Daniel Jun

List of Publications by Year in descending order

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278 papers 6,117 citations

71102 41 h-index 61 g-index

284 all docs

284 docs citations

times ranked

284

4697 citing authors

#	Article	IF	CITATIONS
1	Condensed and Hydrolysable Tannins as Antioxidants Influencing the Health. Mini-Reviews in Medicinal Chemistry, 2008, 8, 436-447.	2.4	218
2	Structural Requirements of Acetylcholinesterase Reactivators. Mini-Reviews in Medicinal Chemistry, 2006, 6, 269-277.	2.4	199
3	Treatment of Organophosphate Intoxication Using Cholinesterase Reactivators: Facts and Fiction. Mini-Reviews in Medicinal Chemistry, 2007, 7, 461-466.	2.4	126
4	In vitro and in vivo evaluation of pyridinium oximes: Mode of interaction with acetylcholinesterase, effect on tabun- and soman-poisoned mice and their cytotoxicity. Toxicology, 2006, 219, 85-96.	4.2	124
5	Phosphatidylinositol 3-Kinase (PI3K) and Phosphatidylinositol 3-Kinase-Related Kinase (PIKK) Inhibitors: Importance of the Morpholine Ring. Journal of Medicinal Chemistry, 2015, 58, 41-71.	6.4	122
6	Tacrine–Trolox Hybrids: A Novel Class of Centrally Active, Nonhepatotoxic Multi-Target-Directed Ligands Exerting Anticholinesterase and Antioxidant Activities with Low In Vivo Toxicity. Journal of Medicinal Chemistry, 2015, 58, 8985-9003.	6.4	121
7	Acetylcholinesterases – the structural similarities and differences. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 417-424.	5.2	110
8	Design of a Potent Reactivator of Tabun-Inhibited AcetylcholinesteraseSynthesis and Evaluation of $(\langle i \rangle E < i \rangle) - 1 - (4 - Carbamoylpyridinium) - 4 - (4 - hydroxyiminomethylpyridinium) - but-2 - ene Dibromide (K203). Journal of Medicinal Chemistry, 2007, 50, 5514 - 5518.$	6.4	100
9	A Resurrection of 7-MEOTA: A Comparison with Tacrine. Current Alzheimer Research, 2013, 10, 893-906.	1.4	92
10	Possible Role of Hydroxylated Metabolites of Tacrine in Drug Toxicity and Therapy of Alzheimers Disease. Current Drug Metabolism, 2008, 9, 332-335.	1.2	83
11	The Antioxidant Additive Approach for Alzheimer's Disease Therapy: New Ferulic (Lipoic) Acid Plus Melatonin Modified Tacrines as Cholinesterases Inhibitors, Direct Antioxidants, and Nuclear Factor (Erythroid-Derived 2)-Like 2 Activators. Journal of Medicinal Chemistry, 2016, 59, 9967-9973.	6.4	83
12	Design and synthesis of new bis-pyridinium oxime reactivators for acetylcholinesterase inhibited by organophosphorous nerve agents. Bioorganic and Medicinal Chemistry Letters, 2005, 15, 2914-2917.	2.2	82
13	Metabolic Pathways of T-2 Toxin. Current Drug Metabolism, 2008, 9, 77-82.	1.2	81
14	Progress in Synthesis of New Acetylcholinesterase Reactivators During the Period 1990-2004. Current Organic Chemistry, 2007, 11, 229-238.	1.6	78
15	SAR study to find optimal cholinesterase reactivator against organophosphorous nerve agents and pesticides. Archives of Toxicology, 2016, 90, 2831-2859.	4.2	75
16	Novel tacrine-tryptophan hybrids: Multi-target directed ligands as potential treatment for Alzheimer's disease. European Journal of Medicinal Chemistry, 2019, 168, 491-514.	5.5	75
17	Acetylcholinesterase and Butyrylcholinesterase – Important Enzymes of Human Body. Acta Medica (Hradec Kralove), 2004, 47, 215-228.	0.5	74
18	Synthesis of the novel series of bispyridinium compounds bearing (E)-but-2-ene linker and evaluation of their reactivation activity against chlorpyrifos-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 622-627.	2.2	65

