

Maria Laura Bolognesi

List of Publications by Citations

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171
papers

7,395
citations

46
h-index

80
g-index

198
ext. papers

8,376
ext. citations

6.1
avg, IF

6.1
L-index

| # | Paper | IF | Citations |
|-----|--|------|-----------|
| 171 | Multi-target-directed ligands to combat neurodegenerative diseases. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 347-72 | 8.3 | 816 |
| 170 | A perspective on multi-target drug discovery and design for complex diseases. <i>Clinical and Translational Medicine</i> , 2018 , 7, 3 | 5.7 | 307 |
| 169 | Insight into the kinetic of amyloid beta (1-42) peptide self-aggregation: elucidation of inhibitors' mechanism of action. <i>ChemBioChem</i> , 2007 , 8, 2152-61 | 3.8 | 289 |
| 168 | Multi-target-directed drug design strategy: from a dual binding site acetylcholinesterase inhibitor to a trifunctional compound against Alzheimer's disease. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 6446-9 | 8.3 | 225 |
| 167 | Rational approach to discover multipotent anti-Alzheimer drugs. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 360-3 | 8.3 | 206 |
| 166 | Tacrine derivatives and Alzheimer's disease. <i>Current Medicinal Chemistry</i> , 2010 , 17, 1825-38 | 4.3 | 180 |
| 165 | Neglected tropical diseases: multi-target-directed ligands in the search for novel lead candidates against Trypanosoma and Leishmania. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 7339-59 | 8.3 | 179 |
| 164 | Inhibition of acetylcholinesterase, beta-amyloid aggregation, and NMDA receptors in Alzheimer's disease: a promising direction for the multi-target-directed ligands gold rush. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 4381-4 | 8.3 | 170 |
| 163 | A small molecule targeting the multifactorial nature of Alzheimer's disease. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 3689-92 | 16.4 | 156 |
| 162 | Propidium-based polyamine ligands as potent inhibitors of acetylcholinesterase and acetylcholinesterase-induced amyloid-beta aggregation. <i>Journal of Medicinal Chemistry</i> , 2005 , 48, 24-7 | 8.3 | 127 |
| 161 | Multitarget drug design strategy: quinone-tacrine hybrids designed to block amyloid- β aggregation and to exert anticholinesterase and antioxidant effects. <i>Journal of Medicinal Chemistry</i> , 2014 , 57, 8576-89 | 8.3 | 122 |
| 160 | Imaging of β amyloid plaques by near infrared fluorescent tracers: a new frontier for chemical neuroscience. <i>Chemical Society Reviews</i> , 2015 , 44, 1807-19 | 58.5 | 116 |
| 159 | Novel class of quinone-bearing polyamines as multi-target-directed ligands to combat Alzheimer's disease. <i>Journal of Medicinal Chemistry</i> , 2007 , 50, 4882-97 | 8.3 | 113 |
| 158 | The Hippo Pathway and YAP/TAZ-TEAD Protein-Protein Interaction as Targets for Regenerative Medicine and Cancer Treatment. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 4857-73 | 8.3 | 109 |
| 157 | Design, synthesis, and biological evaluation of conformationally restricted rivastigmine analogues. <i>Journal of Medicinal Chemistry</i> , 2004 , 47, 5945-52 | 8.3 | 108 |
| 156 | Memoquin: a multi-target-directed ligand as an innovative therapeutic opportunity for Alzheimer's disease. <i>Neurotherapeutics</i> , 2009 , 6, 152-62 | 6.4 | 101 |
| 155 | Alzheimer's disease: new approaches to drug discovery. <i>Current Opinion in Chemical Biology</i> , 2009 , 13, 303-8 | 9.7 | 100 |

