List of Publications by Year in descending order

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DANIEL LUN

#	Article	IF	CITATIONS
1	Monoterpene indole alkaloids from Vinca minor L. (Apocynaceae): Identification of new structural scaffold for treatment of Alzheimer's disease. Phytochemistry, 2022, 194, 113017.	2.9	7
2	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
3	Development of versatile and potent monoquaternary reactivators of acetylcholinesterase. Archives of Toxicology, 2021, 95, 985-1001.	4.2	7
4	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
5	(±)- 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
6	Synthesis of New Biscoumarin Derivatives, In Vitro Cholinesterase Inhibition, Molecular Modelling and Antiproliferative Effect in A549 Human Lung Carcinoma Cells. International Journal of Molecular Sciences, 2021, 22, 3830.	4.1	3
7	Huprine Y – Tryptophan heterodimers with potential implication to Alzheimer's disease treatment. Bioorganic and Medicinal Chemistry Letters, 2021, 43, 128100.	2.2	5
8	Amaryllidaceae Alkaloids of Norbelladine-Type as Inspiration for Development of Highly Selective Butyrylcholinesterase Inhibitors: Synthesis, Biological Activity Evaluation, and Docking Studies. International Journal of Molecular Sciences, 2021, 22, 8308.	4.1	5
9	Synthesis and Decontamination Effect on Chemical and Biological Agents of Benzoxonium-Like Salts. Toxics, 2021, 9, 222.	3.7	2
10	Pursuing the Complexity of Alzheimer's Disease: Discovery of Fluoren-9-Amines as Selective Butyrylcholinesterase Inhibitors and N-Methyl-d-Aspartate Receptor Antagonists. Biomolecules, 2021, 11, 3.	4.0	4
11	Oxime K074 – <i>in vitro</i> and <i>in silico</i> reactivation of acetylcholinesterase inhibited by nerve agents and pesticides. Toxin Reviews, 2020, 39, 157-166.	3.4	5
12	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
13	Encapsulation of oxime K027 into cucurbit[7]uril: In vivo evaluation of safety, absorption, brain distribution and reactivation effectiveness. Toxicology Letters, 2020, 320, 64-72.	0.8	10
14	Simple validated method of LC–MS/MS determination of BZ agent in rat plasma samples. Drug Testing and Analysis, 2020, 12, 431-438.	2.6	4
15	Cysteine-Targeted Insecticides against A. gambiae Acetylcholinesterase Are Neither Selective nor Reversible Inhibitors. ACS Medicinal Chemistry Letters, 2020, 11, 65-71.	2.8	11
16	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
17	Discovery of novel berberine derivatives with balanced cholinesterase and prolyl oligopeptidase inhibition profile. European Journal of Medicinal Chemistry, 2020, 203, 112593.	5.5	24
18	Functionalized aromatic esters of the Amaryllidaceae alkaloid haemanthamine and their in vitro and in silico biological activity connected to Alzheimer's disease. Bioorganic Chemistry, 2020, 100, 103928.	4.1	9

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19	Monitoring of blood cholinesterase activity in workers exposed to nerve agents. , 2020, , 1035-1045.		0
20	Other toxic chemicals as potential chemical warfare agents. , 2020, , 403-412.		1
21	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
22	Global impact of chemical warfare agents used before and after 1945. , 2020, , 27-36.		1
23	Pharmacological prophylaxis against nerve agent poisoning: experimental studies and practical implications. , 2020, , 1091-1101.		0
24	Donepezil and Rivastigmine: Pharmacokinetic Profile and Brain-targeting After Intramuscular Administration in Rats. Iranian Journal of Pharmaceutical Research, 2020, 19, 95-102.	0.5	4
25	DEFINITION OF THE TARGET POPULATION FOR EXTERNAL PACEMAKER AS A KEY ASPECT IN SUCCESSFUL MEDICAL DEVICE HTA PROCESS. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2020, 89, 99-107.	0.5	0
26	Is It the Twilight of BACE1 Inhibitors?. Current Neuropharmacology, 2020, 19, 61-77.	2.9	15
27	OPCW BIOMEDICAL PROFICIENCY TEST IN THE LABORATORY OF ANALYTICAL CHEMISTRY AT THE DEPARTMENT OFÂTOXICOLOGY AND MILITARY PHARMACY. Military Medical Science Letters (Vojenske) Tj ETQ	q1 10 057 84	∔31 ⊕ rgBT /Ο∖
28	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
29	Exploring Structure-Activity Relationship in Tacrine-Squaramide Derivatives as Potent Cholinesterase Inhibitors. Biomolecules, 2019, 9, 379.	4.0	23
30	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
31	Pharmacological and toxicological in vitro and in vivo effect of higher doses of oxime reactivators. Toxicology and Applied Pharmacology, 2019, 383, 114776.	2.8	5
32	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
33	Surface screening, molecular modeling and in vitro studies on the interactions of aflatoxin M1 and human enzymes acetyl- and butyrylcholinesterase. Chemico-Biological Interactions, 2019, 308, 113-119.	4.0	4
34	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
35	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
36	Inhalation of molecular hydrogen prevents ischemia-reperfusion liver damage during major liver resection. Annals of Translational Medicine, 2019, 7, 774-774.	1.7	11