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19	Chemical Aspects of Pharmacological Prophylaxis Against Nerve Agent Poisoning. Current Medicinal Chemistry, 2009, 16, 2977-2986.	2.4	65
20	Targeting copper(II)-induced oxidative stress and the acetylcholinesterase system in Alzheimer's disease using multifunctional tacrine-coumarin hybrid molecules. Journal of Inorganic Biochemistry, 2016, 161, 52-62.	3.5	63
21	Synthesis of monooxime-monocarbamoyl bispyridinium compounds bearing (<i>E</i>)-but-2-ene linker and evaluation of their reactivation activity against tabun- and paraoxon-inhibited acetylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2008, 23, 70-76.	5.2	61
22	Currently Used Cholinesterase Reactivators Against Nerve Agent Intoxication: Comparison of Their Effectivity in Vitro. Drug and Chemical Toxicology, 2007, 30, 31-40.	2.3	56
23	Oxidative stress in organophosphate poisoning: role of standard antidotal therapy. Journal of Applied Toxicology, 2018, 38, 1058-1070.	2.8	56
24	Improvement of acetylcholinesterase-based assay for organophosphates in way of identification by reactivators. Talanta, 2008, 77, 451-454.	5.5	55
25	Isoquinoline Alkaloids from <i>Berberis vulgaris</i> as Potential Lead Compounds for the Treatment of Alzheimer's Disease. Journal of Natural Products, 2019, 82, 239-248.	3.0	55
26	Effective bisquaternary reactivators of tabun-inhibited AChE. Journal of Applied Toxicology, 2005, 25, 491-495.	2.8	54
27	Mycotoxin Assays Using Biosensor Technology: A Review. Drug and Chemical Toxicology, 2007, 30, 253-261.	2.3	53
28	Pseudo-catalytic scavenging: Searching for a suitable reactivator of phosphorylated butyrylcholinesterase. Chemico-Biological Interactions, 2010, 187, 167-171.	4.0	53
29	A newly developed oxime K203 is the most effective reactivator of tabun-inhibited acetylcholinesterase. BMC Pharmacology & Samp; Toxicology, 2018, 19, 8.	2.4	53
30	Synthesis of a novel series of bispyridinium compounds bearing a xylene linker and evaluation of their reactivation activity against chlorpyrifos-inhibited acetylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2005, 20, 409-415.	5.2	52
31	Monooxime reactivators of acetylcholinesterase with (E)-but-2-ene linkerâ€"Preparation and reactivation of tabun- and paraoxon-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry, 2007, 15, 6733-6741.	3.0	52
32	Pretreatment with pyridinium oximes improves antidotal therapy against tabun poisoning. Toxicology, 2006, 228, 41-50.	4.2	51
33	HPLC Analysis of Hlâ€6 Dichloride and Dimethanesulfonateâ€"Antidotes against Nerve Agents and Organophosphorus Pesticides. Analytical Letters, 2007, 40, 2783-2787.	1.8	51
34	Organophosphate hydrolases as catalytic bioscavengers of organophosphorus nerve agents. Toxicology Letters, 2011, 206, 14-23.	0.8	49
35	Prolyl oligopeptidase and its role in the organism: attention to the most promising and clinically relevant inhibitors. Future Medicinal Chemistry, 2017, 9, 1015-1038.	2.3	48
36	Amperometric Biosensors for Real Time Assays of Organophosphates. Sensors, 2008, 8, 5303-5312.	3.8	47