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| 154 | Multitarget drug discovery for Alzheimer's disease: triazinones as BACE-1 and GSK-3 β inhibitors. <i>Angewandte Chemie - International Edition</i> , 2015 , 54, 1578-82 | 16.4 | 87 |
| 153 | Molecular Hybridization as a Tool for Designing Multitarget Drug Candidates for Complex Diseases. <i>Current Topics in Medicinal Chemistry</i> , 2019 , 19, 1694-1711 | 3 | 76 |
| 152 | Tacrine-resveratrol fused hybrids as multi-target-directed ligands against Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2017 , 127, 250-262 | 6.8 | 74 |
| 151 | Acetylcholinesterase noncovalent inhibitors based on a polyamine backbone for potential use against Alzheimer's disease. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 4186-9 | 8.3 | 73 |
| 150 | Exploiting the lipoic acid structure in the search for novel multitarget ligands against Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2011 , 46, 5435-42 | 6.8 | 71 |
| 149 | Multifunctional tacrine derivatives in Alzheimer's disease. <i>Current Topics in Medicinal Chemistry</i> , 2013 , 13, 1771-86 | 3 | 71 |
| 148 | Cystamine-tacrine dimer: a new multi-target-directed ligand as potential therapeutic agent for Alzheimer's disease treatment. <i>Neuropharmacology</i> , 2012 , 62, 997-1003 | 5.5 | 70 |
| 147 | Multitarget-directed drug design strategy: a novel molecule designed to block epidermal growth factor receptor (EGFR) and to exert proapoptotic effects. <i>Journal of Medicinal Chemistry</i> , 2006 , 49, 6642-53 | 8.3 | 70 |
| 146 | Synthesis of a small library of 2-phenoxy-1,4-naphthoquinone and 2-phenoxy-1,4-anthraquinone derivatives bearing anti-trypanosomal and anti-leishmanial activity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008 , 18, 2272-6 | 2.9 | 69 |
| 145 | BACE-1 Inhibitors: From Recent Single-Target Molecules to Multitarget Compounds for Alzheimer's Disease. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 619-637 | 8.3 | 66 |
| 144 | Cardanol-derived AChE inhibitors: Towards the development of dual binding derivatives for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2016 , 108, 687-700 | 6.8 | 66 |
| 143 | MTDL design strategy in the context of Alzheimer's disease: from lipocrine to memoquin and beyond. <i>Current Pharmaceutical Design</i> , 2009 , 15, 601-13 | 3.3 | 64 |
| 142 | Navigating the Chemical Space of Multitarget-Directed Ligands: From Hybrids to Fragments in Alzheimer's Disease. <i>Molecules</i> , 2016 , 21, 466 | 4.8 | 64 |
| 141 | The concept of privileged structures in rational drug design: focus on acridine and quinoline scaffolds in neurodegenerative and protozoan diseases. <i>Expert Opinion on Drug Discovery</i> , 2011 , 6, 251-68 | 6.2 | 63 |
| 140 | Prejunctional muscarinic inhibitory control of acetylcholine release in the human isolated detrusor: involvement of the M4 receptor subtype. <i>British Journal of Pharmacology</i> , 2000 , 129, 493-500 | 8.6 | 63 |
| 139 | Toward a rational design of multitarget-directed antioxidants: merging memoquin and lipoic acid molecular frameworks. <i>Journal of Medicinal Chemistry</i> , 2009 , 52, 7883-6 | 8.3 | 62 |
| 138 | Pharmacological characterization of memoquin, a multi-target compound for the treatment of Alzheimer's disease. <i>PLoS ONE</i> , 2013 , 8, e56870 | 3.7 | 62 |
| 137 | Novel 8-Hydroxyquinoline Derivatives as Multitarget Compounds for the Treatment of Alzheimer's Disease. <i>ChemMedChem</i> , 2016 , 11, 1284-95 | 3.7 | 59 |

