0.9	21
5	Recent Nanocarrier Approaches for Targeted Drug Delivery in Cancer Therapy. <i>Current Molecular Pharmacology</i> , 2021, 14, 350-366.	0.7	9
6	Structure based designing of thiazolidinone-pyrimidine derivatives as ERK2 inhibitors: Synthesis and in vitro evaluation. <i>SAR and QSAR in Environmental Research</i> , 2021, 32, 793-816.	1.0	2
7	Agonist, antagonist and signaling modulators of ABA receptor for agronomic and post-harvest management. <i>Plant Physiology and Biochemistry</i> , 2020, 148, 10-25.	2.8	26
8	Identifying the mechanisms of Î±-synuclein-mediated cytotoxicity in Parkinsonâ€™s disease: new insights from a bioinformatics-based approach. <i>Future Neurology</i> , 2020, 15, .	0.9	4
9	Green synthetic strategies toward thiazoles: a sustainable approach. <i>Chemistry of Heterocyclic Compounds</i> , 2020, 56, 445-454.	0.6	12
10	CuI/DBUâ€“Mediated MBH Reaction of Isatins: A Convenient Synthesis of 3â€“Substitutedâ€“Hydroxyâ€“oxindole. <i>ChemistrySelect</i> , 2020, 5, 3048-3051.	0.7	12
11	An update on chemical classes targeting ERK1/2 for the management of cancer. <i>Future Medicinal Chemistry</i> , 2020, 12, 593-611.	1.1	7
12	Development and characterisation of clobetasol propionate loaded Squarticles as a lipid nanocarrier for treatment of plaque psoriasis. <i>Journal of Microencapsulation</i> , 2020, 37, 341-354.	1.2	13
13	A Summary of Viral Targets and Recently Released PDB IDs of SARS-CoV-2. <i>The Open Virology Journal</i> , 2020, 14, 7-8.	1.8	3
14	Genus Calotropis: A Hub of Medicinally Active Phytoconstituents. <i>Current Traditional Medicine</i> , 2020, 6, 312-331.	0.1	7
15	Isolation of Sinapic Acid from <i>Habenaria intermedia</i> D. Don: A New Chemical Marker for the Identification of Adulteration and Substitution. <i>Current Traditional Medicine</i> , 2020, 6, 380-387.	0.1	4
16	Diversified Synthetic Strategies for Pyrroloindoles: An Overview. <i>Journal of Heterocyclic Chemistry</i> , 2019, 56, 2318-2332.	1.4	17
17	Role of sulphur-heterocycles in medicinal chemistry: An update. <i>European Journal of Medicinal Chemistry</i> , 2019, 180, 486-508.	2.6	226
18	Coumarin Hybrids: Promising Scaffolds in the Treatment of Breast Cancer. <i>Mini-Reviews in Medicinal Chemistry</i> , 2019, 19, 1443-1458.	1.1	16

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19	Structure Based Drug Design: Clinically Relevant HIV-1 Integrase Inhibitors. <i>Current Topics in Medicinal Chemistry</i> , 2019, 18, 2664-2680.	1.0	10
20	Diversity-Oriented Synthetic Approaches for Furoindoline: A Review. <i>Current Organic Synthesis</i> , 2019, 16, 342-368.	0.7	14
21	Enasidenib: First Mutant IDH2 Inhibitor for the Treatment of Refractory and Relapsed Acute Myeloid Leukemia. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2019, 18, 1936-1951.	0.9	21
22	Molecular Docking Studies on Isocytosine Analogues as Xanthine Oxidase Inhibitors. <i>Drug Research</i> , 2018, 68, 395-402.	0.7	9
23	RP-HPLC and UV Method Development for Simultaneous Estimation of Doxofylline, Montelukast and Levocetirizine Dihydrochloride in Pharmaceutical Dosages Form. <i>Analytical Chemistry Letters</i> , 2018, 8, 195-204.	0.4	1
24	Oxaliplatin for Colorectal Cancer Therapy: A Review. <i>Clinical Cancer Drugs</i> , 2018, 5, 13-27.	0.3	5
