

Balraj Saini

List of Publications by Year in descending order

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14
papers

246
citations

1162367

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1125271

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14
times ranked

262
citing authors

#	ARTICLE	IF	CITATIONS
1	Heparanase Inhibitors in Cancer Progression: Recent Advances. <i>Current Pharmaceutical Design</i> , 2021, 27, 43-68.	0.9	7
2	Chemical Constituents and Biological Activities of <i>Cordia Myxa</i> L.: A Review. <i>Natural Products Journal</i> , 2021, 11, .	0.1	0
3	Natural Fused Heterocyclic Flavonoids: Potent Candidates as Anti-Inflammatory and Anti-Allergic Agents in the Treatment of Asthma. <i>Current Bioactive Compounds</i> , 2021, 17, 28-40.	0.2	2
4	Prospecting the Intricate Role of Novel and Potent Biomarkers in Schizophrenia. <i>Current Topics in Medicinal Chemistry</i> , 2021, 21, 1441-1456.	1.0	2
5	Medicinal Potential of Heterocyclic Compounds from Diverse Natural Sources for the Management of Cancer. <i>Mini-Reviews in Medicinal Chemistry</i> , 2020, 20, 942-957.	1.1	12
6	Nitrogen-Containing Heterocycles as Anticancer Agents: An Overview. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2020, 20, 2150-2168.	0.9	120
7	Forced degradation, LC-UV, MSn and LC-MS-TOF studies on azilsartan: Identification of a known and three new degradation impurities. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2016, 120, 202-211.	1.4	13
8	Identification of Four New Degradation Products of Epirubicin Through Forced Degradation, LC-UV, MS ⁿ and LC-MS-TOF Studies. <i>Journal of Chromatographic Science</i> , 2015, 53, bmv083.	0.7	1
9	Isolation and characterization of a degradation product in leflunomide and a validated selective stability-indicating HPLC-UV method for their quantification. <i>Journal of Pharmaceutical Analysis</i> , 2015, 5, 207-212.	2.4	10
10	Degradation Study on Sulfasalazine and a Validated HPLC-UV Method for its Stability Testing. <i>Scientia Pharmaceutica</i> , 2014, 82, 295-306.	0.7	15
11	Thermal characterization and compatibility studies of itraconazole and excipients for development of solid lipid nanoparticles. <i>Journal of Thermal Analysis and Calorimetry</i> , 2014, 115, 2375-2383.	2.0	27
12	MS ⁿ , LC-MS-TOF and LC-PDA studies for identification of new degradation impurities of bupropion. <i>Biomedical Chromatography</i> , 2013, 27, 1387-1397.	0.8	5
13	ESI-MSn and LC-ESI-MS studies to characterize forced degradation products of bosentan and a validated stability-indicating LC-UV method. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2013, 72, 186-197.	1.4	12
14	Characterization of four new photodegradation products of hydroxychloroquine through LC-PDA, ESI-MSn and LC-MS-TOF studies. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2013, 84, 224-231.	1.4	20