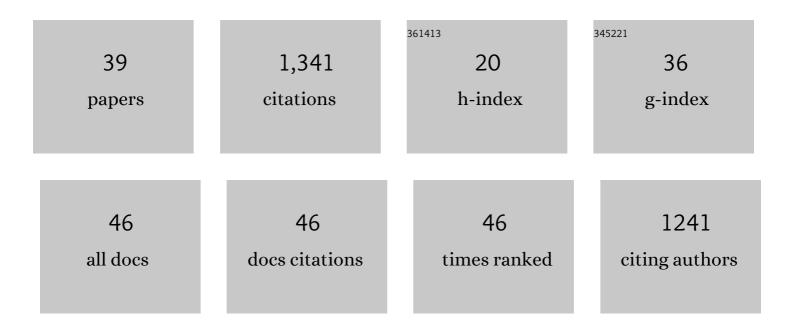


## List of Publications by Year in descending order

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NINC XI

#	Article	IF	CITATIONS
1	A Soluble Base for the Copper-Catalyzed Imidazole N-Arylations with Aryl Halidesâ€. Journal of Organic Chemistry, 2005, 70, 10135-10138.	3.2	166
2	Nitrogen Heterocycles From Furans: The Aza-Achmatowicz Reaction. Synlett, 1998, 1998, 105-114.	1.8	114
3	Discovery and Optimization of a Series of Benzothiazole Phosphoinositide 3-Kinase (PI3K)/Mammalian Target of Rapamycin (mTOR) Dual Inhibitors. Journal of Medicinal Chemistry, 2011, 54, 1789-1811.	6.4	103
4	Structure-Based Design of Novel Class II c-Met Inhibitors: 1. Identification of Pyrazolone-Based Derivatives. Journal of Medicinal Chemistry, 2012, 55, 1858-1867.	6.4	95
5	Design, Synthesis, and Biological Evaluation of Potent c-Met Inhibitors. Journal of Medicinal Chemistry, 2008, 51, 5766-5779.	6.4	86
6	Discovery of a Potent, Selective, and Orally Bioavailable c-Met Inhibitor: 1-(2-Hydroxy-2-methylpropyl)- <i>N</i> -(5-(7-methoxyquinolin-4-yloxy)pyridin-2-yl)-5-methyl-3-oxo-2-phenyl-2,3-d (AMG 458). Journal of Medicinal Chemistry, 2008, 51, 3688-3691.	ihy <b>d</b> r⊕-1 <i< td=""><td>&gt;H8¢i&gt;-pyrazo</td></i<>	>H8¢i>-pyrazo
7	Design of Potent, Orally Available Antagonists of the Transient Receptor Potential Vanilloid 1. Structureâ^'Activity Relationships of 2-Piperazin-1-yl-1H-benzimidazoles. Journal of Medicinal Chemistry, 2006, 49, 3719-3742.	6.4	79
8	Synthesis, chemistry and conformational properties of piperazic acids. Chemical Society Reviews, 1998, 27, 437.	38.1	71
9	Structure-Based Design of Novel Class II c-Met Inhibitors: 2. SAR and Kinase Selectivity Profiles of the Pyrazolone Series. Journal of Medicinal Chemistry, 2012, 55, 1868-1897.	6.4	67
10	Design of a new peptidomimetic agonist for the melanocortin receptors based on the solution structure of the peptide ligand, Ac-Nle-cyclo[Asp-Pro-dPhe-Arg-Trp-Lys]-NH2. Bioorganic and Medicinal Chemistry Letters, 2003, 13, 2337-2340.	2.2	45
11	Elevated Conformational Rigidity in Dipeptides Incorporating Piperazic Acid Derivatives. Journal of the American Chemical Society, 1998, 120, 80-86.	13.7	40
12	Further studies on the chemistry of piperazic acids: New building blocks for β-hydroxy-α-aminoacids through the aza-achmatowicz reaction. Tetrahedron Letters, 1997, 38, 4947-4950.	1.4	39
13	Synthesis of novel melanocortin 4 receptor agonists and antagonists containing a succinamide core. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 377-381.	2.2	38
14	A protection scheme for the preparation of acid chlorides of serine and threonine. Tetrahedron Letters, 1995, 36, 6595-6598.	1.4	34
15	Total Synthesis of Luzopeptin E2. Angewandte Chemie - International Edition, 2000, 39, 2493-2495.	13.8	27
16	Synthesis and evaluation of thiazole carboxamides as vanilloid receptor 1 (TRPV1) antagonists. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 5211-5217.	2.2	25
17	Studies toward Luzopeptins:Â Assembly of the Elusive Serineâ^'PCA Dipeptide. Journal of Organic Chemistry, 1997, 62, 2320-2321.	3.2	24
18	Total synthesis of luzopeptin C. Tetrahedron Letters, 2001, 42, 1907-1909.	1.4	23