25	Development and Validation of UV-Spectrophotometric and RP-HPLC Methods for Simultaneous Estimation of Fexofenadine Hydrochloride, Montelukast Sodium and Ambroxol Hydrochloride in Tablet Dosage Form. <i>Analytical Chemistry Letters</i> , 2018, 8, 829-843.	0.4	5
26	Natural products and their derivatives as cyclooxygenase-2 inhibitors. <i>Future Medicinal Chemistry</i> , 2018, 10, 2471-2492.	1.1	23
27	Metal-catalyzed synthetic strategies toward coumarin derivatives. <i>Chemistry of Heterocyclic Compounds</i> , 2018, 54, 280-291.	0.6	17
28	Drug metabolizing enzymes and their inhibitors' role in cancer resistance. <i>Biomedicine and Pharmacotherapy</i> , 2018, 105, 53-65.	2.5	73
29	Applications of various analytical techniques in quality control of pharmaceutical excipients. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2018, 157, 122-136.	1.4	35
30	Pyrrolopyrimidines: An update on recent advancements in their medicinal attributes. <i>European Journal of Medicinal Chemistry</i> , 2018, 157, 503-526.	2.6	47
31	Polysaccharides based nanomaterials for targeted anti-cancer drug delivery. <i>Journal of Drug Targeting</i> , 2017, 25, 1-16.	2.1	89
32	$\text{I}^2\text{-CD/CuI}$ catalyzed regioselective synthesis of iodo substituted 1,2,3-triazoles, imidazo[1,2-a]-pyridines and benzoimidazo[2,1-b]thiazoles in water and their functionalization. <i>Tetrahedron</i> , 2017, 73, 4295-4306.	1.0	31
33	Design, synthesis and biological evaluation of hydrazone derivatives as anti-proliferative agents. <i>Medicinal Chemistry Research</i> , 2017, 26, 1459-1468.	1.1	5
34	A Highly Stereoselective Chiral Auxiliary-Assisted Reductive Cyclization to Furoindoline. <i>Journal of Heterocyclic Chemistry</i> , 2017, 54, 2696-2702.	1.4	10
35	Recent synthetic and medicinal perspectives of dihydropyrimidinones: A review. <i>European Journal of Medicinal Chemistry</i> , 2017, 132, 108-134.	2.6	193
36	Synthetic and medicinal perspective of thiazolidinones: A review. <i>Bioorganic Chemistry</i> , 2017, 75, 406-423.	2.0	127

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37	Recent synthetic and medicinal perspectives of tryptanthrin. <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 4533-4552.	1.4	112
38	First report of isolation of maleamic acid from natural source <i>Polygonatum cirrhifolium</i> —A potential chemical marker for identification. <i>Journal of Liquid Chromatography and Related Technologies</i> , 2017, 40, 1031-1036.	0.5	3
39	Synthesis, characterization and pharmacological evaluation of (Z)-2-(5-(biphenyl-4-yl)-3-(1-(imino)ethyl)-2,3-dihydro-1,3,4-oxadiazol-2-yl)phenol derivatives as potent antimicrobial and antioxidant agents. <i>Arabian Journal of Chemistry</i> , 2017, 10, S1022-S1031.	2.3	17
40	Chemical and Medicinal Versatility of Substituted 1,4-Dihydropyridines. <i>Current Bioactive Compounds</i> , 2017, 13, 109-120.	0.2	19
41	ICP-MS: Analytical Method for Identification and Detection of Elemental Impurities. <i>Current Drug Discovery Technologies</i> , 2017, 14, 106-120.	0.6	37
42	A convenient synthesis of 4-alkyl-3-benzoylpyrroles from α,β -unsaturated ketones and tosylmethyl isocyanide. <i>Tetrahedron Letters</i> , 2016, 57, 2315-2319.	0.7	23