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37	Synthesis of a novel series of non-symmetrical bispyridinium compounds bearing a xylene linker and evaluation of their reactivation activity against tabun and paraoxon-inhibited acetylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 425-432.	5.2	45
38	Profiling donepezil template into multipotent hybrids with antioxidant properties. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 583-606.	5.2	44
39	From Pyridinium-based to Centrally Active Acetylcholinesterase Reactivators. Mini-Reviews in Medicinal Chemistry, 2014, 14, 215-221.	2.4	44
40	The Acute Toxicity of Acetylcholinesterase Reactivators in Mice in Relation to Their Structure. Neurotoxicity Research, 2006, 9, 291-296.	2.7	43
41	Influence of the Acetylcholinesterase Active Site Protonation on Omega Loop and Active Site Dynamics. Journal of Biomolecular Structure and Dynamics, 2010, 28, 393-403.	3.5	43
42	Amaryllidaceae alkaloids from Narcissus pseudonarcissus L. cv. Dutch Master as potential drugs in treatment of Alzheimer's disease. Phytochemistry, 2019, 165, 112055.	2.9	43
43	Novel series of bispyridinium compounds bearing a (Z)-but-2-ene linkerâ€"Synthesis and evaluation of their reactivation activity against tabun and paraoxon-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry Letters, 2007, 17, 3172-3176.	2.2	40
44	Oxime reactivation of acetylcholinesterase inhibited by toxic phosphorus esters: in vitro kinetics and thermodynamics. Journal of Applied Biomedicine, 2005, 3, 91-99.	1.7	40
45	New methods in synthesis of acetylcholinesterase reactivators and evaluation of their potency to reactivate cyclosarin-inhibited AChE. Chemical Papers, 2006, 60, .	2.2	39
46	<i>In vitro</i> oxime-assisted reactivation of paraoxon-inhibited human acetylcholinesterase and butyrylcholinesterase. Clinical Toxicology, 2009, 47, 545-550.	1.9	38
47	Changes of acetylcholinesterase activity in different rat brain areas following intoxication with nerve agents: Biochemical and histochemical study. Chemico-Biological Interactions, 2007, 165, 14-21.	4.0	37
48	Russian VX: Inhibition and Reactivation of Acetylcholinesterase Compared with VX Agent. Basic and Clinical Pharmacology and Toxicology, 2006, 98, 389-394.	2.5	36
49	Synthesis of asymmetrical bispyridinium compounds bearing cyano-moiety and evaluation of their reactivation activity against tabun and paraoxon-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 5673-5676.	2.2	35
50	7-Methoxytacrine-p-Anisidine Hybrids as Novel Dual Binding Site Acetylcholinesterase Inhibitors for Alzheimer's Disease Treatment. Molecules, 2015, 20, 22084-22101.	3.8	35
51	Design, Synthesis and in vitro Evaluation of Indolotacrine Analogues as Multitargetâ€Directed Ligands for the Treatment of Alzheimer's Disease. ChemMedChem, 2016, 11, 1264-1269.	3.2	35
52	Alkaloids from Narcissus poeticus cv. Pink Parasol of various structural types and their biological activity. Archives of Pharmacal Research, 2018, 41, 208-218.	6.3	35
53	Novel Tacrine-Scutellarin Hybrids as Multipotent Anti-Alzheimer's Agents: Design, Synthesis and Biological Evaluation. Molecules, 2017, 22, 1006.	3.8	32
54	New quaternary pyridine aldoximes as casual antidotes against nerve agents intoxications. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2005, 149, 75-82.	0.6	32