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| 136 | Novel tacrine-grafted Ugi adducts as multipotent anti-Alzheimer drugs: a synthetic renewal in tacrine-ferulic acid hybrids. <i>ChemMedChem</i> , 2015 , 10, 523-39 | 3.7 | 56 |
| 135 | Two diseases, one approach: multitarget drug discovery in Alzheimer's and neglected tropical diseases. <i>MedChemComm</i> , 2014 , 5, 853-861 | 5 | 56 |
| 134 | Multitargeted drugs discovery: balancing anti-amyloid and anticholinesterase capacity in a single chemical entity. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011 , 21, 2655-8 | 2.9 | 55 |
| 133 | Tau-Centric Multitarget Approach for Alzheimer's Disease: Development of First-in-Class Dual Glycogen Synthase Kinase 3 and Tau-Aggregation Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2018 , 61, 7640-7656 | 8.3 | 53 |
| 132 | Structure-activity relationships of acetylcholinesterase noncovalent inhibitors based on a polyamine backbone. 4. Further investigation on the inner spacer. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 7308-12 | 8.3 | 52 |
| 131 | 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 | 6.8 | 49 |
| 130 | From dual binding site acetylcholinesterase inhibitors to multi-target-directed ligands (MTDLs): a step forward in the treatment of Alzheimer's disease. <i>Mini-Reviews in Medicinal Chemistry</i> , 2008 , 8, 960-7 ³⁻² | | 48 |
| 129 | 3,4-Dihydro-1,3,5-triazin-2(1H)-ones as the First Dual BACE-1/GSK-3 β Fragment Hits against Alzheimer's Disease. <i>ACS Chemical Neuroscience</i> , 2015 , 6, 1665-82 | 5.7 | 47 |
| 128 | Naphthoquinone derivatives exert their antitrypanosomal activity via a multi-target mechanism. <i>PLoS Neglected Tropical Diseases</i> , 2013 , 7, e2012 | 4.8 | 46 |
| 127 | Polyamines in drug discovery: from the universal template approach to the multitarget-directed ligand design strategy. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 5906-14 | 8.3 | 44 |
| 126 | Binding profile of the selective muscarinic receptor antagonist tripitramine. <i>European Journal of Pharmacology</i> , 1994 , 268, 459-62 | | 44 |
| 125 | From combinations to multitarget-directed ligands: A continuum in Alzheimer's disease polypharmacology. <i>Medicinal Research Reviews</i> , 2021 , 41, 2606-2633 | 14.4 | 41 |
| 124 | Prazosin-related compounds. Effect of transforming the piperazinyloquinazoline moiety into an aminomethyltetrahydroacridine system on the affinity for alpha1-adrenoreceptors. <i>Journal of Medicinal Chemistry</i> , 2003 , 46, 4895-903 | 8.3 | 41 |
| 123 | Synthesis and biological activity of some methoctramine-related tetraamines bearing a 11-acetyl-5,11-dihydro-6H-pyrido[2,3-b][1,4]-benzodiazepin-6-one moiety as antimuscarinics: a second generation of highly selective M2 muscarinic receptor antagonists. <i>Journal of Medicinal Chemistry</i> , 1993 , 36, 3734-7 | 8.3 | 41 |
| 122 | Changing paradigm to target microglia in neurodegenerative diseases: from anti-inflammatory strategy to active immunomodulation. <i>Expert Opinion on Therapeutic Targets</i> , 2016 , 20, 627-40 | 6.4 | 40 |
| 121 | Recent advances in alpha1-adrenoreceptor antagonists as pharmacological tools and therapeutic agents. <i>Current Topics in Medicinal Chemistry</i> , 2007 , 7, 147-62 | 3 | 40 |
| 120 | Toward the development of dual-targeted glyceraldehyde-3-phosphate dehydrogenase/trypanothione reductase inhibitors against <i>Trypanosoma brucei</i> and <i>Trypanosoma cruzi</i> . <i>ChemMedChem</i> , 2014 , 9, 371-82 | 3.7 | 39 |
| 119 | New tacrine dimers with antioxidant linkers as dual drugs: Anti-Alzheimer's and antiproliferative agents. <i>European Journal of Medicinal Chemistry</i> , 2017 , 138, 761-773 | 6.8 | 38 |