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37	Oxime K203: a drug candidate for the treatment of tabun intoxication. Archives of Toxicology, 2019, 93, 673-691.	4.2	19
38	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
39	Novel quinazolin-4-one derivatives as potentiating agents of doxorubicin cytotoxicity. Bioorganic Chemistry, 2019, 82, 204-210.	4.1	2
40	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
41	N-alkylated Tacrine Derivatives as Potential Agents in Alzheimer's Disease Therapy. Current Alzheimer Research, 2019, 16, 333-343.	1.4	5
42	Oxidative stress in organophosphate poisoning: role of standard antidotal therapy. Journal of Applied Toxicology, 2018, 38, 1058-1070.	2.8	56
43	Synthesis, Biological Assessment and Molecular Modeling of Racemic <i>QuinoPyranoTacrines</i> for Alzheimer's Disease Therapy. ChemistrySelect, 2018, 3, 461-466.	1.5	10
44	Profiling donepezil template into multipotent hybrids with antioxidant properties. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 583-606.	5.2	44
45	The influence of modulators of acetylcholinesterase on the resistance of mice against soman and on the effectiveness of antidotal treatment of soman poisoning in mice. Journal of Applied Biomedicine, 2018, 16, 10-14.	1.7	4
46	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
47	Purin-6-one and pyrrolo[2,3-d]pyrimidin-4-one derivatives as potentiating agents of doxorubicin cytotoxicity. Future Medicinal Chemistry, 2018, 10, 2029-2038.	2.3	2
48	Molecular Modeling Studies on the Interactions of Aflatoxin B1 and Its Metabolites with Human Acetylcholinesterase. Part II: Interactions with the Catalytic Anionic Site (CAS). Toxins, 2018, 10, 389.	3.4	5
49	Novel Group of AChE Reactivators—Synthesis, In Vitro Reactivation and Molecular Docking Study. Molecules, 2018, 23, 2291.	3.8	13
50	Synthesis, Biological Evaluation, and Docking Studies of Novel Bisquaternary Aldoxime Reactivators on Acetylcholinesterase and Butyrylcholinesterase Inhibited by Paraoxon. Molecules, 2018, 23, 1103.	3.8	11
51	Simultaneous determination of malondialdehyde and 3â€nitrotyrosine in cultured human hepatoma cells by liquid chromatography–mass spectrometry. Biomedical Chromatography, 2018, 32, e4349.	1.7	12
52	A newly developed oxime K203 is the most effective reactivator of tabun-inhibited acetylcholinesterase. BMC Pharmacology & amp; Toxicology, 2018, 19, 8.	2.4	53
53	Development of small bisquaternary cholinesterase inhibitors as drugs for pre-treatment of nerve agent poisonings. Drug Design, Development and Therapy, 2018, Volume 12, 505-512.	4.3	4
54	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

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55	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
56	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
57	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
58	Development of information andÂmanagement system for laboratory based on open source licensed software withÂsecurity logs extension. Journal of Intelligent and Fuzzy Systems, 2017, 32, 1497-1508.	1.4	2
59	Bis-isoquinolinium and bis-pyridinium acetylcholinesterase inhibitors: in vitro screening of probes for novel selective insecticides. RSC Advances, 2017, 7, 39279-39291.	3.6	6
60	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
61	Novel Tacrine-Scutellarin Hybrids as Multipotent Anti-Alzheimer's Agents: Design, Synthesis and Biological Evaluation. Molecules, 2017, 22, 1006.	3.8	32
62	Cholinesterase and Prolyl Oligopeptidase Inhibitory Activities of Alkaloids from Argemone platyceras (Papaveraceae). Molecules, 2017, 22, 1181.	3.8	19
63	Development of 2-Methoxyhuprine as Novel Lead for Alzheimer's Disease Therapy. Molecules, 2017, 22, 1265.	3.8	26
64	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
65	HLö-7 - A REVIEW OF ACETYLCHOLINESTERASE REACTIVATOR AGAINST ORGANOPHOSPHOROUS INTOXICATION. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2017, 86, 70-83.	0.5	2
66	Design, Synthesis and in vitro Evaluation of Indolotacrine Analogues as Multitargetâ€Ðirected Ligands for the Treatment of Alzheimer's Disease. ChemMedChem, 2016, 11, 1264-1269.	3.2	35
67	Device Security Implementation Model based on Internet of Things for a Laboratory Environment. IFAC-PapersOnLine, 2016, 49, 419-424.	0.9	8
68	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
69	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
70	SAR study to find optimal cholinesterase reactivator against organophosphorous nerve agents and pesticides. Archives of Toxicology, 2016, 90, 2831-2859.	4.2	75
71	In vitro characterization of acetylcholinesterase reactivators: The cytotoxicity and oxidative stress induction. Toxicology Letters, 2016, 258, S126-S127.	0.8	0
72	Biological mechanisms of sulfur mustard toxicity: Dose and time-dependent study. Toxicology Letters, 2016, 258, S254-S255.	0.8	0

