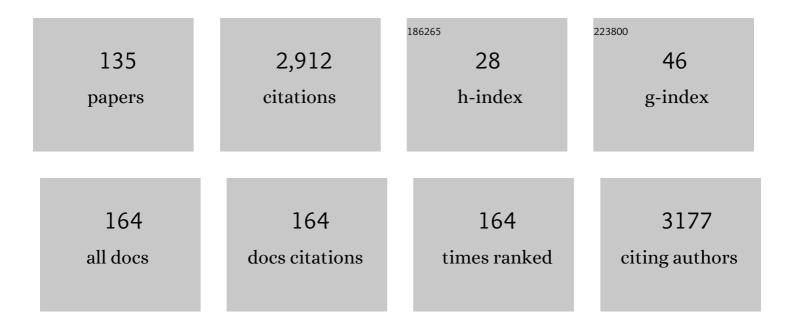
## Patrick Dallemagne

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4715494/publications.pdf Version: 2024-02-01



PATRICK DALLEMACHE

#	Article	IF	CITATIONS
1	Drug repositioning: a brief overview. Journal of Pharmacy and Pharmacology, 2020, 72, 1145-1151.	2.4	185
2	Synthesis, Antimalarial Activity, and Molecular Modeling of New Pyrrolo[1,2-a]quinoxalines, Bispyrrolo[1,2-a]quinoxalines, Bispyrido[3,2-e]pyrrolo[1,2-a]pyrazines, and Bispyrrolo[1,2-a]thieno[3,2-e]pyrazines. Journal of Medicinal Chemistry, 2004, 47, 1997-2009.	6.4	151
3	hERG toxicity assessment: Useful guidelines for drug design. European Journal of Medicinal Chemistry, 2020, 195, 112290.	5.5	121
4	New aromatase inhibitors. Synthesis and biological activity of aryl-substituted pyrrolizine and indolizine derivatives. Bioorganic and Medicinal Chemistry, 2000, 8, 945-955.	3.0	114
5	Novel Multitarget-Directed Ligands (MTDLs) with Acetylcholinesterase (AChE) Inhibitory and Serotonergic Subtype 4 Receptor (5-HT <sub>4</sub> R) Agonist Activities As Potential Agents against Alzheimer's Disease: The Design of Donecopride. Journal of Medicinal Chemistry, 2015, 58, 3172-3187.	6.4	100
6	Design of donecopride, a dual serotonin subtype 4 receptor agonist/acetylcholinesterase inhibitor with potential interest for Alzheimer's disease treatment. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E3825-30.	7.1	96
7	Suzuki-type cross-coupling reaction of 3-iodoindazoles with aryl boronic acids: A general and flexible route to 3-arylindazoles. Tetrahedron, 1999, 55, 6917-6922.	1.9	79
8	Synthesis of new pyrrolo[1,2-a]quinoxalines: potential non-peptide glucagon receptor antagonists. European Journal of Medicinal Chemistry, 1998, 33, 293-308.	5.5	78
9	Therapeutic Potential of 5-HT <sub>6</sub> Receptor Agonists. Journal of Medicinal Chemistry, 2015, 58, 7901-7912.	6.4	72
10	Synthesis and biological evaluation of novel pyrrolopyrrolizinones as anticancer agents. Bioorganic and Medicinal Chemistry, 2006, 14, 8162-8175.	3.0	58
11	Synthesis of novel pyrazolopyrrolopyrazines, potential analogs of sildenafil. Journal of Heterocyclic Chemistry, 2001, 38, 1045-1050.	2.6	54
12	Synthesis and biological evaluation of five-Membered heterocycles fused to cyclopenta[ c ]thiophene as new antitumor agents. Bioorganic and Medicinal Chemistry, 2003, 11, 1161-1167.	3.0	51
13	Multifaceted properties of 1,4-dimethylcarbazoles: Focus on trimethoxybenzamide and trimethoxyphenylurea derivatives as novel human topoisomerase II inhibitors. European Journal of Pharmaceutical Sciences, 2017, 96, 263-272.	4.0	49
14	Modulating 5-HT <sub>4</sub> and 5-HT <sub>6</sub> receptors in Alzheimer's disease treatment. Future Medicinal Chemistry, 2017, 9, 781-795.	2.3	46
15	Pharmacotechnical Development of a Nasal Drug Delivery Composite Nanosystem Intended for Alzheimer's Disease Treatment. Pharmaceutics, 2020, 12, 251.	4.5	43
16	Matrix Metalloproteinases as New Targets in Alzheimer's Disease: Opportunities and Challenges. Journal of Medicinal Chemistry, 2020, 63, 10705-10725.	6.4	42
17	Synergistic effect of acetylcholinesterase inhibition (donepezil) and 5-HT4 receptor activation (RS67333) on object recognition in mice. Behavioural Brain Research, 2012, 230, 304-308.	2.2	39
