

# Vendula Sepsova

## List of Publications by Year in descending order

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56  
papers

1,441  
citations

393982

19  
h-index

329751

37  
g-index

56  
all docs

56  
docs citations

56  
times ranked

1947  
citing authors

#	ARTICLE	IF	CITATIONS
1	Toxicity, pharmacokinetics, and effectiveness of the ortho-chlorinated bispyridinium oxime, K870. <i>Food and Chemical Toxicology</i> , 2022, 167, 113236.	1.8	1
2	Tacrine and its 7-methoxy derivate; time-change concentration in plasma and brain tissue and basic toxicological profile in rats. <i>Drug and Chemical Toxicology</i> , 2021, 44, 207-214.	1.2	6
3	Development of versatile and potent monoquatarnary reactivators of acetylcholinesterase. <i>Archives of Toxicology</i> , 2021, 95, 985-1001.	1.9	7
4	Tacrine – Benzothiazoles: Novel class of potential multitarget anti-Alzheimer's drugs dealing with cholinergic, amyloid and mitochondrial systems. <i>Bioorganic Chemistry</i> , 2021, 107, 104596.	2.0	17
5	7-phenoxytacrine is a dually acting drug with neuroprotective efficacy in vivo. <i>Biochemical Pharmacology</i> , 2021, 186, 114460.	2.0	12
6	( $\Delta\pm$ )- <b>BIGI-3h</b> : Pentatarget-Directed Ligand combining Cholinesterase, Monoamine Oxidase, and Glycogen Synthase Kinase 3 $\beta$ Inhibition with Calcium Channel Antagonism and Antiaggregating Properties for Alzheimer's Disease. <i>ACS Chemical Neuroscience</i> , 2021, 12, 1328-1342.	1.7	21
7	Phenothiazine-Tacrine Heterodimers: Pursuing Multitarget Directed Approach in Alzheimer's Disease. <i>ACS Chemical Neuroscience</i> , 2021, 12, 1698-1715.	1.7	16
8	Design and synthesis of novel tacrine-indole hybrids as potential multitarget-directed ligands for the treatment of Alzheimer's disease. <i>Future Medicinal Chemistry</i> , 2021, 13, 785-804.	1.1	5
9	Huprine Y – Tryptophan heterodimers with potential implication to Alzheimer's disease treatment. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2021, 43, 128100.	1.0	5
10	Pyridostigmine bromide and its relation to Gulf War illness. <i>Toxin Reviews</i> , 2020, 39, 138-146.	1.5	2
11	Synthesis, <i>in vitro</i> screening and molecular docking of isoquinolinium-5-carbaldoximes as acetylcholinesterase and butyrylcholinesterase reactivators. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2020, 35, 478-488.	2.5	15
12	Encapsulation of oxime K027 into cucurbit[7]uril: In vivo evaluation of safety, absorption, brain distribution and reactivation effectiveness. <i>Toxicology Letters</i> , 2020, 320, 64-72.	0.4	10
13	Cysteine-Targeted Insecticides against <i>A. gambiae</i> Acetylcholinesterase Are Neither Selective nor Reversible Inhibitors. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 65-71.	1.3	11
14	Discovery of novel berberine derivatives with balanced cholinesterase and prolyl oligopeptidase inhibition profile. <i>European Journal of Medicinal Chemistry</i> , 2020, 203, 112593.	2.6	24
15	Exploring Structure-Activity Relationship in Tacrine-Squaramide Derivatives as Potent Cholinesterase Inhibitors. <i>Biomolecules</i> , 2019, 9, 379.	1.8	23
16	Pharmacological and toxicological in vitro and in vivo effect of higher doses of oxime reactivators. <i>Toxicology and Applied Pharmacology</i> , 2019, 383, 114776.	1.3	5
17	Tacroximes: novel unique compounds for the recovery of organophosphorus-inhibited acetylcholinesterase. <i>Future Medicinal Chemistry</i> , 2019, 11, 2625-2634.	1.1	6
18	Novel tacrine-tryptophan hybrids: Multi-target directed ligands as potential treatment for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2019, 168, 491-514.	2.6	75