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| 118 | Design, synthesis, and biological evaluation of symmetrically and unsymmetrically substituted methoctramine-related polyamines as muscular nicotinic receptor noncompetitive antagonists. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 5212-23 | 8.3 | 37 |
| 117 | A Fluorescent Styrylquinoline with Combined Therapeutic and Diagnostic Activities against Alzheimer's and Prion Diseases. <i>ACS Medicinal Chemistry Letters</i> , 2013 , 4, 225-9 | 4.3 | 36 |
| 116 | 2-Phenoxy-1,4-naphthoquinones: From a Multitarget Antitrypanosomal to a Potential Antitumor Profile. <i>Journal of Medicinal Chemistry</i> , 2015 , 58, 6422-34 | 8.3 | 35 |
| 115 | Bis(7)-tacrine derivatives as multitarget-directed ligands: Focus on anticholinesterase and antiamyloid activities. <i>ChemMedChem</i> , 2010 , 5, 1215-20 | 3.7 | 35 |
| 114 | Privileged structure-guided synthesis of quinazoline derivatives as inhibitors of trypanothione reductase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 3031-5 | 2.9 | 34 |
| 113 | Discovery of a class of diketopiperazines as antiprion compounds. <i>ChemMedChem</i> , 2010 , 5, 1324-34 | 3.7 | 34 |
| 112 | Polyamine conjugation of curcumin analogues toward the discovery of mitochondria-directed neuroprotective agents. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 7264-8 | 8.3 | 32 |
| 111 | Lipoic acid, a lead structure for multi-target-directed drugs for neurodegeneration. <i>Mini-Reviews in Medicinal Chemistry</i> , 2006 , 6, 1269-74 | 3.2 | 32 |
| 110 | Universal template approach to drug design: polyamines as selective muscarinic receptor antagonists. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 4150-60 | 8.3 | 32 |
| 109 | Conjugation of quinones with natural polyamines: toward an expanded antitrypanosomatid profile. <i>Journal of Medicinal Chemistry</i> , 2012 , 55, 10490-500 | 8.3 | 31 |
| 108 | From Companion Diagnostics to Theranostics: A New Avenue for Alzheimer's Disease?. <i>Journal of Medicinal Chemistry</i> , 2016 , 59, 7759-70 | 8.3 | 31 |
| 107 | Tacripyrimidines, the first tacrine-dihydropyrimidine hybrids, as multi-target-directed ligands for Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2018 , 155, 839-846 | 6.8 | 30 |
| 106 | Parallel synthesis, evaluation, and preliminary structure-activity relationship of 2,5-diamino-1,4-benzoquinones as a novel class of bivalent anti-prion compound. <i>Journal of Medicinal Chemistry</i> , 2010 , 53, 8197-201 | 8.3 | 30 |
| 105 | Docking Ligands on Protein Surfaces: The Case Study of Prion Protein. <i>Journal of Chemical Theory and Computation</i> , 2009 , 5, 2565-73 | 6.4 | 30 |
| 104 | Small-molecule theranostic probes: a promising future in neurodegenerative diseases. <i>International Journal of Cell Biology</i> , 2013 , 2013, 150952 | 2.6 | 29 |
| 103 | Multitarget-directed ligands: innovative chemical probes and therapeutic tools against Alzheimer's disease. <i>Current Topics in Medicinal Chemistry</i> , 2011 , 11, 2797-806 | 3 | 29 |
| 102 | Parallel synthesis and cytotoxicity evaluation of a polyamine-quinone conjugates library. <i>Journal of Medicinal Chemistry</i> , 2008 , 51, 5463-7 | 8.3 | 29 |
| 101 | In vitro characterization of tripitramine, a polymethylene tetraamine displaying high selectivity and affinity for muscarinic M2 receptors. <i>British Journal of Pharmacology</i> , 1995 , 114, 1507-17 | 8.6 | 29 |