18	3-(Dipropylamino)-5-hydroxybenzofuro[2,3-f]quinazolin-1(2H)-one (DPA-HBFQ-1) plays an inhibitory role on breast cancer cell growth and progression. European Journal of Medicinal Chemistry, 2016, 107, 275-287.	5.5	39

#	Article	IF	CITATIONS
19	Synthesis and biological evaluation as AChE inhibitors of new indanones and thiaindanones related to donepezil. European Journal of Medicinal Chemistry, 2005, 40, 1222-1245.	5.5	36
20	Chronic activation of 5-HT4 receptors or blockade of 5-HT6 receptors improve memory performances. Behavioural Brain Research, 2015, 293, 10-17.	2.2	36
21	Synthesis and evaluation of cytotoxic activities of new guanidines derived from carbazoles. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 467-472.	2.2	35
22	Synthesis and inÂvitro evaluation of 4-trichloromethylpyrrolo[1,2-a]quinoxalines as new antiplasmodial agents. European Journal of Medicinal Chemistry, 2014, 83, 26-35.	5.5	35
23	Novel multitarget-directed ligands targeting acetylcholinesterase and Ïf1 receptors as lead compounds for treatment of Alzheimer's disease: Synthesis, evaluation, and structural characterization of their complexes with acetylcholinesterase. European Journal of Medicinal Chemistry, 2019, 162, 234-248.	5.5	35
24	Design, synthesis, and pharmacological evaluation of multitarget-directed ligands with both serotonergic subtype 4 receptor (5-HT4R) partial agonist and 5-HT6R antagonist activities, as potential treatment of Alzheimer's disease. European Journal of Medicinal Chemistry, 2016, 121, 283-293.	5.5	33
25	Benzothienoquinazolinones as new multi-target scaffolds: Dual inhibition of human Topoisomerase I and tubulin polymerization. European Journal of Medicinal Chemistry, 2019, 181, 111583.	5.5	32
26	Synthesis and biological evaluation of cyclopenta[ c ]thiophene related compounds as new antitumor agents. Bioorganic and Medicinal Chemistry, 2002, 10, 2185-2191.	3.0	30
27	Evidence for new non-steroidal human aromatase inhibitors and comparison with equine aromatase inhibition for an understanding of the mammalian active site. European Journal of Medicinal Chemistry, 1998, 33, 451-462.	5.5	29
28	Discovery of new thienopyrimidinone derivatives displaying antimalarial properties toward both erythrocytic and hepatic stages of Plasmodium. European Journal of Medicinal Chemistry, 2015, 95, 16-28.	5.5	29
29	First synthesis of arylpyrrolo- and pyrazolopyrrolizinones as useful agents with potential biological interest. Tetrahedron Letters, 2004, 45, 6353-6355.	1.4	27
30	Synthesis and PreliminaryIn Vitro Evaluation of Antimycobacterial Activity of New Pyrrolo[1,2-a]quinoxaline-carboxylic Acid Hydrazide Derivatives. Journal of Enzyme Inhibition and Medicinal Chemistry, 2004, 19, 489-495.	5.2	27
31	Synthesis of New 2-(Aminomethyl)-4-phenylpyrrolo[1,2-a]-quinoxalines and their Preliminary In-vivo Central Dopamine Antagonist Activity Evaluation in Mice. Journal of Pharmacy and Pharmacology, 2010, 52, 1369-1375.	2.4	27
32	Virtual Screening Discovery of New Acetylcholinesterase Inhibitors Issued from CERMN Chemical Library. Journal of Chemical Information and Modeling, 2010, 50, 422-428.	5.4	24
33	Design and synthesis of a new type of non steroidal human aromatase inhibitors. Bioorganic and Medicinal Chemistry Letters, 1998, 8, 1041-1044.	2.2	23