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55	Cholinesterase Reactivators: The Fate and Effects in the Organism Poisoned with Organophosphates/Nerve Agents. Current Drug Metabolism, 2007, 8, 803-809.	1.2	31
56	Potency of several oximes to reactivate human acetylcholinesterase and butyrylcholinesterase inhibited by paraoxon in vitro. Chemico-Biological Interactions, 2008, 175, 421-424.	4.0	31
57	The summary on non-reactivation cholinergic properties of oxime reactivators: the interaction with muscarinic and nicotinic receptors. Archives of Toxicology, 2013, 87, 711-719.	4.2	31
58	TLC Analysis of Intermediates Arising During the Preparation of Oxime HI-6 Dimethanesulfonate. Journal of Chromatographic Science, 2008, 46, 316-319.	1.4	30
59	Cholinesterase Biosensor Construction – A Review. Protein and Peptide Letters, 2008, 15, 795-798.	0.9	30
60	Isoquinoline Alkaloids from <i>Fumaria officinalis</i> L. and Their Biological Activities Related to <i>Alzheimer</i> 's Disease. Chemistry and Biodiversity, 2016, 13, 91-99.	2.1	30
61	Synthesis of Bispyridinium Compounds Bearing Propane Linker and Evaluation of their Reactivation Activity against Tabun- and Paraoxon-Inhibited Acetylcholinesterase. Letters in Organic Chemistry, 2006, 3, 831-835.	0.5	30
62	The influence of simvastatin, atorvastatin and high-cholesterol diet on acetylcholinesterase activity, amyloid beta and cholesterol synthesis in rat brain. Steroids, 2009, 74, 13-19.	1.8	29
63	Acetylcholinesterase and butyrylcholinesterase-important enzymes of human body. Acta Medica (Hradec Kralove), 2004, 47, 215-28.	0.5	29
64	<i>In Vitro</i> Antiplatelet Activity of Flavonoids from <i>Leuzea Carthamoides</i> Drug and Chemical Toxicology, 2008, 31, 27-35.	2.3	28
65	Design, synthesis and in vitro testing of 7-methoxytacrine-amantadine analogues: a novel cholinesterase inhibitors for the treatment of Alzheimer's disease. Medicinal Chemistry Research, 2015, 24, 2645-2655.	2.4	28
66	Progress in acetylcholinesterase reactivators and in the treatment of organophosphorus intoxication: a patent review (2006–2016). Expert Opinion on Therapeutic Patents, 2017, 27, 971-985.	5.0	28
67	Prophylaxis Against Nerve Agent Intoxications. Defence Science Journal, 2006, 56, 775-784.	0.8	28
68	Could oxime HI-6 really be considered as "broad-spectrum" antidote?. Journal of Applied Biomedicine, 2009, 7, 143-149.	1.7	28
69	Potency of new structurally different oximes to reactivate cyclosarin-inhibited human brain acetylcholinesterases. Journal of Enzyme Inhibition and Medicinal Chemistry, 2006, 21, 663-666.	5.2	26
70	Reactivation of DFP- and paraoxon-inhibited acetylcholinesterases by pyridinium oximes. Chemico-Biological Interactions, 2008, 175, 365-367.	4.0	26
71	Oxime K027: novel low-toxic candidate for the universal reactivator of nerve agent- and pesticide-inhibited acetylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 509-512.	5.2	26
72	Development of 2-Methoxyhuprine as Novel Lead for Alzheimer's Disease Therapy. Molecules, 2017, 22, 1265.	3.8	26

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73	Monoquaternary pyridinium salts with modified side chain—synthesis and evaluation on model of tabun- and paraoxon-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry, 2008, 16, 8218-8223.	3.0	25
74	In Vitro Ability of Currently Available Oximes to Reactivate Organophosphate Pesticide-Inhibited Human Acetylcholinesterase and Butyrylcholinesterase. International Journal of Molecular Sciences, 2011, 12, 2077-2087.	4.1	25
75	Tannins and their Influence on Health. , 2014, , 159-208.		25
76	Reactivation of organophosphate-inhibited acetylcholinesterase by quaternary pyridinium aldoximes. Neurotoxicity Research, 2004, 6, 565-570.	2.7	24
77	Bis-pyridiumaldoxime reactivators connected with CH2O(CH2)nOCH2 linkers between pyridinium rings and their reactivity against VX. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 4852-4855.	2.2	24
78	An attempt to assess functionally minimal acetylcholinesterase activity necessary for survival of rats intoxicated with nerve agents. Chemico-Biological Interactions, 2008, 175, 281-285.	4.0	24
79	Discovery of novel berberine derivatives with balanced cholinesterase and prolyl oligopeptidase inhibition profile. European Journal of Medicinal Chemistry, 2020, 203, 112593.	5.5	24
80	Prophylaxis and Post-exposure Treatment of Intoxications Caused by Nerve Agents and Organophosphorus Pesticides. Mini-Reviews in Medicinal Chemistry, 2013, 13, 2102-2115.	2.4	24
81	The Reactivating and Therapeutic Efficacy of Oximes to Counteract Russian VX Poisonings. International Journal of Toxicology, 2006, 25, 397-401.	1.2	23
82	Amperometric Biosensor for Evaluation of Competitive Cholinesterase Inhibition by the Reactivator $Hla \in 6$. Analytical Letters, 2007, 40, 2351-2359.	1.8	23
83	A comparison of reactivating efficacy of newly developed oximes (K074, K075) and currently available oximes (obidoxime, HI-6) in soman, cyclosarin and tabun-poisoned rats. Chemico-Biological Interactions, 2008, 175, 425-427.	4.0	23
84	Isoquinoline alkaloids as prolyl oligopeptidase inhibitors. Fìtoterapìâ, 2015, 103, 192-196.	2.2	23
85	Isolation of Amaryllidaceae alkaloids from Nerine bowdenii W. Watson and their biological activities. RSC Advances, 2016, 6, 80114-80120.	3.6	23
86	Exploring Structure-Activity Relationship in Tacrine-Squaramide Derivatives as Potent Cholinesterase Inhibitors. Biomolecules, 2019, 9, 379.	4.0	23
87	In Vitro and In Silico Acetylcholinesterase Inhibitory Activity of Thalictricavine and Canadine and Their Predicted Penetration across the Blood-Brain Barrier. Molecules, 2019, 24, 1340.	3.8	23
88	Inhibition of Acetylcholinesterase in Different Structures of the Rat Brain Following Soman Intoxication Pretreated with Huperzine A. International Journal of Molecular Sciences, 2007, 8, 1165-1176.	4.1	22
89	Synthesis and In Vitro Evaluation of N-(Bromobut-3-en-2-yl)-7-methoxy-1,2,3,4-tetrahydroacridin-9-amine as a Cholinesterase Inhibitor with Regard to Alzheimer's Disease Treatment. Molecules, 2010, 15, 8804-8812.	3.8	22
90	A 7-methoxytacrine–4-pyridinealdoxime hybrid as a novel prophylactic agent with reactivation properties in organophosphate intoxication. Toxicology Research, 2016, 5, 1012-1016.	2.1	22