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19	Combination of Memantine and 6-Chlorotacrine as Novel Multi-Target Compound against Alzheimer's Disease. <i>Current Alzheimer Research</i> , 2019, 16, 821-833.	0.7	17
20	Synthesis and biological assessment of Kojotacrines as new agents for Alzheimer's disease therapy. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019, 34, 163-170.	2.5	19
21	N-alkylated Tacrine Derivatives as Potential Agents in Alzheimer's Disease Therapy. <i>Current Alzheimer Research</i> , 2019, 16, 333-343.	0.7	5
22	Design, Synthesis, and Biological Evaluation of 1-Benzylamino-2-hydroxyalkyl Derivatives as New Potential Disease-Modifying Multifunctional Anti-Alzheimer's Agents. <i>ACS Chemical Neuroscience</i> , 2018, 9, 1074-1094.	1.7	47
23	Synthesis, Biological Assessment and Molecular Modeling of Racemic QuinopyranoTacrines for Alzheimer's Disease Therapy. <i>ChemistrySelect</i> , 2018, 3, 461-466.	0.7	10
24	Investigation of New Orexin 2 Receptor Modulators Using In Silico and In Vitro Methods. <i>Molecules</i> , 2018, 23, 2926.	1.7	6
25	Development of small bisquaternary cholinesterase inhibitors as drugs for pre-treatment of nerve agent poisonings. <i>Drug Design, Development and Therapy</i> , 2018, Volume 12, 505-512.	2.0	4
26	In vitro and in silico Evaluation of Non-Quaternary Reactivators of AChE as Antidotes of Organophosphorus Poisoning - a New Hope or a Blind Alley?. <i>Medicinal Chemistry</i> , 2018, 14, 281-292.	0.7	19
27	Tacrine-resveratrol fused hybrids as multi-target-directed ligands against Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2017, 127, 250-262.	2.6	95
28	A comparison of the reactivating and therapeutic efficacy of two novel bispyridinium oximes (K305,) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 Biomedicine, 2017, 15, 49-53.	0.6	3
29	Multi-target-directed therapeutic potential of 7-methoxytacrine-adamantylamine heterodimers in the Alzheimer's disease treatment. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2017, 1863, 607-619.	1.8	37
30	Novel Tacrine-Scutellarin Hybrids as Multipotent Anti-Alzheimer's Agents: Design, Synthesis and Biological Evaluation. <i>Molecules</i> , 2017, 22, 1006.	1.7	32
31	Development of 2-Methoxyhuprine as Novel Lead for Alzheimer's Disease Therapy. <i>Molecules</i> , 2017, 22, 1265.	1.7	26
32	The Evaluation of the Reactivating and Neuroprotective Efficacy of Two Newly Prepared Bispyridinium Oximes (K305, K307) in Tabun-Poisoned Rats: A Comparison with Trimedoxime and the Oxime K203. <i>Molecules</i> , 2017, 22, 1152.	1.7	8
33	Design, Synthesis and in vitro Evaluation of Indolotacrine Analogues as Multitarget-Directed Ligands for the Treatment of Alzheimer's Disease. <i>ChemMedChem</i> , 2016, 11, 1264-1269.	1.6	35
34	Targeting copper(II)-induced oxidative stress and the acetylcholinesterase system in Alzheimer's disease using multifunctional tacrine-coumarin hybrid molecules. <i>Journal of Inorganic Biochemistry</i> , 2016, 161, 52-62.	1.5	63
35	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. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9967-9973.	2.9	83
36	Interaction of 7-Methoxytacrine-Adamantylamine Cholinesterase Inhibitors with Nicotinic and Muscarinic Acetylcholine Receptors. <i>Biophysical Journal</i> , 2015, 108, 430a.	0.2	0

