

# Subramaniam AnandaThangadurai

## List of Publications by Year in descending order

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13  
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1477746

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1281420

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13  
times ranked

313  
citing authors

#	ARTICLE	IF	CITATIONS
1	Indazole: a medically important heterocyclic moiety. Medicinal Chemistry Research, 2012, 21, 1509-1523.	1.1	127
2	Synthesis, antimicrobial activity and QSAR studies of new 2,3-disubstituted-3,3a,4,5,6,7-hexahydro-2H-indazoles. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 2960-2964.	1.0	35
3	Development and validation of RP-HPLC method for the estimation of Erlotinib in pharmaceutical formulation. Arabian Journal of Chemistry, 2017, 10, S1138-S1144.	2.3	16
4	Synthesis, antimicrobial evaluation and QSAR studies of 3-ethoxy-4-hydroxybenzylidene/4-nitrobenzylidene hydrazides. Chinese Chemical Letters, 2011, 22, 1293-1296.	4.8	12
5	Development and validation of Ketorolac Tromethamine in eye drop formulation by RP-HPLC method. Arabian Journal of Chemistry, 2017, 10, S928-S935.	2.3	11
6	3,4-Disubstituted-1,2,3,4,5,6,7,8-Octahydroquinazoline-2-thiones: Synthesis, Antimicrobial Evaluation and QSAR Investigations Using Hansch Analysis. Archiv Der Pharmazie, 2008, 341, 231-239.	2.1	7
7	Validation and Method Development of Tadalafil in Bulk and Tablet Dosage Form by RP-HPLC. Drug Research, 2015, 65, 82-85.	0.7	6
8	Synthesis and Antiviral Activity of Dihydropyrimidines - Ciprofloxacin Mannich bases Against Various Viral Strains. Anti-Infective Agents, 2015, 13, 154-165.	0.1	4
9	RP-HPLC Method Development and Validation for the simultaneous estimation of Atazanavir sulphate and Ritonavir in bulk and tablet dosage form. Journal of Pharmaceutical Chemistry, 2014, 1, 50.	0.2	2
10	Method Development and Validation of Hydrochlorothiazide and Quinapril in bulk and tablet dosage form by RP-HPLC. Journal of Pharmaceutical Chemistry, 2014, 1, 10.	0.2	1
11	Formulation, method development and validation of water soluble vitamins B1, B2 & B6 in bulk and tablet dosage form by HPTLC method. Journal of Pharmaceutical Chemistry, 2018, 5, 1-4.	0.2	1
12	Analytical Method Development and Validation of a Dissolution Method for Aliskiren Hemifumarate and Hydrochlorthiazide in Bulk and in Combined Tablet Dosage Form by UPLC. Analytical Chemistry Letters, 2017, 7, 86-96.	0.4	0
13	Development and validation of RP-HPLC/LUV methods for the estimation of Risedronate sodium in pure and pharmaceutical dosage form. Journal of Pharmaceutical Chemistry, 2019, 6, .	0.2	0