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91	Cytotoxicity of acetylcholinesterase reactivators evaluated <i>in vitro</i> and its relation to their structure. Drug and Chemical Toxicology, 2019, 42, 252-256.	2.3	22
92	The wide-spectrum antimicrobial effect of novel N-alkyl monoquaternary ammonium salts and their mixtures; the QSAR study against bacteria. European Journal of Medicinal Chemistry, 2020, 206, 112584.	5 . 5	22
93	In vitro reactivation potency of acetylcholinesterase reactivators â€" K074 and K075 â€" to reactivate tabun-inhibited human brain cholinesterases. Neurotoxicity Research, 2007, 11, 101-106.	2.7	21
94	Evaluation of natural antioxidants of <i>Leuzea carthamoides</i> as a result of a screening study of 88 plant extracts from the European Asteraceae and Cichoriaceae. Journal of Enzyme Inhibition and Medicinal Chemistry, 2008, 23, 218-224.	5.2	21
95	Evaluation of natural substances from <i>Evolvulus alsinoides</i> L. with the purpose of determining their antioxidant potency. Journal of Enzyme Inhibition and Medicinal Chemistry, 2008, 23, 574-578.	5.2	21
96	Synthesis, in vitro acetylcholinesterase inhibitory activity and molecular docking of new acridine-coumarin hybrids. International Journal of Biological Macromolecules, 2017, 104, 333-338.	7.5	21
97	(±)- BIGI-3h : Pentatarget-Directed Ligand combining Cholinesterase, Monoamine Oxidase, and Glycogen Synthase Kinase 3β Inhibition with Calcium Channel Antagonism and Antiaggregating Properties for Alzheimer's Disease. ACS Chemical Neuroscience, 2021, 12, 1328-1342.	3.5	21
98	Acetylcholinesterase Inhibitors and Drugs Acting on Muscarinic Receptors- Potential Crosstalk of Cholinergic Mechanisms During Pharmacological Treatment. Current Neuropharmacology, 2017, 15, 637-653.	2.9	21
99	Synthesis, Structure, and Cholinergic Effect of Novel Neuroprotective Compounds Bearing the Tacrine Pharmacophore. Heterocycles, 2008, 76, 1219.	0.7	20
100	Activity of cholinesterases in a young and healthy middle-European population: Relevance for toxicology, pharmacology and clinical praxis. Toxicology Letters, 2017, 277, 24-31.	0.8	20
101	Alkaloids of Zephyranthes citrina (Amaryllidaceae) and their implication to Alzheimer's disease: Isolation, structural elucidation and biological activity. Bioorganic Chemistry, 2021, 107, 104567.	4.1	20
102	Alkaloids from Chlidanthus fragrans and their acetylcholinesterase, butyrylcholinesterase and prolyl oligopeptidase activities. Natural Product Communications, 2013, 8, 1541-4.	0.5	20
103	Preparation of Benzalkonium Salts Differing in the Length of a Side Alkyl Chain. Molecules, 2007, 12, 2341-2347.	3.8	19
104	RP-HPLC determination of the lipophilicity of bispyridinium reactivators of acetylcholinesterase bearing a but-2-ene connecting linker. Analytical and Bioanalytical Chemistry, 2008, 391, 367-372.	3.7	19
105	The Influence of Acetylcholinesterase Reactivators on Selected Hepatic Functions in Rats. Basic and Clinical Pharmacology and Toxicology, 2008, 103, 119-123.	2.5	19
106	Preparation and characterization of methoxy polyethylene glycol-conjugated phosphotriesterase as a potential catalytic bioscavenger against organophosphate poisoning. Chemico-Biological Interactions, 2010, 187, 380-383.	4.0	19
107	Cholinesterase and Prolyl Oligopeptidase Inhibitory Activities of Alkaloids from Argemone platyceras (Papaveraceae). Molecules, 2017, 22, 1181.	3.8	19
108	Oxime K203: a drug candidate for the treatment of tabun intoxication. Archives of Toxicology, 2019, 93, 673-691.	4.2	19