43	$\text{In}(\text{OTf})_3$ catalysed an expeditious synthesis of β -carboline—imidazo[1,2-a]pyridine and imidazo[1,2-a]pyrazine conjugates. <i>RSC Advances</i> , 2016, 6, 43881-43891.	1.7	28
44	$\text{In}(\text{OTf})_3 \cdot \text{HBF}_4$ Assisted Multicomponent Approach for One-Pot Synthesis of Pyrazolopyridinone Fused Imidazopyridines. <i>ChemistrySelect</i> , 2016, 1, 4696-4703.	0.7	20
45	Water-Promoted Regiospecific Azidolysis and Copper-Catalyzed Azide—Alkyne Cycloaddition: One-Pot Synthesis of 3-Hydroxy-1-alkyl-3-[(4-aryl/alkyl-1H-1,2,3-triazol-1-yl)methyl]indolin-2-ones. <i>Journal of Organic Chemistry</i> , 2016, 81, 9757-9764.	1.7	40
46	Natural product inspired design and synthesis of β -carboline and β -lactone based molecular hybrids. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 8154-8166.	1.5	31
47	AlCl_3 /Cyclohexane Mediated Electrophilic Activation of Isothiocyanates: An Efficient Synthesis of Thioamides. <i>ChemistrySelect</i> , 2016, 1, 3228-3231.	0.7	18
48	ZrCl_4 Catalysed Diastereoselective Synthesis of Spirocarbocyclic Oxindoles via [4+2] Cycloaddition. <i>ChemistrySelect</i> , 2016, 1, 2409-2412.	0.7	12
49	Metal-free 1,3-dipolar cycloaddition approach towards the regioselective synthesis of β -carboline and isoxazole based molecular hybrids. <i>RSC Advances</i> , 2016, 6, 88066-88076.	1.7	12
50	Impact of Artificial Neural Networks in QSAR and Computational Modeling. , 2016, , 153-179.		15
51	Chemistry and Bioactivities of Aristeromycins: An Overview. <i>Current Topics in Medicinal Chemistry</i> , 2016, 16, 3258-3273.	1.0	16
52	Medicinal Attributes of Imidazo[1,2-a]pyridine Derivatives: An Update. <i>Current Topics in Medicinal Chemistry</i> , 2016, 16, 2963-2994.	1.0	92
53	Synthetic and Medicinal Prospective of Structurally Modified Curcumins. <i>Current Topics in Medicinal Chemistry</i> , 2016, 17, 148-161.	1.0	30
54	Simultaneous Estimation of Hydrochlorothiazide, Hydralazine Hydrochloride, and Reserpine Using PCA, NAS, and NAS-PCA. <i>Scientia Pharmaceutica</i> , 2015, 83, 599-610.	0.7	2

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55	Recent Advances in the Chemistry and Biology of Benzothiazoles. <i>Archiv Der Pharmazie</i> , 2015, 348, 155-178.	2.1	109
56	Riociguat as a treatment regime for pulmonary arterial hypertension: a review. <i>Clinical and Experimental Hypertension</i> , 2015, 37, 323-331.	0.5	1
57	Application of RP-HPLC method in dissolution testing and statistical evaluation by NASSAM for simultaneous estimation of tertiary combined dosages forms. <i>Journal of Pharmaceutical Analysis</i> , 2015, 5, 307-315.	2.4	9
58	The critical role of bisphosphonates to target bone cancer metastasis: an overview. <i>Journal of Drug Targeting</i> , 2015, 23, 1-15.	2.1	66
59	Diversity-oriented synthesis of fused-imidazole derivatives via Groebke-Blackburn-Bienayme reaction: a review. <i>Tetrahedron</i> , 2015, 71, 183-232.	1.0	114
60	Mechanism of Adefovir, Tenofovir and Entecavir Resistance: Molecular Modeling Studies of How A Novel Anti-HBV Agent (FMCA) Can Overcome the Drug Resistance. <i>Current Medicinal Chemistry</i> , 2015, 22, 3922-3932.	1.2	10
61	Recent Advances in Polymer Drug Conjugates. <i>Mini-Reviews in Medicinal Chemistry</i> , 2015, 15, 751-761.	1.1	8