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73	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
74	Isolation of Amaryllidaceae alkaloids from Nerine bowdenii W. Watson and their biological activities. RSC Advances, 2016, 6, 80114-80120.	3.6	23
75	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
76	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
77	Novel caffeine derivatives with antiproliferative activity. RSC Advances, 2016, 6, 32534-32539.	3.6	12
78	Application of Artificial Neural Networks in Condition Based Predictive Maintenance. Studies in Computational Intelligence, 2016, , 75-86.	0.9	17
79	Small Molecules Targeting Ataxia Telangiectasia and Rad3-Related (ATR) Kinase: An Emerging way to Enhance Existing Cancer Therapy. Current Cancer Drug Targets, 2016, 16, 200-208.	1.6	11
80	TRISOXIME - a Bulky Trisquaternary Reactivator of Acetylcholinesterase. Letters in Drug Design and Discovery, 2016, 13, 372-375.	0.7	1
81	Preparation of 7â€Methoxy Tacrine Dimer Analogs and Their <i>In vitro/In silico</i> Evaluation as Potential Cholinesterase Inhibitors. Bulletin of the Korean Chemical Society, 2015, 36, 1654-1660.	1.9	9
82	Novel Cholinesterase Reactivators. , 2015, , 1071-1087.		4
83	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
84	Cholinergic properties ofÂnew 7-methoxytacrine-donepezil derivatives. General Physiology and Biophysics, 2015, 34, 189-200.	0.9	17
85	Pharmacological Prophylaxis Against Nerve Agent Poisoning. , 2015, , 979-987.		2
86	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
87	The biomedical data collecting system. , 2015, , .		2
88	Global Impact of Chemical Warfare Agents Used Before and After 1945. , 2015, , 17-25.		2
89	Other Toxic Chemicals as Potential Chemical Warfare Agents. , 2015, , 337-345.		3
90	Isoquinoline alkaloids as prolyl oligopeptidase inhibitors. Fìtoterapìâ, 2015, 103, 192-196.	2.2	23

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91	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
92	Development of Information and Management System for Laboratory Based on Open Source Licensed Software. Lecture Notes in Computer Science, 2015, , 377-387.	1.3	6
93	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
94	Universality of Oxime K203 for Reactivation of Nerve Agent-Inhibited AChE. Medicinal Chemistry, 2015, 11, 683-686.	1.5	7
95	Impact of tacrine and 7-methoxytacrine on gastric myoelectrical activity assessed using electrogastrography in experimental pigs. Neuroendocrinology Letters, 2015, 36 Suppl 1, 150-5.	0.2	2
96	Chemical Composition of Bioactive Alkaloid Extracts from Some Narcissus Species and Varieties and their Biological Activity. Natural Product Communications, 2014, 9, 1934578X1400900.	0.5	5
97	Revised NMR Data for 9-O-Demethylgalanthine: An Alkaloid from Zephyranthes robusta (Amaryllidaceae) and its Biological Activity. Natural Product Communications, 2014, 9, 1934578X1400900.	0.5	6
98	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
99	Tannins and their Influence on Health. , 2014, , 159-208.		25
100	Pharmacotherapy of Alzheimer's Disease: Current State and Future Perspectives. , 2014, , 3-39.		5
101	Trithiocyanurate Complexes of Iron, Manganese and Nickel and Their Anticholinesterase Activity. Molecules, 2014, 19, 4338-4354.	3.8	8
102	In vitro evaluation of reactivating efficacy of newly developed oximes for preparation of "pseudocatalytic scavenger―based on butyrylcholinesterase. Toxicology Letters, 2014, 229, S167-S168.	0.8	0
103	The system of instant access to the life biomedical data. , 2014, , .		2
104	From Pyridinium-based to Centrally Active Acetylcholinesterase Reactivators. Mini-Reviews in Medicinal Chemistry, 2014, 14, 215-221.	2.4	44
105	The development of ataxia telangiectasia mutated kinase inhibitors. Mini-Reviews in Medicinal Chemistry, 2014, 14, 1-1.	2.4	18
106	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
107	Chemical composition of bioactive alkaloid extracts from some Narcissus species and varieties and their biological activity. Natural Product Communications, 2014, 9, 1151-5.	0.5	9
108	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