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109	Synthesis and biological assessment of KojoTacrines as new agents for Alzheimer's disease therapy. Journal of Enzyme Inhibition and Medicinal Chemistry, 2019, 34, 163-170.	5.2	19
110	In vitro and in silico Evaluation of Non-Quaternary Reactivators of AChE as Antidotes of Organophosphorus Poisoning - a New Hope or a Blind Alley?. Medicinal Chemistry, 2018, 14, 281-292.	1.5	19
111	The development of ataxia telangiectasia mutated kinase inhibitors. Mini-Reviews in Medicinal Chemistry, 2014, 14, 1-1.	2.4	18
112	Novel Bisquaternary Oximesâ€"Reactivation of Acetylcholinesterase and Butyrylcholinesterase Inhibited by Paraoxon. Molecules, 2009, 14, 4915-4921.	3.8	17
113	The effect of oxime reactivators on muscarinic receptors: Functional and binding examinations. Environmental Toxicology and Pharmacology, 2011, 31, 364-370.	4.0	17
114	Cholinergic properties ofÂnew 7-methoxytacrine-donepezil derivatives. General Physiology and Biophysics, 2015, 34, 189-200.	0.9	17
115	Tacrine – Benzothiazoles: Novel class of potential multitarget anti-Alzheimeŕs drugs dealing with cholinergic, amyloid and mitochondrial systems. Bioorganic Chemistry, 2021, 107, 104596.	4.1	17
116	Application of Artificial Neural Networks in Condition Based Predictive Maintenance. Studies in Computational Intelligence, 2016, , 75-86.	0.9	17
117	A Comparison of the Potency of the Oxime HLö-7 and Currently Used Oximes (HI-6, Pralidoxime,) Tj ETQq1 1 (Acta Medica (Hradec Kralove), 2005, 48, 81-86.	0.784314 r _. 0.5	gBT Overlock 17
118	In vitro reactivation potency of some acetylcholinesterase reactivators against sarin- and cyclosarin-induced inhibitions. Journal of Applied Toxicology, 2005, 25, 296-300.	2.8	16
119	Time-Course Changes of Acetylcholinesterase Activity in Blood and Some Tissues in Rats After Intoxication by Russian VX. Neurotoxicity Research, 2009, 16, 356-360.	2.7	16
120	Reactivation of Human Acetylcholinesterase and Butyrylcholinesterase Inhibited by Leptophos-Oxon with Different Oxime Reactivators in Vitro. International Journal of Molecular Sciences, 2010, 11, 2856-2863.	4.1	16
121	Molecular modeling studies on the interactions of aflatoxin B1 and its metabolites with the peripheral anionic site of human acetylcholinesterase. Journal of Biomolecular Structure and Dynamics, 2019, 37, 2041-2048.	3.5	16
122	Aflatoxin Assay Using an Amperometric Sensor Strip and Acetylcholinesterase as Recognition Element. Sensor Letters, 2008, 6, 450-453.	0.4	16
123	Targeted Synthesis of 1-(4-Hydroxyiminomethylpyridinium)-3-pyridiniumpropane Dibromide – A New Nerve Agent Reactivator. Molecules, 2007, 12, 1964-1972.	3.8	15
124	Photometric microplate assay for estimation of the efficacy of paraoxon-inhibited acetylcholinesterase reactivation. Journal of Enzyme Inhibition and Medicinal Chemistry, 2008, 23, 781-784.	5.2	15
125	Synthesis, antimicrobial evaluation and molecular modeling of 5-hydroxyisoquinolinium salt series; the effect of the hydroxyl moiety. Bioorganic and Medicinal Chemistry, 2016, 24, 841-848.	3.0	15
126	Synthesis, <i>inÂvitro</i> screening and molecular docking of isoquinolinium-5-carbaldoximes as acetylcholinesterase and butyrylcholinesterase reactivators. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 478-488.	5.2	15