62	A Novel Multiple Tyrosine-kinase Targeted Agent to Explore the Future Perspectives of Anti-Angiogenic Therapy for the Treatment of Multiple Solid Tumors: Cabozantinib. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2014, 15, 37-47.	0.9	37
63	Chemometrics: A new scenario in herbal drug standardization. <i>Journal of Pharmaceutical Analysis</i> , 2014, 4, 223-233.	2.4	130
64	Design, synthesis and ex-vivo release studies of colon-specific polyphosphazene-anticancer drug conjugates. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 1104-1114.	1.4	28
65	Chemometrics tools used in analytical chemistry: An overview. <i>Talanta</i> , 2014, 123, 186-199.	2.9	288
66	A Comprehensive Review on Combretastatin Analogues as Tubulin Binding Agents. <i>Current Organic Chemistry</i> , 2014, 18, 2462-2512.	0.9	21
67	A Compendium of Techniques for the Analysis of Pharmaceutical Impurities. <i>Current Pharmaceutical Analysis</i> , 2014, 10, 145-160.	0.3	4
68	A Novel Integrase Targeting Agent to Explore the Future Prospective of HIV Eradication: Dolutegravir. <i>Current HIV Research</i> , 2014, 12, 325-338.	0.2	5
69	2-Fluoro-6-methylene-carbocyclic adenosine phosphoramidate (FMCAP) prodrug: In vitro anti-HBV activity against the lamivudine-entecavir resistant triple mutant and its mechanism of action. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 503-506.	1.0	15
70	Design, synthesis and ex vivo evaluation of colon-specific azo based prodrugs of anticancer agents. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013, 23, 5332-5338.	1.0	30
71	Synthesis and Characterization of 3-Substituted Amides of 17 α -Aza-Dhomo-4-androsten-17-one as Potent 5-Reductase Inhibitors and Antimicrobial Agents. <i>Current Topics in Medicinal Chemistry</i> , 2013, 13, 2047-2061.	1.0	2
72	Design, Synthesis and In-Vitro Cytotoxicity of Novel Platinum (II) Complexes with Phthalate as the Leaving Group. <i>Letters in Drug Design and Discovery</i> , 2013, 10, 872-878.	0.4	5

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73	Structure-Activity Relationship Studies on Clinically Relevant HIV-1 NNRTIs. <i>Current Medicinal Chemistry</i> , 2012, 19, 5364-5380.	1.2	30
74	Synthesis of Entecavir and Its Novel Class of Analogs. <i>Current Protocols in Nucleic Acid Chemistry</i> , 2011, 47, Unit 14.7.1-17.	0.5	5
75	Characterization of immunologic properties of a second HLA-A2 epitope from a granule protease in CML patients and HLA-A2 transgenic mice. <i>Blood</i> , 2011, 118, 2159-2169.	0.6	14
76	Antiviral activity of novel 2- β -fluoro-6- β -methylene-carbocyclic adenosine against wild-type and drug-resistant hepatitis B virus mutants. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 6328-6331.	1.0	13
77	Synthesis and antiviral activity of cyclopropyl-spirocarbocyclic adenosine, (4 R,5 S,6 R,7 R) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tj ETQq1 1 0.784314 rgBT /Overlock 10 <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 3982-3985.	1.0	21
78	Structure-activity relationships of carbocyclic 6-benzylthioinosine analogues as subversive substrates of <i>Toxoplasma gondii</i> adenosine kinase. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 3403-3412.	1.4	16
79	($\hat{\alpha}$) ⁺ -Carbodine: Enantiomeric synthesis and in vitro antiviral activity against various strains of influenza virus including H5N1 (avian influenza) and novel 2009 H1N1 (swine flu). <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010, 20, 2601-2604.	1.0	10