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| 100 | Structure-activity relationship and site of binding of polyamine derivatives at the nicotinic acetylcholine receptor. <i>FEBS Journal</i> , 2000 , 267, 110-20 | | 28 |
| 99 | Nutritional and Pharmacological Strategies to Regulate Microglial Polarization in Cognitive Aging and Alzheimer's Disease. <i>Frontiers in Aging Neuroscience</i> , 2017 , 9, 175 | 5.3 | 27 |
| 98 | Complementary medicinal chemistry-driven strategies toward new antitrypanosomal and antileishmanial lead drug candidates. <i>FEMS Immunology and Medical Microbiology</i> , 2010 , 58, 51-60 | | 27 |
| 97 | The bivalent ligand approach as a tool for improving the in vitro anti-Alzheimer multitarget profile of dimebon. <i>ChemMedChem</i> , 2013 , 8, 1276-81 | 3.7 | 26 |
| 96 | Design, synthesis, and biological and crystallographic evaluation of novel inhibitors of Plasmodium falciparum enoyl-ACP-reductase (PfFabI). <i>Journal of Medicinal Chemistry</i> , 2013 , 56, 7516-26 | 8.3 | 26 |
| 95 | WB 4101-related compounds. 2. Role of the ethylene chain separating amine and phenoxy units on the affinity for alpha(1)-adrenoreceptor subtypes and 5-HT(1A) receptors. <i>Journal of Medicinal Chemistry</i> , 1999 , 42, 4214-24 | 8.3 | 26 |
| 94 | Design, synthesis, and biological activity of prazosin-related antagonists. Role of the piperazine and furan units of prazosin on the selectivity for alpha1-adrenoreceptor subtypes. <i>Journal of Medicinal Chemistry</i> , 1998 , 41, 4844-53 | 8.3 | 25 |
| 93 | Progress in acetylcholinesterase inhibitors for Alzheimer's disease: an update. <i>Expert Opinion on Therapeutic Patents</i> , 2008 , 18, 387-401 | 6.8 | 24 |
| 92 | Hexahydrochromeno[4,3-b]pyrrole derivatives as acetylcholinesterase inhibitors. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 105-9 | 8.3 | 24 |
| 91 | Design, synthesis, and biological activity of methoctramine-related tetraamines bearing an 11-acetyl-5,11-dihydro-6H-pyrido[2,3-b][1,4] benzodiazepin-6-one moiety: structural requirements for optimum occupancy of muscarinic receptor subtypes as revealed by symmetrical and unsymmetrical polyamines. <i>Journal of Medicinal Chemistry</i> , 1994 , 37, 3363-72 | 8.3 | 24 |
| 90 | Approaches for discovering anti-prion compounds: lessons learned and challenges ahead. <i>Expert Opinion on Drug Discovery</i> , 2015 , 10, 389-97 | 6.2 | 23 |
| 89 | Location of the polyamine binding site in the vestibule of the nicotinic acetylcholine receptor ion channel. <i>Journal of Biological Chemistry</i> , 2001 , 276, 6151-60 | 5.4 | 23 |
| 88 | Sustainable production of pharmaceutical, nutraceutical and bioactive compounds from biomass and waste. <i>Chemical Society Reviews</i> , 2021 , 50, 11191-11207 | 58.5 | 23 |
| 87 | A small chemical library of 2-aminoimidazole derivatives as BACE-1 inhibitors: Structure-based design, synthesis, and biological evaluation. <i>European Journal of Medicinal Chemistry</i> , 2012 , 48, 206-13 | 6.8 | 22 |
| 86 | Synthesis of monomeric derivatives to probe memoquin's bivalent interactions. <i>Journal of Medicinal Chemistry</i> , 2011 , 54, 8299-304 | 8.3 | 22 |
| 85 | Structure-activity relationships of memoquin: Influence of the chain chirality in the multi-target mechanism of action. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4312-5 | 2.9 | 22 |
| 84 | A new EGFR inhibitor induces apoptosis in colon cancer cells. <i>Biochemical and Biophysical Research Communications</i> , 2007 , 354, 409-13 | 3.4 | 21 |
| 83 | Trypanocidal Activity of Quinoxaline 1,4 Di-N-oxide Derivatives as Trypanothione Reductase Inhibitors. <i>Molecules</i> , 2017 , 22, | 4.8 | 20 |