34	Rational design of novel benzisoxazole derivatives with acetylcholinesterase inhibitory and serotoninergic 5-HT4 receptors activities for the treatment of Alzheimer's disease. Scientific Reports, 2020, 10, 3014.	3.3	23
35	Study of substrate specificity of human aromatase by site directed mutagenesis. FEBS Journal, 2002, 269, 1393-1405.	0.2	22
36	Synthesis of a Novel Class of Non-Peptide NK-2 Receptor Ligand, Derived from 1-Phenyl-3-pyrrol-1-ylindan-2-carboxamides. Bioorganic and Medicinal Chemistry, 2002, 10, 1043-1050.	3.0	22

#	Article	IF	CITATIONS
37	Synthesis of new dipyrrolo- and furopyrrolopyrazinones related to tripentones and their biological evaluation as potential kinases (CDKs1–5, GSK-3) inhibitors. European Journal of Medicinal Chemistry, 2009, 44, 708-716.	5.5	22
38	Recent Advances in Phenanthroindolizidine and Phenanthroquinolizidine Derivatives with Anticancer Activities. Anti-Cancer Agents in Medicinal Chemistry, 2015, 15, 1080-1091.	1.7	22
39	One-pot cyclization of alkoxy-3-aminoindan-1-ones Tetrahedron Letters, 1991, 32, 6327-6328.	1.4	21
40	A convenient synthesis of dihydro- and tetrahydro-1,3-thiazine derivatives from β-aryl-β-amino acids. Tetrahedron Letters, 2004, 45, 1503-1505.	1.4	21
41	Indenopyrazole oxime ethers: Synthesis and β1-adrenergic blocking activity. European Journal of Medicinal Chemistry, 2015, 92, 672-681.	5.5	21
42	New piperazine multi-effect drugs prevent neurofibrillary degeneration and amyloid deposition, and preserve memory in animal models of Alzheimer's disease. Neurobiology of Disease, 2019, 129, 217-233.	4.4	21
43	Design, synthesis and biological evaluation of novel indano- and thiaindano-pyrazoles with potential interest for Alzheimer's disease. MedChemComm, 2013, 4, 925.	3.4	20
44	Inhibiting Acetylcholinesterase to Activate Pleiotropic Prodrugs with Therapeutic Interest in Alzheimer's Disease. Molecules, 2019, 24, 2786.	3.8	20
45	A Novel in vivo Anti-amnesic Agent, Specially Designed to Express Both Acetylcholinesterase (AChE) Inhibitory, Serotonergic Subtype 4 Receptor (5-HT4R) Agonist and Serotonergic Subtype 6 Receptor (5-HT6R) Inverse Agonist Activities, With a Potential Interest Against Alzheimer's Disease. Frontiers in Aging Neuroscience, 2019, 11, 148.	3.4	20
46	Pleiotropic prodrugs: Design of a dual butyrylcholinesterase inhibitor and 5-HT6 receptor antagonist with therapeutic interest in Alzheimer's disease. European Journal of Medicinal Chemistry, 2021, 210, 113059.	5.5	20
47	An Efficient Synthesis of New Phenylpyrrolizine and Phenylpyrrolopyrazine Derivatives. Heterocycles, 1993, 36, 2129.	0.7	20
48	Donecopride, a Swiss army knife with potential against Alzheimer's disease. British Journal of Pharmacology, 2020, 177, 1988-2005.	5.4	19
49	An expedient synthesis of 6-arylpiperidine-2,4-diones by chain-extension of β-aryl-β-aminoacids. Tetrahedron Letters, 2001, 42, 8997-8999.	1.4	18
50	Hydrogenative desulphurization of thienopyrrolizinones: An easy and selective access to (Z)-phenethylidenepyrrolizinones with in vitro cytotoxic activity. European Journal of Medicinal Chemistry, 2010, 45, 1146-1150.	5.5	18
51	Screening and evaluation of antiviral compounds against Equid alpha-herpesviruses using an impedance-based cellular assay. Virology, 2019, 526, 105-116.	2.4	18