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109	Preparation, in vitro evaluation and molecular modelling of pyridinium–quinolinium/isoquinolinium non-symmetrical bisquaternary cholinesterase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6663-6666.	2.2	11
110	Hyaluronidase: Its effects on HI-6 dichloride and dimethanesulphonate pharmacokinetic profile in pigs. Toxicology Letters, 2013, 220, 167-171.	0.8	6
111	Alkaloids from Chlidanthus fragrans and their Acetylcholinesterase, Butyrylcholinesterase and Prolyl Oligopeptidase Activities. Natural Product Communications, 2013, 8, 1934578X1300801.	0.5	14
112	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
113	A Resurrection of 7-MEOTA: A Comparison with Tacrine. Current Alzheimer Research, 2013, 10, 893-906.	1.4	92
114	SCREENING OF BLOOD-BRAIN BARRIER PENETRATION USING THE IMMOBILIZED ARTIFICIAL MEMBRANE PHOSPHATIDYLCHOLINE COLUMN CHROMATOGRAPHY AT THE PHYSIOLOGICAL PH. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2013, 82, 55-62.	0.5	1
115	SELECTIVE MONITORING OF ENZYMATIC ACTIVITY OF ACETYLCHOLINESTERASE BY FLOW INJECTION ANALYSIS WITH MASS SPECTROMETRIC DETECTION. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2013, 82, 120-125.	0.5	Ο
116	Intravenous application of HI-6 salts (dichloride and dimethansulphonate) in pigs: comparison with pharmacokinetics profile after intramuscular administration. Neuroendocrinology Letters, 2013, 34 Suppl 2, 74-8.	0.2	0
117	Impact of paraoxon followed by acetylcholinesterase reactivator HI-6 on gastric myoelectric activity in experimental pigs. Neuroendocrinology Letters, 2013, 34 Suppl 2, 79-83.	0.2	4
118	Alkaloids from Chlidanthus fragrans and their acetylcholinesterase, butyrylcholinesterase and prolyl oligopeptidase activities. Natural Product Communications, 2013, 8, 1541-4.	0.5	20
119	The interaction of standard oxime reactivators with hemicholinium-3 sensitive choline carriers. Toxicology Letters, 2012, 212, 315-319.	0.8	7
120	The effect of oxime reactivators on muscarinic receptors: Functional and binding examinations. Environmental Toxicology and Pharmacology, 2011, 31, 364-370.	4.0	17
121	Organophosphate hydrolases as catalytic bioscavengers of organophosphorus nerve agents. Toxicology Letters, 2011, 206, 14-23.	0.8	49
122	TLC analysis of twelve different salts of oxime HI-6 — Reactivator of nerve agent inhibited AChE. Journal of Planar Chromatography - Modern TLC, 2011, 24, 105-107.	1.2	0
123	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
124	ON THE UNIVERSALITY OF OXIME HLö-7 - ANTIDOTE FOR CASE OF THE NERVE AGENT POISONING. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2011, 80, 80-84.	0.5	4
125	HUMAN SERUM BUTYRYLCHOLINESTERASE AS A PROPHYLAXIS AGAINST RUSSIAN VX. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2011, 80, 97-102.	0.5	0
126	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

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127	Synthesis of (2E)-2-methyl-3-(4-{[4-(quinolin-2-ylmethoxy)phenyl]sulfanyl}phenyl)prop-2-enoic acid (VUFB 20609) and 2-methyl-3-(4-{[4-(quinolin-2-ylmethoxy)phenyl]sulfanyl}phenyl)propanoic acid (VUFB) Tj	ETQq12.140.78	43014 rgBT /(
128	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
129	Pseudo-catalytic scavenging: Searching for a suitable reactivator of phosphorylated butyrylcholinesterase. Chemico-Biological Interactions, 2010, 187, 167-171.	4.0	53
130	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
131	New Bisquaternary Isoquinolinium Inhibitors of Brain Cholinesterases - Synthesis and Anticholinesterase Activity. Letters in Drug Design and Discovery, 2010, 7, 1-4.	0.7	0
132	High-Performance Liquid Chromatography Analysis of By-Products and Intermediates Arising During the Synthesis of the Acetylcholinesterase Reactivator HI-6. Journal of Chromatographic Science, 2010, 48, 694-696.	1.4	11
133	Characterization of the anticholinergic properties of obidoxime; functional examinations of the rat atria and the urinary bladder. Toxicology Mechanisms and Methods, 2010, 20, 428-433.	2.7	10
134	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
135	Cholinesterase Reactivators as Prophylactics Against Nerve Agents. Current Bioactive Compounds, 2010, 6, 2-8.	0.5	10
136	The effect of trimedoxime on acetylcholinesterase and on the cholinergic system of the rat bladder. Journal of Applied Biomedicine, 2010, 8, 87-92.	1.7	3
137	Novel acetylcholinesterase reactivator K112 and its cholinergic properties. Biomedicine and Pharmacotherapy, 2010, 64, 541-545.	5.6	12
138	Reactivation of VX-inhibited AChE by novel oximes having two oxygen atoms in the linker. Environmental Toxicology and Pharmacology, 2010, 30, 85-87.	4.0	7
139	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
140	New antioxidant flavonoid isolated from <i>Leuzea carthamoides</i> . Journal of Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 143-145.	5.2	5
141	A comparison of tabun-inhibited rat brain acetylcholinesterase reactivation by three oximes (HI-6,) Tj ETQq1 Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 790-797.	1 0.784314 rg 5.2	BT /Overlock 6
142	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
143	Why acetylcholinesterase reactivators do not work in butyrylcholinesterase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2010, 25, 318-322.	5.2	14
144	New Bisquaternary Isoquinolinium Inhibitors of Brain Cholinesterases - Synthesis and Anticholinesterase Activity. Letters in Drug Design and Discovery, 2010, 7, 1-4.	0.7	4