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| 82 | Synthetic polyamines: an overview of their multiple biological activities. <i>Amino Acids</i> , 2010 , 38, 383-92 | 3.5 | 20 |
| 81 | Polymethylene tetraamine backbone as template for the development of biologically active polyamines. <i>Medicinal Research Reviews</i> , 2003 , 23, 200-33 | 14.4 | 19 |
| 80 | Analysis of the muscarinic receptor subtype mediating inhibition of the neurogenic contractions in rabbit isolated vas deferens by a series of polymethylene tetra-amines. <i>British Journal of Pharmacology</i> , 2001 , 132, 1009-16 | 8.6 | 19 |
| 79 | Lewis Acid-Catalyzed Generation of C?C and C?N Bonds on EDeficient Heterocyclic Substrates. <i>Advanced Synthesis and Catalysis</i> , 2015 , 357, 185-195 | 5.6 | 18 |
| 78 | Sustainable Drug Discovery of Multi-Target-Directed Ligands for Alzheimer's Disease. <i>Journal of Medicinal Chemistry</i> , 2021 , 64, 4972-4990 | 8.3 | 18 |
| 77 | Repurposing of Drugs Targeting YAP-TEAD Functions. <i>Cancers</i> , 2018 , 10, | 6.6 | 18 |
| 76 | Crassiflorone derivatives that inhibit Trypanosoma brucei glyceraldehyde-3-phosphate dehydrogenase (TbGAPDH) and Trypanosoma cruzi trypanothione reductase (TcTR) and display trypanocidal activity. <i>European Journal of Medicinal Chemistry</i> , 2017 , 141, 138-148 | 6.8 | 17 |
| 75 | Quinones bearing non-steroidal anti-inflammatory fragments as multitarget ligands for Alzheimer's disease. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2013 , 23, 6254-8 | 2.9 | 17 |
| 74 | Search for selective antagonists at alpha 1-adrenoreceptors: neutral or negative antagonism?. <i>Il Farmaco</i> , 1998 , 53, 278-86 | | 17 |
| 73 | Heterocyclic inhibitors of AChE acylation and peripheral sites. <i>Il Farmaco</i> , 2005 , 60, 465-73 | | 17 |
| 72 | Stereocontrolled transformations of cyclohexadienone derivatives to access stereochemically rich and natural product-inspired architectures. <i>Organic and Biomolecular Chemistry</i> , 2020 , 18, 8526-8571 | 3.9 | 17 |
| 71 | N-1,2,3-triazole-isatin derivatives for cholinesterase and Eamyloid aggregation inhibition: A comprehensive bioassay study. <i>Bioorganic Chemistry</i> , 2020 , 98, 103753 | 5.1 | 16 |
| 70 | Histone deacetylases as targets for the treatment of neurodegenerative disorders: Challenges and future opportunities. <i>Medicinal Research Reviews</i> , 2020 , 40, 2177-2211 | 14.4 | 16 |
| 69 | Synthesis and evaluation of a library of 2,5-bisdiamino-benzoquinone derivatives as probes to modulate protein-protein interactions in prions. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2010 , 20, 1866-8 | 2.9 | 16 |
| 68 | Opioid antagonist activity of naltrexone-derived bivalent ligands: importance of a properly oriented molecular scaffold to guide "address" recognition at kappa opioid receptors. <i>Journal of Medicinal Chemistry</i> , 1996 , 39, 1816-22 | 8.3 | 15 |
| 67 | Discovery of Sustainable Drugs for Neglected Tropical Diseases: Cashew Nut Shell Liquid (CNSL)-Based Hybrids Target Mitochondrial Function and ATP Production in Trypanosoma brucei. <i>ChemMedChem</i> , 2019 , 14, 621-635 | 3.7 | 14 |
| 66 | Neuroregeneration versus neurodegeneration: toward a paradigm shift in Alzheimer's disease drug discovery. <i>Future Medicinal Chemistry</i> , 2017 , 9, 995-1013 | 4.1 | 14 |
| 65 | Progress in acetylcholinesterase inhibitors for Alzheimer's disease. <i>Expert Opinion on Therapeutic Patents</i> , 2006 , 16, 811-823 | 6.8 | 14 |