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127	In vitro reactivation potency of bispyridinium (E)-but-2-ene linked acetylcholinesterase reactivators against tabun-inhibited acetylcholinesterase. Journal of Applied Biomedicine, 2007, 5, 25-30.	1.7	15
128	Is It the Twilight of BACE1 Inhibitors?. Current Neuropharmacology, 2020, 19, 61-77.	2.9	15
129	Revised NMR data for 9-O-demethylgalanthine: an alkaloid from Zephyranthes robusta (Amaryllidaceae) and its biological activity. Natural Product Communications, 2014, 9, 787-8.	0.5	15
130	Inhibition of blood cholinesterases following intoxication with VX and its derivatives. Journal of Applied Toxicology, 2007, 27, 458-463.	2.8	14
131	Fluctuation in the ergosterol and deoxynivalenol content in barley and malt during malting process. Analytical and Bioanalytical Chemistry, 2010, 397, 109-114.	3.7	14
132	Why acetylcholinesterase reactivators do not work in butyrylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 318-322.	5. 2	14
133	Alkaloids from Chlidanthus fragrans and their Acetylcholinesterase, Butyrylcholinesterase and Prolyl Oligopeptidase Activities. Natural Product Communications, 2013, 8, 1934578X1300801.	0.5	14
134	Benzothiazolyl Ureas are Low Micromolar and Uncompetitive Inhibitors of 17β-HSD10 with Implications to Alzheimer's Disease Treatment. International Journal of Molecular Sciences, 2020, 21, 2059.	4.1	14
135	Strategy for the development of new acetylcholinesterase reactivators - antidotes used for treatment of nerve agent poisonings. Biomedical Papers of the Medical Faculty of the University Palacký, Olomouc, Czechoslovakia, 2005, 149, 429-431.	0.6	14
136	Twelve Different HI-6 Salts and their Potency to Reactivate Cyclosarin Inhibited AChE In Vitro. Letters in Drug Design and Discovery, 2007, 4, 510-512.	0.7	13
137	Evaluation of the antioxidant activity of several naturally occurring coumarins and their synthesized analogues by "ferric reducing antioxidant power―assay. Journal of Enzyme Inhibition and Medicinal Chemistry, 2014, 29, 49-54.	5.2	13
138	Novel Group of AChE Reactivatorsâ€"Synthesis, In Vitro Reactivation and Molecular Docking Study. Molecules, 2018, 23, 2291.	3.8	13
139	Prediction of a new broad-spectrum reactivator capable of reactivating acetylcholinesterase inhibited by nerve agents. Journal of Applied Biomedicine, 2005, 3, 139-145.	1.7	13
140	Comparison ofin vitro potency of oximes (pralidoxime, obidoxime, HI-6) to reactivate sarin-inhibited acetylcholinesterase in various parts of pig brain. Journal of Applied Toxicology, 2005, 25, 271-276.	2.8	12
141	In VitroEvaluation of Acetylcholinesterase Reactivators as Potential Antidotes Against Tabun Nerve Agent Poisonings. Drug and Chemical Toxicology, 2006, 29, 443-449.	2.3	12
142	A comparison of reactivating efficacy of newly developed oximes (K074, K075) and currently available oximes (obidoxime, HI-6) in cyclosarin-and tabun-poisoned rats. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 297-300.	5.2	12
143	Changes of Cholinesterase Activities in the Rat Blood and Brain After Sarin Intoxication Pretreated with Butyrylcholinesterase. Drug and Chemical Toxicology, 2007, 30, 351-359.	2.3	12
144	Bisquaternary Oximes as Reactivators of Tabun-Inhibited Human Brain Cholinesterases: An in vitro Study. Basic and Clinical Pharmacology and Toxicology, 2007, 101, 25-28.	2.5	12

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