## Anamika Sharma

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	<i>In situ</i> Fmoc removal – a sustainable solid-phase peptide synthesis approach. Green Chemistry, 2022, 24, 4887-4896.	4.6	6
2	1,3,5-Triazine as core for the preparation of dendrons. Arkivoc, 2021, 2020, 64-73.	0.3	2
3	The Antiproliferative and Apoptotic Effect of a Novel Synthesized S-Triazine Dipeptide Series, and Toxicity Screening in Zebrafish Embryos. Molecules, 2021, 26, 1170.	1.7	7
4	s-Triazine: A Privileged Structure for Drug Discovery and Bioconjugation. Molecules, 2021, 26, 864.	1.7	31
5	Synthesis, crystal structure, spectroscopic and photophysical studies of novel fluorinated quinazoline derivatives. Journal of Molecular Structure, 2021, 1231, 129951.	1.8	1
6	Exploiting azido-dichloro-triazine as a linker for regioselective incorporation of peptides through their N, O, S functional groups. Bioorganic Chemistry, 2020, 104, 104334.	2.0	3
7	Disulfide-Based Protecting Groups for the Cysteine Side Chain. Organic Letters, 2020, 22, 9644-9647.	2.4	10
8	Novel 4,6-Disubstituted s-Triazin-2-yl Amino Acid Derivatives as Promising Antifungal Agents. Journal of Fungi (Basel, Switzerland), 2020, 6, 237.	1.5	8
9	Insights into the chemistry of the amphibactin–metal (M3+) interaction and its role in antibiotic resistance. Scientific Reports, 2020, 10, 21049.	1.6	3
10	Protocol for synthesis of di- and tri-substituted s-triazine derivatives. MethodsX, 2020, 7, 100825.	0.7	2
11	Synthesis and characterisation of thiobarbituric acid enamine derivatives, and evaluation of their α-glucosidase inhibitory and anti-glycation activity. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 692-701.	2.5	17
12	Breaking a Couple: Disulfide Reducing Agents. ChemBioChem, 2020, 21, 1947-1954.	1.3	39
13	Crystal Structure and Theoretical Investigation of Thiobarbituric Acid Derivatives as Nonlinear Optical (NLO) Materials. Crystals, 2020, 10, 442.	1.0	2
14	Barbiturate- and Thiobarbituarte-Based <i>s</i> -Triazine Hydrazone Derivatives with Promising Antiproliferative Activities. ACS Omega, 2020, 5, 15805-15811.	1.6	21
15	Phenol as a Modulator in the Chemical Reactivity of 2,4,6-Trichloro-1,3,5-triazine: Rules of the Game II. Australian Journal of Chemistry, 2020, 73, 352.	0.5	5
16	s-Triazine: A Multidisciplinary and International Journey. Chemistry Proceedings, 2020, 3, .	0.1	0
17	Scope and Limitations of γ-Valerolactone (GVL) as a Green Solvent to be Used with Base for Fmoc Removal in Solid Phase Peptide Synthesis. Molecules, 2019, 24, 4004.	1.7	20
18	Investigating Triorthogonal Chemoselectivity. Effect of Azide Substitution on the Triazine Core. Organic Letters, 2019, 21, 7888-7892.	2.4	9

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19	Design and synthesis of mono-and di-pyrazolyl-s-triazine derivatives, their anticancer profile in human cancer cell lines, and in vivo toxicity in zebrafish embryos. Bioorganic Chemistry, 2019, 87, 457-464.	2.0	37
20	2-(Dibenzylamino)butane-1,4-dithiol (DABDT), a Friendly Disulfide-Reducing Reagent Compatible with a Broad Range of Solvents. Organic Letters, 2019, 21, 10111-10114.	2.4	7
21	OctaGel Resin - A New PEG-PS-based Solid Support for Solid-Phase Peptide Synthesis. Letters in Organic Chemistry, 2019, 16, 935-940.	0.2	4
22	Efficient Route for Synthesis of Enamines from 1,3-Alkyl-2-Thioxodihydropyrimidine-4,6(1H,5H)-dione Enols. Letters in Organic Chemistry, 2019, 16, 538-540.	0.2	0
23	<i>N</i> â€methylation in amino acids and peptides: Scope and limitations. Biopolymers, 2018, 109, e23110.	1.2	41
24	Crystal structure, spectroscopic studies and theoretical studies of thiobarbituric acid derivatives: understanding the hydrogen-bonding patterns. Acta Crystallographica Section C, Structural Chemistry, 2018, 74, 1703-1714.	0.2	4
25	Perfluorophenyl Derivatives as Unsymmetrical Linkers for Solid Phase Conjugation. Frontiers in Chemistry, 2018, 6, 589.	1.8	5
26	Exploring the Orthogonal Chemoselectivity of 2,4,6-Trichloro-1,3,5-Triazine (TCT) as a Trifunctional Linker With Different Nucleophiles: Rules of the Game. Frontiers in Chemistry, 2018, 6, 516.	1.8	30
27	Exploiting the Thiobarbituric Acid Scaffold for Antibacterial Activity. ChemMedChem, 2018, 13, 1923-1930.	1.6	12
28	Tetrahydropyranyl: A Nonâ€aromatic, Mildâ€Acid‣abile Group for Hydroxyl Protection in Solidâ€Phase Peptide Synthesis. ChemistryOpen, 2017, 6, 206-210.	0.9	4
29	Understanding Tetrahydropyranyl as a Protecting Group in Peptide Chemistry. ChemistryOpen, 2017, 6, 168-177.	0.9	15
30	Novel pyrazolyl-s-triazine derivatives, molecular structure and antimicrobial activity. Journal of Molecular Structure, 2017, 1145, 244-253.	1.8	45
31	Fmoc-Amox, A Suitable Reagent for the Introduction of Fmoc. Organic Process Research and Development, 2017, 21, 1533-1541.	1.3	3
32	Synthesis, Characterization, and Tautomerism of 1,3-Dimethyl Pyrimidine-2,4,6-Trione s-Triazinyl Hydrazine/Hydrazone Derivatives. Journal of Chemistry, 2017, 2017, 1-10.	0.9	7
33	Dual Inhibition of AChE and BChE with the C-5 Substituted Derivative of Meldrum's Acid: Synthesis, Structure Elucidation, and Molecular Docking Studies. Crystals, 2017, 7, 211.	1.0	18
34	Synthesis, Crystal Structure and DFT Studies of 1,3-Dimethyl-5-propionylpyrimidine-2,4,6(1H,3H,5H)-trione. Crystals, 2017, 7, 31.	1.0	6
35	Implications of N-capped urea/thiourea and C-capped 3-(1-piperazinyl)-1,2-benzisothiazole with bridging Gly-Val/Phe-Gly-Val-Pro as therapeutic targets. European Journal of Medicinal Chemistry, 2014, 87, 657-661.	2.6	3
36	tert-Butyl 1,5-bis(4-(benzo[d]isothiazol-3-yl)piperazin-1-yl)-1,5-dioxopentan-2-ylcarbamate urea/thiourea derivatives as potent H+/K+-ATPase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 4096-4098.	1.0	18

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37	Ureas/Thioureas of Benzo[ <i>d</i> ]isothiazole Analog Conjugated Glutamic Acid: Synthesis and Biological Evaluation. Archiv Der Pharmazie, 2013, 346, 359-366.	2.1	15
38	Novel urea and thiourea derivatives of thiazole-glutamic acid conjugate as potential inhibitors of microbes and fungi. Russian Journal of Bioorganic Chemistry, 2013, 39, 656-664.	0.3	4