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| 64 | Structure-activity relationships of methoctramine-related polyamines as muscular nicotinic receptor noncompetitive antagonists. 2. Role of polymethylene chain lengths separating amine functions and of substituents on the terminal nitrogen atoms. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 1860-78 | 8.3 | 14 |
| 63 | A Focused Library of Psychotropic Analogues with Neuroprotective and Neuroregenerative Potential. <i>ACS Chemical Neuroscience</i> , 2019 , 10, 279-294 | 5.7 | 14 |
| 62 | Tacrine-O-protected phenolics heterodimers as multitarget-directed ligands against Alzheimer's disease: Selective subnanomolar BuChE inhibitors. <i>European Journal of Medicinal Chemistry</i> , 2019 , 181, 111550 | 6.8 | 13 |
| 61 | Molecular basis for covalent inhibition of glyceraldehyde-3-phosphate dehydrogenase by a 2-phenoxy-1,4-naphthoquinone small molecule. <i>Chemical Biology and Drug Design</i> , 2017 , 90, 225-235 | 2.9 | 12 |
| 60 | Solvent- and chromatography-free amination of deficient nitrogen heterocycles under microwave irradiation. A fast, efficient and green route to 9-aminoacridines, 4-aminoquinolines and 4-aminoquinazolines and its application to the synthesis of the drugs amsacrine and bistacrine. <i>Tetrahedron</i> , 2013 , 69, 1024-1030 | 2.4 | 12 |
| 59 | Multi-target-directed ligands as innovative tools to combat trypanosomatid diseases. <i>Current Topics in Medicinal Chemistry</i> , 2011 , 11, 2824-33 | 3 | 12 |
| 58 | Sequential virtual screening approach to the identification of small organic molecules as potential BACE-1 inhibitors. <i>Chemical Biology and Drug Design</i> , 2011 , 77, 268-71 | 2.9 | 12 |
| 57 | Analogues of prazosin that bear a benextramine-related polyamine backbone exhibit different antagonism toward alpha1-adrenoreceptor subtypes. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 362-71 | 8.3 | 12 |
| 56 | Quinone-amino acid conjugates targeting Leishmania amino acid transporters. <i>PLoS ONE</i> , 2014 , 9, e107994 | 3.7 | 12 |
| 55 | Development of a Focused Library of Triazole-Linked Privileged-Structure-Based Conjugates Leading to the Discovery of Novel Phenotypic Hits against Protozoan Parasitic Infections. <i>ChemMedChem</i> , 2018 , 13, 678-683 | 3.7 | 11 |
| 54 | Binding profile of benextramine at neuropeptide Y receptor subtypes in rat brain areas. <i>European Journal of Pharmacology</i> , 1994 , 265, 93-8 | 5.3 | 11 |
| 53 | Neurodegenerative drug discovery: building on the past, looking to the future. <i>Future Medicinal Chemistry</i> , 2017 , 9, 707-709 | 4.1 | 10 |
| 52 | Novel Sustainable-by-Design HDAC Inhibitors for the Treatment of Alzheimer's Disease. <i>ACS Medicinal Chemistry Letters</i> , 2019 , 10, 671-676 | 4.3 | 10 |
| 51 | Hybrid lipoic acid derivatives to attack prion disease on multiple fronts. <i>ChemMedChem</i> , 2011 , 6, 601-5 | 3.7 | 10 |
| 50 | Accelerating Drug Discovery Efforts for Trypanosomatid Infections Using an Integrated Transnational Academic Drug Discovery Platform. <i>SLAS Discovery</i> , 2019 , 24, 346-361 | 3.4 | 9 |
| 49 | Presynaptic M3 muscarinic cholinceptors mediate inhibition of excitatory synaptic transmission in area CA1 of rat hippocampus. <i>Brain Research</i> , 2015 , 1629, 260-9 | 3.7 | 9 |
| 48 | Structure-activity relationships of methoctramine-related polyamines as muscular nicotinic receptor noncompetitive antagonists. 3. Effect of inserting the tetraamine backbone into a macrocyclic structure. <i>Journal of Medicinal Chemistry</i> , 2002 , 45, 3286-95 | 8.3 | 9 |
| 47 | Design, synthesis, and biological activity of methoctramine-related polyamines as putative G(i) protein activators. <i>Journal of Medicinal Chemistry</i> , 2001 , 44, 4035-8 | 8.3 | 9 |