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145	Novel Nucleophilic Compounds with Oxime Group as Reactivators of Paraoxon-Inhibited Cholinesterases. Letters in Drug Design and Discovery, 2010, 7, 260-264.	0.7	3
146	<i>In vitro</i> reactivation of trichlorfon-inhibited butyrylcholinesterase using HI-6, obidoxime, pralidoxime and K048. Journal of Enzyme Inhibition and Medicinal Chemistry, 2009, 24, 680-683.	5.2	5
147	Preparation and In Vitro Evaluation of Monoquaternary Inhibitors of Brain Cholinesterases. Letters in Organic Chemistry, 2009, 6, 500-503.	0.5	1
148	Effect of acetylcholinesterase oxime-type reactivators K-48 and HI-6 on human liver microsomal cytochromes P450 invitro. Chemico-Biological Interactions, 2009, 180, 449-453.	4.0	6
149	Methylacridinium and its Cholinergic Properties. Neurotoxicity Research, 2009, 16, 372-377.	2.7	6
150	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
151	Reactivation of Human Brain Homogenate Cholinesterases Inhibited by Tabun using Newly Developed Oximes K117 and K127. Basic and Clinical Pharmacology and Toxicology, 2009, 105, 207-210.	2.5	8
152	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
153	Development of new antidotes of organophosphate intoxications: Oxime-assisted reactivation of dimethoxy- and diethoxy-phosphorylated human butyrylcholinesterase for construction of "pseudo catalytic―bioscavengers. Toxicology Letters, 2009, 189, S216.	0.8	5
154	Commercially available antidotes of organophosphate poisonings (pralidoxime, obidoxime, methoxime,) Tj ETQq inhibited by selected organophosphate pesticides. Toxicology Letters, 2009, 189, S217.	0 0 0 rgBT 0.8	/Overlock 10 0
155	Global Impact of Chemical Warfare Agents Used Before and After 1945. , 2009, , 17-24.		11
156	Other Toxic Chemicals as Potential Chemical Warfare Agents. , 2009, , 331-338.		1
157	<i>In vitro</i> oxime-assisted reactivation of paraoxon-inhibited human acetylcholinesterase and butyrylcholinesterase. Clinical Toxicology, 2009, 47, 545-550.	1.9	38
158	In vitro identification of novel acetylcholinesterase reactivators. Toxin Reviews, 2009, 28, 238-244.	3.4	8
159	Novel Bisquaternary Oximes—Reactivation of Acetylcholinesterase and Butyrylcholinesterase Inhibited by Paraoxon. Molecules, 2009, 14, 4915-4921.	3.8	17
160	Chemical Aspects of Pharmacological Prophylaxis Against Nerve Agent Poisoning. Current Medicinal Chemistry, 2009, 16, 2977-2986.	2.4	65
161	Novel Oximes. , 2009, , 997-1021.		6
162	Could oxime HI-6 really be considered as "broad-spectrum" antidote?. Journal of Applied Biomedicine, 2009, 7, 143-149.	1.7	28

#	Article	IF	CITATIONS
163	Inhibition of blood cholinesterases by nerve agents in vitro. Journal of Applied Biomedicine, 2009, 7, 201-206.	1.7	4
164	Pharmacological Prophylaxis Against Nerve Agent Poisoning: Experimental Studies and Practical Implications. , 2009, , 977-984.		0
165	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
166	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
167	Preparation and antiplatelet activity of glycidic acid derivatives. Chemical Papers, 2008, 62, .	2.2	0
168	The Influence of Acetylcholinesterase Reactivators on Selected Hepatic Functions in Rats. Basic and Clinical Pharmacology and Toxicology, 2008, 103, 119-123.	2.5	19
169	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
170	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
171	Reactivation of DFP- and paraoxon-inhibited acetylcholinesterases by pyridinium oximes. Chemico-Biological Interactions, 2008, 175, 365-367.	4.0	26
172	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
173	Aflatoxin assay using an acetylcholinesterase based biosensor. Toxicology Letters, 2008, 180, S73.	0.8	0
174	Amperometric Biosensors for Real Time Assays of Organophosphates. Sensors, 2008, 8, 5303-5312.	3.8	47
175	DPPH Radical Scavenging Activity of Several Naturally Occurring Coumarins and Their Synthesized Analogs Measured by the SIA Method. Toxicology Mechanisms and Methods, 2008, 18, 413-418.	2.7	12
176	<i>In Vitro</i> Antiplatelet Activity of Flavonoids from <i>Leuzea Carthamoides</i> . Drug and Chemical Toxicology, 2008, 31, 27-35.	2.3	28
177	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
178	TLC Analysis of Intermediates Arising During the Preparation of Oxime HI-6 Dimethanesulfonate. Journal of Chromatographic Science, 2008, 46, 316-319.	1.4	30
179	Improvement of acetylcholinesterase-based assay for organophosphates in way of identification by reactivators. Talanta, 2008, 77, 451-454.	5.5	55
180	Daphnia intoxicated by nerve agent tabun can be treated using human antidotes. Environmental Toxicology and Pharmacology, 2008, 25, 329-333.	4.0	7