52	Tripentones: A Promising Series of Potent Anti-Cancer Agents. Anti-Cancer Agents in Medicinal Chemistry, 2009, 9, 369-380.	1.7	18
53	A Convenient Route to 6-Aminocyclopenta[c]thiophen-4-one Derivatives. Heterocycles, 1993, 36, 287.	0.7	17
54	MR 20492 and MR 20494: two indolizinone derivatives that strongly inhibit human aromatase. Journal of Steroid Biochemistry and Molecular Biology, 1999, 70, 59-71.	2.5	17

#	Article	IF	CITATIONS
55	MR22388, a novel anti-cancer agent with a strong FLT-3 ITD kinase affinity. Cancer Letters, 2013, 331, 92-98.	7.2	16
56	New arylhexahydropyrimidinediones: Synthesis, benzodiazepine receptor affinity and anticonvulsant activity. European Journal of Medicinal Chemistry, 1996, 31, 335-339.	5.5	15
57	Efficient synthesis of 2-aryl-6-methyl-2,3-dihydro-1H-pyridin-4-ones. Tetrahedron Letters, 2000, 41, 681-683.	1.4	15
58	First synthesis of methyl 3-amino-4-(het)aryl-1H-pyrrole-2-carboxylates as useful scaffolds in medicinal chemistry. Tetrahedron, 2004, 60, 2267-2270.	1.9	14
59	Synthesis of new 4-[2-(alkylamino)ethylthio]pyrrolo[1,2- <i>a</i> ]quinoxaline and 5-[2-(alkylamino)ethylthio]pyrrolo[1,2- <i>a</i> ]thieno[3,2- <i>e</i> ]pyrazine derivatives, as potential bacterial multidrug resistance pump inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 620-631.	5.2	14
60	Synthesis of dual AChE/5-HT4 receptor multi-target directed ligands. MedChemComm, 2012, 3, 627.	3.4	14
61	First Synthesis of Isothianinhydrin, the Second Thiophene Isostere of Ninhydrin. Synlett, 1999, 1999, 1450-1452.	1.8	13
62	First Synthesis of Racemic Trans Propargylamino-Donepezil, a Pleiotrope Agent Able to Both Inhibit AChE and MAO-B, with Potential Interest against Alzheimer's Disease. Molecules, 2021, 26, 80.	3.8	13
63	A Convenient Rearrangement of 1-Phenylpyrrole-2-carboxaldehydes into Their 3-Isomers. Synthetic Communications, 1994, 24, 1855-1857.	2.1	12
64	One-pot synthesis of novel poly-substituted phenanthrenes. Tetrahedron, 2010, 66, 2803-2808.	1.9	12
65	Nâ€substituted Piperazinopyridylsteroid Derivatives as Abiraterone Analogues Inhibit Growth and Induce Proâ€apoptosis in Human Hormoneâ€independent Prostate Cancer Cell Lines. Chemical Biology and Drug Design, 2013, 82, 620-629.	3.2	12
66	Novel benzylidenephenylpyrrolizinones with pleiotropic activities potentially useful in Alzheimer's disease treatment. European Journal of Medicinal Chemistry, 2016, 114, 365-379.	5.5	12
67	Novel multi target-directed ligands targeting 5-HT4 receptors with in cellulo antioxidant properties as promising leads in Alzheimer's disease. European Journal of Medicinal Chemistry, 2019, 182, 111596.	5.5	12
68	Disproportionality analysis in VigiBase as a drug repositioning method for the discovery of potentially useful drugs in Alzheimer's disease. British Journal of Clinical Pharmacology, 2021, 87, 2830-2837.	2.4	12
69	Therapeutic modulators of the serotonin 5-HT4 receptor: a patent review (2014-present). Expert Opinion on Therapeutic Patents, 2020, 30, 495-508.	5.0	12
70	A Convenient Route to 6-Aminocyclopenta[b]thiophene Derivatives. Heterocycles, 1987, 26, 3233.	0.7	12
71	Synthesis and Cytotoxic Activity against L1210 Leukemia of New Aminocyclopenta(c)thiophenones Chemical and Pharmaceutical Bulletin, 1994, 42, 1605-1608.	1.3	11