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| 46 | Modulation of prion by small molecules: from monovalent to bivalent and multivalent ligands. <i>Current Topics in Medicinal Chemistry</i> , 2013 , 13, 2491-503 | 3 | 9 |
| 45 | Structure-activity relationships and mechanistic studies of novel mitochondria-targeted, leishmanicidal derivatives of the 4-aminostyrylquinoline scaffold. <i>European Journal of Medicinal Chemistry</i> , 2019 , 171, 38-53 | 6.8 | 8 |
| 44 | Rational approach to an antiprion compound with a multiple mechanism of action. <i>Future Medicinal Chemistry</i> , 2015 , 7, 2113-20 | 4.1 | 8 |
| 43 | Novel multi target-directed ligands targeting 5-HT receptors with in cellulo antioxidant properties as promising leads in Alzheimer's disease. <i>European Journal of Medicinal Chemistry</i> , 2019 , 182, 111596 | 6.8 | 8 |
| 42 | Synthesis of new lipoic acid conjugates and evaluation of their free radical scavenging and neuroprotective activities. <i>Chemical Biology and Drug Design</i> , 2014 , 83, 688-96 | 2.9 | 8 |
| 41 | Synthesis, muscarinic blocking activity and molecular modeling studies of 4-DAMP-related compounds. <i>Bioorganic and Medicinal Chemistry</i> , 1995 , 3, 267-77 | 3.4 | 8 |
| 40 | Multitarget ligands and theranostics: sharpening the medicinal chemistry sword against prion diseases. <i>Future Medicinal Chemistry</i> , 2014 , 6, 1017-29 | 4.1 | 7 |
| 39 | Design, synthesis, and biological evaluation of substituted 2,3-dihydro-1H-cyclopenta[b]quinolin-9-ylamine related compounds as fructose-1,6-bisphosphatase inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2006 , 14, 7846-53 | 3.4 | 7 |
| 38 | Design, synthesis and biological evaluation of ambenonium derivatives as AChE inhibitors. <i>Il Farmaco</i> , 2003 , 58, 917-28 | | 7 |
| 37 | The Mu.Ta.Lig. Chemotheca: A Community-Populated Molecular Database for Multi-Target Ligands Identification and Compound-Repurposing. <i>Frontiers in Chemistry</i> , 2018 , 6, 130 | 5 | 6 |
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