#	Article	IF	CITATIONS
181	Efficacy and dosing of antidotes applied to Daphnia intoxicated by nerve agent tabun. Environmental Toxicology and Pharmacology, 2008, 26, 283-289.	4.0	4
182	Synthesis, Structure, and Cholinergic Effect of Novel Neuroprotective Compounds Bearing the Tacrine Pharmacophore. Heterocycles, 2008, 76, 1219.	0.7	20
183	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
184	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
185	In Vitro Comparison of Two Most Promising H-Oximes (HI-6 and HLö-7) and Currently Commercially Available Reactivators Pralidoxime and Obidoxime in Reactivation of Cyclosarin-Inhibited Human Cholinesterases. Toxicology Mechanisms and Methods, 2008, 18, 329-333.	2.7	1
186	High-performance Liquid Chromatography Analysis of Four Leuzea carthamoides Flavonoids. Journal of Chromatographic Science, 2008, 46, 162-164.	1.4	4
187	Planar Ni(II) 1,2-dithiolenes involving bidentate <i>N</i> -donor ligands. Journal of Coordination Chemistry, 2008, 61, 3525-3533.	2.2	7
188	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
189	Sensor System Based on Acetylcholinesterase in Homogenous Phase for Analysis of Paraoxon. Analytical Letters, 2008, 41, 2214-2223.	1.8	8
190	Metabolic Pathways of T-2 Toxin. Current Drug Metabolism, 2008, 9, 77-82.	1.2	81
191	Potency of Novel Oximes to Reactivate Sarin Inhibited Human Cholinesterases. Drug and Chemical Toxicology, 2008, 31, 1-9.	2.3	11
192	Condensed and Hydrolysable Tannins as Antioxidants Influencing the Health. Mini-Reviews in Medicinal Chemistry, 2008, 8, 436-447.	2.4	218
193	Cholinesterase Biosensor Construction – A Review. Protein and Peptide Letters, 2008, 15, 795-798.	0.9	30
194	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
195	Aflatoxin Assay Using an Amperometric Sensor Strip and Acetylcholinesterase as Recognition Element. Sensor Letters, 2008, 6, 450-453.	0.4	16
196	Preparation of Oxime HI-6 (Dichloride and Dimethanesulphonate) - Antidote against Nerve Agents. Defence Science Journal, 2008, 58, 399-404.	0.8	10
197	Optimization of acetylcholinesterase immobilization onto screen printed platinum electrode. Journal of Applied Biomedicine, 2008, 6, 27-30.	1.7	3
198	Prevention of the production of T-2 toxin and methods of minimizing its toxic effects. Kontakt, 2008, 10, 200-208.	0.2	0

#	Article	IF	CITATIONS
199	The effect of HI-6 on cholinesterases and on the cholinergic system of the rat bladder. Neuroendocrinology Letters, 2008, 29, 759-62.	0.2	6
200	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
201	Potency of Five Structurally Different Acetylcholinesterase Reactivators to Reactivate Human Brain Cholinesterases Inhibited by Cyclosarin. Clinical Toxicology, 2007, 45, 512-515.	1.9	8
202	Treatment of Organophosphate Intoxication Using Cholinesterase Reactivators:Facts and Fiction. Mini-Reviews in Medicinal Chemistry, 2007, 7, 461-466.	2.4	126
203	Progress in Synthesis of New Acetylcholinesterase Reactivators During the Period 1990-2004. Current Organic Chemistry, 2007, 11, 229-238.	1.6	78
204	Structural Factors Influencing Potency of Currently Used Acetylcholinesterase Reactivators for Treatment of Cyclosarin Intoxications. Current Pharmaceutical Design, 2007, 13, 3445-3452.	1.9	10
205	Cholinesterase Reactivators: The Fate and Effects in the Organism Poisoned with Organophosphates/Nerve Agents. Current Drug Metabolism, 2007, 8, 803-809.	1.2	31
206	Structure-Activity Relationship of Acetylcholinesterase Reactivators -Antidotes Against Nerve Agents. Letters in Organic Chemistry, 2007, 4, 212-217.	0.5	10
207	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
208	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
209	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
210	Targeted Synthesis of 1-(4-Hydroxyiminomethylpyridinium)-3-pyridiniumpropane Dibromide – A New Nerve Agent Reactivator. Molecules, 2007, 12, 1964-1972.	3.8	15
211	Acetylcholinesterases – the structural similarities and differences. Journal of Enzyme Inhibition and Medicinal Chemistry, 2007, 22, 417-424.	5.2	110
212	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
213	HPLC Analysis of HIâ€6 Dichloride and Dimethanesulfonate—Antidotes against Nerve Agents and Organophosphorus Pesticides. Analytical Letters, 2007, 40, 2783-2787.	1.8	51
214	Mycotoxin Assays Using Biosensor Technology: A Review. Drug and Chemical Toxicology, 2007, 30, 253-261.	2.3	53
215	Amperometric Biosensor for Evaluation of Competitive Cholinesterase Inhibition by the Reactivator Hlâ€6. Analytical Letters, 2007, 40, 2351-2359.	1.8	23
216	Design of a Potent Reactivator of Tabun-Inhibited AcetylcholinesteraseSynthesis and Evaluation of (<i>E</i>)-1-(4-Carbamoylpyridinium)-4-(4-hydroxyiminomethylpyridinium)-but-2-ene Dibromide (K2O3). Journal of Medicinal Chemistry, 2007, 50, 5514-5518.	6.4	100