72	An efficient route to 6-(het)aryl-2-methyl-2,3-dihydro-1H-pyridin-4-ones as potential nAChRs ligands. Tetrahedron, 2004, 60, 4861-4865.	1.9	11

#	Article	IF	CITATIONS
73	Synthesis and Study of the Stability of 3b,4,4a,5-Tetrahydrothieno[2',3':5,4]cyclopenta[2,3-b]aziridine Derivatives. Heterocycles, 1987, 26, 1449.	0.7	11
74	Pyrrolothieno[1,4]diazepines part III: Synthesis of amino, hydrazino and mercapto derivatives. Journal of Heterocyclic Chemistry, 1996, 33, 87-91.	2.6	10
75	FRIEDEL-CRAFTS ACYLATION WITH MALONIC ACIDS IN POLYPHOSPHORIC ACID. Organic Preparations and Procedures International, 1999, 31, 324-328.	1.3	10
76	Facile Diastereoselective Routes to Highly Functionalized Cyclopenta[c]thiophenes as Useful Scaffolds in Medicinal Chemistry. Synthesis, 2002, 2002, 1091-1095.	2.3	10
77	Benzylphenylpyrrolizinones with Antiâ€amyloid and Radical Scavenging Effects, Potentially Useful in Alzheimer's Disease Treatment. ChemMedChem, 2017, 12, 913-916.	3.2	10
78	A convenient route to new phenyltetrahydroindolizines. Journal of Heterocyclic Chemistry, 1996, 33, 1689-1694.	2.6	9
79	First and mild synthesis of fluorene-9-malonic acid and some substituted derivatives via the intramolecular hydroarylation of 2-phenylbenzylidenemalonic acids. Tetrahedron, 2011, 67, 2548-2554.	1.9	9
80	One-pot synthesis of new aza- and diaza-aminophenanthrenes. Tetrahedron, 2011, 67, 5806-5810.	1.9	9
81	Synthesis of novel 7-oxo and 7-hydroxy trifluoroallocolchicinoids with cytotoxic effect. Bioorganic and Medicinal Chemistry, 2012, 20, 2614-2623.	3.0	9
82	Active Targeted Nanoemulsions for Repurposing of Tegaserod in Alzheimer's Disease Treatment. Pharmaceutics, 2021, 13, 1626.	4.5	9
83	Pyrrolothieno[1,4]diazepines: Synthesis of alkoxy derivatives. Journal of Heterocyclic Chemistry, 1995, 32, 1719-1724.	2.6	8
84	A NEW EFFICIENT SYNTHESIS OF 3-AMINO-1-PHENYLPYRROLE. Organic Preparations and Procedures International, 1995, 27, 236-239.	1.3	8
85	A First Cyclopenta [c] thiophene Dimer as a New Bivalent Potent Cytotoxic Derivative. Journal of Enzyme Inhibition and Medicinal Chemistry, 2002, 17, 439-442.	5.2	8
86	Synthesis and Biological Evaluation of Thienopyrrolizines, a New Family of CDK/GSK-3 Inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2004, 19, 585-593.	5.2	8
87	Synthesis of Novel Pyrazolopyrrolizinones as Prospective Anticancer Agents. Heterocycles, 2006, 68, 2063.	0.7	8
88	An expedient one-pot synthesis of novel 10-substituted 9-aminophenanthrenes. Tetrahedron Letters, 2009, 50, 5704-5708.	1.4	8
89	Intramolecular Cyclisation of β-Aryl-β-Amino Acids in the Design of Novel Heterocyclic Systems with Therapeutic Interest: An Unfailing Source of Diversity. Current Medicinal Chemistry, 2010, 17, 4342-4369.	2.4	8
90	Desirable drug–drug interactions or when a matter of concern becomes a renewed therapeutic strategy. Drug Discovery Today, 2021, 26, 315-328.	6.4	8

#	Article	IF	CITATIONS
91	A New and Efficient Synthesis of 4-Arylimidazolidin-2-ones. Heterocycles, 1991, 32, 1301.	0.7	8
92	Pyrrolothieno[1,4]diazepines Part 1 : Synthesis and Study of the Reaction Pathway. Heterocycles, 1995, 41, 515.	0.7	8
93	Thienocyclopenta[2,3-b]aziridin-5-one: Cleavage in Acidic Medium. Heterocycles, 1992, 34, 1317.	0.7	7
94	Synthesis of new 6â€(4â€chlorophenyl)perhydroâ€1,3â€diazepineâ€2,4â€diones <i>via</i> ureidobutyric acids. Journal of Heterocyclic Chemistry, 1998, 35, 535-539.	2.6	7