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37	7-Methoxytacrine-p-Anisidine Hybrids as Novel Dual Binding Site Acetylcholinesterase Inhibitors for Alzheimer's Disease Treatment. <i>Molecules</i> , 2015, 20, 22084-22101.	1.7	35
38	Cholinergic properties of new 7-methoxytacrine-donepezil derivatives. <i>General Physiology and Biophysics</i> , 2015, 34, 189-200.	0.4	17
39	A Comparison of the Reactivating and Therapeutic Efficacy of Two Newly Developed Oximes (K727 and Tj ETQq1 1 0.784314 rgBT /O Pharmacology and Toxicology, 2015, 116, 367-371.	1.2	10
40	A comparison of the reactivating and therapeutic efficacy of two novel bispyridinium oximes (K727,) Tj ETQq0 0 0 rgBT /Overlock 10 TF 5 Methods, 2015, 25, 229-233.	1.3	5
41	Tacrine-Trox Hybrids: A Novel Class of Centrally Active, Nonhepatotoxic Multi-Target-Directed Ligands Exerting Anticholinesterase and Antioxidant Activities with Low In Vivo Toxicity. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 8985-9003.	2.9	121
42	A comparison of the reactivating and therapeutic efficacy of two novel bispyridinium oximes (K920,) Tj ETQq0 0 0 rgBT /Overlock 10 TF 5 Biomedicine, 2015, 13, 299-304.	0.6	1
43	The evaluation of the reactivating and therapeutic efficacy of two novel oximes (K361 and K378) in comparison with the oxime K203 and trimedoxime in tabun-poisoned rats and mice. <i>Toxicology Mechanisms and Methods</i> , 2014, 24, 173-178.	1.3	6
44	The Evaluation of Prophylactic Efficacy of Newly Developed Reversible Inhibitors of Acetylcholinesterase in Soman-Poisoned Mice - A Comparison with Commonly Used Pyridostigmine. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2014, 115, 571-576.	1.2	7
45	A comparison of the reactivating and therapeutic efficacy of two novel oximes K378 and K458 with currently available oximes in rats and mice poisoned with sarin. <i>Journal of Applied Biomedicine</i> , 2014, 12, 155-160.	0.6	1
46	7-MEOTA-donepezil like compounds as cholinesterase inhibitors: Synthesis, pharmacological evaluation, molecular modeling and QSAR studies. <i>European Journal of Medicinal Chemistry</i> , 2014, 82, 426-438.	2.6	80
47	Outcomes of Alzheimer's disease therapy with acetylcholinesterase inhibitors and memantine. <i>Expert Opinion on Drug Safety</i> , 2014, 13, 759-74.	1.0	209
48	A comparison of the reactivating efficacy of a novel bispyridinium oxime K203 with currently available oximes in VX agent-poisoned rats. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2013, 28, 753-757.	2.5	3
49	The evaluation of the reactivating and therapeutic efficacy of three novel bispyridinium oximes (K454,) Tj ETQq1 1 0.784314 rgBT /Over Toxicology Mechanisms and Methods, 2013, 23, 94-98.	1.3	4
50	Oximes: Inhibitors of Human Recombinant Acetylcholinesterase. A Structure-Activity Relationship (SAR) Study. <i>International Journal of Molecular Sciences</i> , 2013, 14, 16882-16900.	1.8	38
51	Acetylcholinesterase Reactivators (HI-6, Obidoxime, Trimedoxime, K027, K075, K127, K203, K282): Structural Evaluation of Human Serum Albumin Binding and Absorption Kinetics. <i>International Journal of Molecular Sciences</i> , 2013, 14, 16076-16086.	1.8	12
52	A Resurrection of 7-MEOTA: A Comparison with Tacrine. <i>Current Alzheimer Research</i> , 2013, 10, 893-906.	0.7	92
53	A comparison of the reactivating and therapeutic efficacy of the newly developed bispyridinium oxime K203 with currently available oximes, in sarin poisoned rats and mice. <i>Journal of Applied Biomedicine</i> , 2011, 9, 225-230.	0.6	11
54	The Benefit of Combinations of Oximes for the Reactivating and Therapeutic Efficacy of Antidotal Treatment of Sarin Poisoning in Rats and Mice. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2011, 109, 30-34.	1.2	9

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55	A Comparison of the Reactivating and Therapeutic Efficacy of Chosen Combinations of Oximes With Individual Oximes Against VX in Rats and Mice. International Journal of Toxicology, 2011, 30, 562-567.	0.6	8
56	OXIMES AS INHIBITORS OF ACETYLHOLINESTERASE - A STRUCTURE-ACTIVITY RELATIONSHIP (SAR) STUDY. Military Medical Science Letters (Vojenske Zdravotnicke Listy), 2011, 80, 178-186.	0.2	2