#	Article	IF	CITATIONS
217	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
218	Two Step Synthesis of a Non-symmetric Acetylcholinesterase Reactivator. Molecules, 2007, 12, 1755-1761.	3.8	9
219	Preparation of Benzalkonium Salts Differing in the Length of a Side Alkyl Chain. Molecules, 2007, 12, 2341-2347.	3.8	19
220	Inhibition of blood cholinesterases following intoxication with VX and its derivatives. Journal of Applied Toxicology, 2007, 27, 458-463.	2.8	14
221	Monooxime reactivators of acetylcholinesterase with (E)-but-2-ene linker—Preparation and reactivation of tabun- and paraoxon-inhibited acetylcholinesterase. Bioorganic and Medicinal Chemistry, 2007, 15, 6733-6741.	3.0	52
222	Novel series of bispyridinium compounds bearing a (Z)-but-2-ene linker—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
223	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
224	Comparison of Reactivating and Therapeutic Efficacy of Two Salts of the Oxime HI-6 against Tabun, Soman and Cyclosarin in Rats. Basic and Clinical Pharmacology and Toxicology, 2007, 101, 328-332.	2.5	12
225	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
226	Structure $\hat{a} \in \hat{a}$ activity relationships for in vitro oxime reactivation of chlorpyrifos-inhibited acetylcholinesterase. Chemical Papers, 2007, 61, .	2.2	3
227	In vitro reactivation potency of acetylcholinesterase reactivators — K074 and K075 — to reactivate tabun-inhibited human brain cholinesterases. Neurotoxicity Research, 2007, 11, 101-106.	2.7	21
228	Potency of new structurally different oximes to reactivate cyclosarin inhibited-human brain acetylcholinesterases. Toxicology, 2007, 233, 231-232.	4.2	1
229	Phosphotriesterase modified by poly[N-(2-hydroxypropyl)methacrylamide]. Toxicology, 2007, 233, 235.	4.2	4
230	Biohazards of Protein Biotoxins. Defence Science Journal, 2007, 57, 825-837.	0.8	3
231	TLC identification of benzalkonium bromide homologs. Journal of Planar Chromatography - Modern TLC, 2007, 20, 283-285.	1.2	1
232	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
233	Assay of radical scavenging activity of antidotes against chemical warfare agents by DPPH test using sequential injection technique. Journal of Applied Biomedicine, 2007, 5, 81-84.	1.7	11
234	Amperometric biosensor for pesticide methamidophos assay. Acta Medica (Hradec Kralove), 2007, 50, 239-41.	0.5	2