95	Synthesis and preliminary behavioural evaluation in mice of new 3-aryl-3-pyrrol-1-ylpropanamides, analogues of FGIN-1–27 and FGIN-1–43. Journal of Pharmacy and Pharmacology, 2010, 53, 1561-1568.	2.4	7
96	A chemical screen identifies two novel small compounds that alter Arabidopsis thaliana pollen tube growth. BMC Plant Biology, 2019, 19, 152.	3.6	7
97	Phenanthrolinic analogs of quinolones show antibacterial activity against M.Âtuberculosis. European Journal of Medicinal Chemistry, 2020, 207, 112821.	5.5	7
98	Two Antagonistic Microtubule Targeting Drugs Act Synergistically to Kill Cancer Cells. Cancers, 2020, 12, 2196.	3.7	7
99	Effective cerebral antihypoxic activity of new aminocyclopentanones. European Journal of Medicinal Chemistry, 1992, 27, 961-965.	5.5	6
100	Synthesis of new thienocyclopenta[3,2â€ <i>d</i> ]â€oxazole and thiazole derivatives. Journal of Heterocyclic Chemistry, 1993, 30, 799-802.	2.6	6
101	Pyirolothieno[1,4]diazepines. Part <b>IV</b> . First synthesis of pyrroloâ€{1,2â€ <i>a</i> ]thieno[2,3â€ <i>f</i> ][1,4]diazepine Derivatives. Journal of Heterocyclic Chemistry, 1996, 33, 1743-1749.	2.6	6
102	Synthesis and CNS Activity of New 3-Amino-3-arylpropionic Acid Derivatives. Pharmacy and Pharmacology Communications, 1999, 5, 217-223.	0.3	6
103	Synthesis, reactivity and biological evaluation of novel halogenated tripentones. Bioorganic and Medicinal Chemistry, 2009, 17, 7783-7788.	3.0	6
104	An unusual boron tribromide-mediated, one-pot bromination/cyclization reaction. Application to the synthesis of a highly strained cyclopenta[1,3]cyclopropa[1,2-b]pyrrolizin-8-one. Tetrahedron Letters, 2013, 54, 1133-1136.	1.4	6
105	6-Sulfonylbenzothiazolones as potential scaffolds for the design of 5-HT6 ligands. European Journal of Medicinal Chemistry, 2015, 92, 807-817.	5.5	6
106	Synthesis and evaluation of novel serotonin 4 receptor radiotracers for single photon emission computed tomography. European Journal of Medicinal Chemistry, 2016, 116, 90-101.	5.5	6
107	Facing the complexity of Alzheimer's disease. Future Medicinal Chemistry, 2020, 12, 175-177.	2.3	6
108	Identification of antiviral compounds against equid herpesvirus-1 using real-time cell assay screening: Efficacy of decitabine and valganciclovir alone or in combination. Antiviral Research, 2020, 183, 104931.	4.1	6

#	Article	IF	CITATIONS
109	Synthesis of 7-Amino-4,5,6,7-tetrahydrothieno[3,4-c]pyrid-4-ones. Heterocycles, 1997, 45, 527.	0.7	6
110	First Synthesis of 1-Phenyl-3-pyrrol-1-ylindan-2-carboxylic Acid, a New Scaffold of Potential Non-peptide Endothelin Receptor Antagonists. Synlett, 1999, 1999, 1263-1264.	1.8	5
111	Synthesis and Initial Results for MAO-B Inhibition by New N-Propargyl-3-pyrrol-1-ylindanamine Derivatives, Analogues of Rasagiline. Journal of Enzyme Inhibition and Medicinal Chemistry, 2003, 18, 147-153.	5.2	5
112	Replication of Equine arteritis virus is efficiently suppressed by purine and pyrimidine biosynthesis inhibitors. Scientific Reports, 2020, 10, 10100.	3.3	5
113	Synthesis of 4-Aminothieno[2,3-c]pyrid-7-one from 4-Aminocyclopenta[b]thiophen-6-one. Heterocycles, 1988, 27, 1637.	0.7	5
114	FIRST SYNTHESIS OF 4-AMINO-4,5-DIHYDRO-1-PHENYLCYCLOPENTA[ b]PYRROL-6(1H)-ONES. Heterocyclic Communications, 1994, 1, .	1.2	4