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#	Article	IF	CITATIONS
19	Melanocortin subtype-4 receptor agonists containing a piperazine core with substituted aryl sulfonamides. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 1623-1627.	2.2	21
20	Interaction of Cu(II) 3,5-diisopropylsalicylate with human serum albuminan evaluation of spectroscopic data. BioMetals, 1998, 11, 21-26.	4.1	20
21	N-Aryl-γ-lactams as integrin αvβ3 antagonists. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 2905-2909.	2.2	20
22	4-Phenyl tetrahydroisoquinolines as dual norepinephrine and dopamine reuptake inhibitors. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 7219-7222.	2.2	20
23	Synthesis and chemical properties of PCA, an unusual amino acid in luzopeptins. Journal of the Chemical Society Chemical Communications, 1994, , 1867.	2.0	17
24	Regio-controlled synthesis of N-substituted imidazoles. Tetrahedron Letters, 2005, 46, 7315-7319.	1.4	17
25	Studies toward luzopeptins: a key tripeptide subunit selectively deprotectable under neutral conditions. Tetrahedron Letters, 1999, 40, 3693-3696.	1.4	12
26	Noncovalent Interactions of Fluorine with Amide and CH <sub>2</sub> Groups in <i>N</i> -Phenyl γ-Lactams: Covalently Identical Fluorine Atoms in Nonequivalent Chemical Environments. Journal of Organic Chemistry, 2018, 83, 11586-11594.	3.2	12
27	19F NMR spectroscopy as a tool to detect rotations in fluorine substituted phenyl compounds. Tetrahedron, 2020, 76, 131679.	1.9	7
28	Drug interaction of ningetinib and gefitinib involving CYP1A1 and efflux transporters in nonâ€small cell lung cancer patients. British Journal of Clinical Pharmacology, 2021, 87, 2098-2110.	2.4	7
29	A unified approach to peptin antibiotics. Journal of Heterocyclic Chemistry, 1999, 36, 1409-1419.	2.6	6
30	Studies on structural requirements for atropisomerism in N -phenyl γ-lactams. Tetrahedron, 2018, 74, 2991-2998.	1.9	5
31	Facile synthesis of bicyclic 1-arylpyrazol-5-ones. Tetrahedron Letters, 2014, 55, 142-147.	1.4	3
32	Electrostatic effects on 19F NMR chemical shifts in N-phenyl Î <sup>3</sup> -lactam derivatives. Tetrahedron, 2022, 113, 132733.	1.9	3
33	Interaction Of Metal Ions With D-Glucose In Glassy StateA FT-IR Study. , 1989, , .		1
34	Editorial [Hot Topic: Melanocortin Receptor Ligands and Their Potential Therapeutic Uses (Guest) Tj ETQq0 0 0 r	gBT /Overl 2.1	ock 10 Tf 50
35	Selective β-Hydroxyethylation at the N-1 Position of a Pyrazolone: Synthesis of Benzyl 1-(β-Hydroxyethyl)-5-methyl-3-oxo-2-phenyl-2,3-dihydro-1H-pyrazole-4-carboxylate. Synlett, 2008, 2008, 1005-1008.	1.8	1

Practical synthesis of bicyclic pyrazol-5-one derivatives. Tetrahedron Letters, 2017, 58, 46-49.

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#	Article	IF	CITATIONS
37	FT-IR Spectroscopy As A Tool For The Study Of Metal Ions/D- Galactose Complexes. , 1989, , .		Ο
38	Synthesis of Novel Melanocortin 4 Receptor Agonists and Antagonists Containing a Succinamide Core ChemInform, 2004, 35, no.	0.0	0
39	Regio-Controlled Synthesis of N-Substituted Imidazoles ChemInform, 2006, 37, no.	0.0	0