80	Modified H5 promoter improves stability of insert genes while maintaining immunogenicity during extended passage of genetically engineered MVA vaccines. <i>Vaccine</i> , 2010, 28, 1547-1557.	1.7	42
81	Mamu-A*01/Kb transgenic and MHC Class I knockout mice as a tool for HIV vaccine development. <i>Virology</i> , 2009, 387, 16-28.	1.1	2
82	Breaking Immune Tolerance to Granule Proteases with Full-Length Antigen Vaccine in Humanized Transgenic Mice Reveals Alternative Antigen Processing and Immunodominance Hierarchy Applicable to Clinical Immunotherapy. <i>Blood</i> , 2009, 114, 2054-2054.	0.6	0
83	Design and synthesis of 2-(2,6-dibromophenyl)-3-heteroaryl-1,3-thiazolidin-4-ones as anti-HIV agents. <i>European Journal of Medicinal Chemistry</i> , 2008, 43, 2800-2806.	2.6	97
84	A novel approach to evaluate the immunogenicity of viral antigens of clinical importance in HLA transgenic murine models. <i>Immunology Letters</i> , 2008, 120, 108-116.	1.1	8
85	Non-nucleoside inhibitors of the hepatitis C virus NS5B RNA-dependant RNA polymerase: 2-Aryl-3-heteroaryl-1,3-thiazolidin-4-one derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 6110-6114.	1.0	36
86	2-(2,6-dihalo-phenyl)-3-heteroaryl-2-methyl-1,3-thiazolidin-4-ones: Anti-HIV agents. <i>Chemical Biology and Drug Design</i> , 2008, 72, 147-154.	1.5	21
87	Design, synthesis, and evaluation of 2-aryl-3-heteroaryl-1,3-thiazolidin-4-ones as anti-HIV agents. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 1725-1731.	1.4	164
88	Synthesis and evaluation of 2-(2,6-dihalophenyl)-3-pyrimidinyl-1,3-thiazolidin-4-one analogues as anti-HIV-1 agents. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 3134-3142.	1.4	119
89	Molecular Surface Features in Modeling the HIV-1 RT Inhibitory Activity of 2-(2,6-Disubstituted) Tj ETQq1 1 0.784314 rgBT /Overlock 10 398-406.	1.5	11
90	Synthesis and Biological Evaluation of 2, 3-Diaryl substituted-1, 3-thiazolidin-4-ones as Anti-HIV Agents. <i>Medicinal Chemistry</i> , 2007, 3, 355-363.	0.7	22

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91	2-(2,6-Dichlorophenyl)-3-(quinolin-2-yl)thiazolidin-4-one. Acta Crystallographica Section E: Structure Reports Online, 2006, 62, o5349-o5351.	0.2	1
92	Molecular docking studies on 4-thiazolidinones as HIV-1 RT inhibitors. Journal of Molecular Modeling, 2006, 13, 155-161.	0.8	21
93	2-(Aryl)-3-furan-2-ylmethyl-thiazolidin-4-ones as selective HIV-RT Inhibitors. Bioorganic and Medicinal Chemistry, 2005, 13, 6771-6776.	1.4	158
94	Synthesis and QSAR Studies on Thiazolidinones as Anti-HIV Agents. Combinatorial Chemistry and High Throughput Screening, 2005, 8, 439-443.	0.6	28
95	Topological Descriptors in Modeling the HIV Inhibitory Activity of 2-Aryl-3- pyridyl-thiazolidin-4-ones. Combinatorial Chemistry and High Throughput Screening, 2005, 8, 431-437.	0.6	36
96	An expeditious synthesis of thiazolidinones and tetathiazanones. Journal of Chemical Research, 2004, 2004, 368-369.	0.6	51
97	CP-MLR/PLS Directed Structure-Activity Modeling of the HIV-1 RT Inhibitory Activity of 2,3-Diaryl-1,3-thiazolidin-4-ones. QSAR and Combinatorial Science, 2004, 23, 234-244.	1.5	58