115	A CONVENIENT SYNTHESIS OF NEW HALOTHIENYL Î <sup>2</sup> -AMINOACIDS AS VERSATILE BUILDING BLOCKS. Organic Preparations and Procedures International, 1997, 29, 488-494.	1.3	4
116	Pyrrolothieno[1,4]diazepines. Part V. Study of their chemical reactivity and first synthesis of oxazino[4,3-c]pyrrolo[1,2-a]thieno[2,3-f] [1,4]diazepines. Journal of Heterocyclic Chemistry, 1997, 34, 1219-1225.	2.6	4
117	Synthesis and biological evaluation of new Donepezil-like Thiaindanones as AChE inhibitors. Journal of Enzyme Inhibition and Medicinal Chemistry, 2008, 23, 696-703.	5.2	4
118	Development of Novel Potential Pleiotropic Compounds of Interest in Alzheimer's Disease Treatment through Rigidification Strategy. Molecules, 2021, 26, 2536.	3.8	4
119	Synthesis and Evaluation of the CNS Activity of New 4-Alkoxyphenylimidazolidin-2-ones Chemical and Pharmaceutical Bulletin, 1998, 46, 711-714.	1.3	3
120	Synthesis and preliminaryin vivoevaluation of new 2-Aryl-6-methyl-1,2-dihydro-1H-pyridin-4-ones and 2-Aryl-6-methylpiperidin-4-ols, as potential anti-amnesiant agents. Journal of Enzyme Inhibition and Medicinal Chemistry, 2005, 20, 551-556.	5.2	3
121	Screening of potential antiviral molecules against equid herpesvirus-1 using cellular impedance measurement: Dataset of 2,891 compounds Data in Brief, 2020, 33, 106492.	1.0	3
122	5-HT4R modulators: a patent landscape. Pharmaceutical Patent Analyst, 2021, 10, 179-181.	1.1	3
123	A Versatile and Efficient Synthesis of 2-Alkyl and 2-Aryl-6-alkyl-2,3-dihydro-1H-pyridin-4-ones. Synthesis, 2002, 2002, 1740-1746.	2.3	2
124	Advances in prodrug design for Alzheimer's disease: the state of the art. Expert Opinion on Drug Discovery, 2022, 17, 325-341.	5.0	2
125	A NEW TYPE OF TRIFLUOROACETYLAMINO ANCHIMERIC ASSISTANCE IN A CYCLOPENTANE RING Heterocyclic Communications, 1996, 2, .	1.2	0
126	Synthesis and Pharmacological Evaluation of New 3-Aryl-3-hydroxyaminopropionic Acids. Pharmacy and Pharmacology Communications, 1999, 5, 239-242.	0.3	0

#	Article	IF	CITATIONS
127	Pharmacological Evaluation of New Baclofen Derivatives. Pharmacy and Pharmacology Communications, 1999, 5, 243-247.	0.3	0
128	A Convenient Synthesis of Dihydro- and Tetrahydro-1,3-thiazine Derivatives from β-Aryl-β-amino Acids ChemInform, 2004, 35, no.	0.0	0
129	First Synthesis of Methyl 3-Amino-4-(het)aryl-1H-pyrrole-2-carboxylates as Useful Scaffolds in Medicinal Chemistry ChemInform, 2004, 35, no.	0.0	0
130	An Efficient Route to 6-(Het)aryl-2-methyl-2,3-dihydro-1H-pyridin-4-ones as Potential nAChRs Ligands ChemInform, 2004, 35, no.	0.0	0
131	First Synthesis of Arylpyrrolo- and Pyrazolopyrrolizinones as Useful Agents with Potential Biological Interest ChemInform, 2004, 35, no.	0.0	0
132	Synthesis of new bisaryl cyclopentathiophene and thieno-cyclopentoxazolidine derivatives as potential cytotoxic agents. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 632-637.	5.2	0
133	P4-206: A NOVEL MTDL APPROACH TO ALZHEIMER DISEASE: 5-HT4 RECEPTOR AGONISTS WITH ACETYLCHOLINESTERASE INHIBITORY ACTIVITIES. , 2014, 10, P863-P864.		0
134	Edema Factor Of Bacillus Anthracis Interacting with its Inhibitors. Biophysical Journal, 2019, 116, 482a-483a.	0.5	0
135	Aminothiaindanone as an Accessible Scaffold for a Three-Point Chemical Diversity. Synthesis, 0, 53, .	2.3	0