#	Article	IF	CITATIONS
235	In Vitro Potency of H Oximes (HI-6, HLö-7), the Oxime BI-6, and Currently Used Oximes (Pralidoxime,) Tj ETQq1 of Toxicology and Environmental Health - Part A: Current Issues, 2006, 69, 1431-1440.	1 0.78431 2.3	4 rgBT /Ove 9
236	In VitroEvaluation of Acetylcholinesterase Reactivators as Potential Antidotes Against Tabun Nerve Agent Poisonings. Drug and Chemical Toxicology, 2006, 29, 443-449.	2.3	12
237	New group of xylene linker-containing acetylcholinesterase reactivators as antidotes against the nerve agent cyclosarin. Journal of Enzyme Inhibition and Medicinal Chemistry, 2006, 21, 515-519.	5.2	8
238	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
239	Structure-activity relationship for the reactivators of acetylcholinesterase inhibited by nerve agent VX. Toxicology Letters, 2006, 164, S51-S52.	0.8	2
240	In vitro comparison of eight acetylcholinesterase reactivators to reactivate VX inhibited ACHE. Toxicology Letters, 2006, 164, S120.	0.8	2
241	Russian VX: Inhibition and Reactivation of Acetylcholinesterase Compared with VX Agent. Basic and Clinical Pharmacology and Toxicology, 2006, 98, 389-394.	2.5	36
242	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
243	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
244	Pretreatment with pyridinium oximes improves antidotal therapy against tabun poisoning. Toxicology, 2006, 228, 41-50.	4.2	51
245	New methods in synthesis of acetylcholinesterase reactivators and evaluation of their potency to reactivate cyclosarin-inhibited AChE. Chemical Papers, 2006, 60, .	2.2	39
246	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
247	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
248	The Acute Toxicity of Acetylcholinesterase Reactivators in Mice in Relation to Their Structure. Neurotoxicity Research, 2006, 9, 291-296.	2.7	43
249	Reactivation of sarin-inhibited pig brain acetylcholinesterase using oxime antidotes. Journal of Medical Toxicology, 2006, 2, 141-146.	1.5	7
250	Effect of some acetylcholinesterase reactivators on human platelet aggregationin vitro. Journal of Applied Toxicology, 2006, 26, 258-261.	2.8	7
251	Evaluation of reactivation test in anaesthetized dogs with experimental intoxication with nerve agents. Journal of Applied Toxicology, 2006, 26, 439-443.	2.8	9
252	Structural Requirements of Acetylcholinesterase Reactivators. Mini-Reviews in Medicinal Chemistry, 2006, 6, 269-277.	2.4	199

#	Article	IF	CITATIONS
253	The Reactivating and Therapeutic Efficacy of Oximes to Counteract Russian VX Poisonings. International Journal of Toxicology, 2006, 25, 397-401.	1.2	23
254	Toxicity of the nerve agent tabun toDaphnia magna, a new experimental species in military toxicology. Chemistry and Ecology, 2006, 22, 175-180.	1.6	8
255	Botulinum Toxin; Bioterror and Biomedicinal Agent. Defence Science Journal, 2006, 56, 189-197.	0.8	5
256	Prophylaxis Against Nerve Agent Intoxications. Defence Science Journal, 2006, 56, 775-784.	0.8	28
257	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
258	Reactivation of acetycholinesterase inhibited by the pesticide chlorpyrifos. Journal of Applied Biomedicine, 2006, 4, 147-151.	1.7	4
259	Chemical terrorism. Kontakt, 2006, 8, 123-127.	0.2	3
260	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
261	(09) Comparison of the potency of newly developed and currently available oximes to reactivate nerve agent-inhibited acetylcholinesterase in vitro and in vivo. Chemico-Biological Interactions, 2005, 157-158, 367-368.	4.0	9
262	(22) Comparison of ability of some oximes to reactivate sarin-inhibited brain acetylcholinesterase from different species. Chemico-Biological Interactions, 2005, 157-158, 385-387.	4.0	6
263	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
264	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
265	Effective bisquaternary reactivators of tabun-inhibited AChE. Journal of Applied Toxicology, 2005, 25, 491-495.	2.8	54
266	In-Vitro Searching for a New Potent Reactivator of Acetylcholinesterase Inhibited by Nerve Agent VX. Letters in Drug Design and Discovery, 2005, 2, 23-25.	0.7	8
267	Bispyridinium Oximes As Antidotal Treatment of Cyclosarin Poisoning—In Vitro and In Vivo Testing. International Journal of Toxicology, 2005, 24, 399-402.	1.2	8
268	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
269	A Comparison of the Potency of the Oxime HLö-7 and Currently Used Oximes (HI-6, Pralidoxime,) Tj ETQq1 1 0 Acta Medica (Hradec Kralove), 2005, 48, 81-86.	.784314 r 0.5	gBT /Overloci 17
270	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

#	Article	IF	CITATIONS
271	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
272	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
273	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
274	Reactivation of organophosphate-inhibited acetylcholinesterase by quaternary pyridinium aldoximes. Neurotoxicity Research, 2004, 6, 565-570.	2.7	24
275	Acetylcholinesterase and Butyrylcholinesterase – Important Enzymes of Human Body. Acta Medica (Hradec Kralove), 2004, 47, 215-228.	0.5	74
276	Preparation of 2-(4-{[4-(Quinolin-2-ylmethoxy)phenyl]sulfanyl}phenyl) Propionic Acid (VUFB 20615) and 2-Methyl-2-(4-{[4-(quinolin-2- ylmethoxy)Phenyl]sulfanyl}phenyl)Propionic Acid (VUFB 20623) as Potential Antileukotrienic Agents. Current Organic Chemistry, 2004, 8, 1235-1243.	1.6	10
277	Acetylcholinesterase and butyrylcholinesteraseimportant enzymes of human body. Acta Medica (Hradec Kralove), 2004, 47, 215-28.	0.5	29
278	Progress in Antidotes (Acetylcholinesterase Reactivators) Against Organophosphorus Pesticides. , 0, ,		5