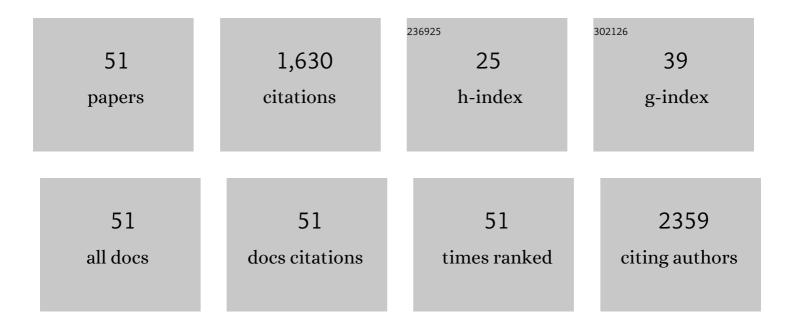
Alba EspargarÃ³

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Design, Synthesis, and In Vitro, In Silico and In Cellulo Evaluation of New Pyrimidine and Pyridine Amide and Carbamate Derivatives as Multi-Functional Cholinesterase Inhibitors. Pharmaceuticals, 2022, 15, 673.	3.8	3
2	Azobioisosteres of Curcumin with Pronounced Activity against Amyloid Aggregation, Intracellular Oxidative Stress, and Neuroinflammation. Chemistry - A European Journal, 2021, 27, 6015-6027.	3.3	4
3	Dual Inhibitors of Amyloid-β and Tau Aggregation with Amyloid-β Disaggregating Properties: Extended <i>In Cellulo</i> , <i>In Silico</i> , and Kinetic Studies of Multifunctional Anti-Alzheimer's Agents. ACS Chemical Neuroscience, 2021, 12, 2057-2068.	3.5	36
4	Dual Effect of Prussian Blue Nanoparticles on Aβ40 Aggregation: β-Sheet Fibril Reduction and Copper Dyshomeostasis Regulation. Biomacromolecules, 2021, 22, 430-440.	5.4	11
5	New Pyrimidine and Pyridine Derivatives as Multitarget Cholinesterase Inhibitors: Design, Synthesis, and <i>In Vitro</i> and <i>In Cellulo</i> Evaluation. ACS Chemical Neuroscience, 2021, 12, 4090-4112.	3.5	16
6	Centrally Active Multitarget Anti-Alzheimer Agents Derived from the Antioxidant Lead CR-6. Journal of Medicinal Chemistry, 2020, 63, 9360-9390.	6.4	25
7	Pharmacophore Modeling and 3D-QSAR Study of Indole and Isatin Derivatives as Antiamyloidogenic Agents Targeting Alzheimer's Disease. Molecules, 2020, 25, 5773.	3.8	9
8	Thiosemicarbazone Derivatives as Inhibitors of Amyloid-Î ² Aggregation: Effect of Metal Coordination. Inorganic Chemistry, 2020, 59, 6978-6987.	4.0	20
9	On the Binding of Congo Red to Amyloid Fibrils. Angewandte Chemie - International Edition, 2020, 59, 8104-8107.	13.8	36
10	On the Binding of Congo Red to Amyloid Fibrils. Angewandte Chemie, 2020, 132, 8181-8184.	2.0	11
11	A novel class of multitarget anti-Alzheimer benzohomoadamantane‒chlorotacrine hybrids modulating cholinesterases and glutamate NMDA receptors. European Journal of Medicinal Chemistry, 2019, 180, 613-626.	5.5	26
12	Synthesis, In Vitro Profiling, and In Vivo Efficacy Studies of a New Family of Multitarget Anti-Alzheimer Compounds. Proceedings (mdpi), 2019, 22, .	0.2	0
13	Amyloid Pan-inhibitors: One Family of Compounds To Cope with All Conformational Diseases. ACS Chemical Neuroscience, 2019, 10, 1311-1317.	3.5	14
14	Bacterial Inclusion Bodies for Anti-Amyloid Drug Discovery: Current and Future Screening Methods. Current Protein and Peptide Science, 2019, 20, 563-576.	1.4	7
15	Combined in Vitro Cell-Based/in Silico Screening of Naturally Occurring Flavonoids and Phenolic Compounds as Potential Anti-Alzheimer Drugs. Journal of Natural Products, 2017, 80, 278-289.	3.0	68
16	Design, synthesis and multitarget biological profiling of second-generation anti-Alzheimer rhein–huprine hybrids. Future Medicinal Chemistry, 2017, 9, 965-981.	2.3	40
17	Evidence of Protein Adsorption in Pegylated Liposomes: Influence of Liposomal Decoration. Nanomaterials, 2017, 7, 37.	4.1	19
18	Key Points Concerning Amyloid Infectivity and Prion-Like Neuronal Invasion. Frontiers in Molecular Neuroscience, 2016, 9, 29,	2.9	19

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19	Histidineâ€Rich Oligopeptides To Lessen Copperâ€Mediated Amyloidâ€Î² Toxicity. Chemistry - A European Journal, 2016, 22, 7268-7280.	3.3	25
20	Investigation into the stability and reactivity of the pentacyclic alkaloid dehydroevodiamine and the benz-analog thereof. Tetrahedron, 2016, 72, 2535-2543.	1.9	9
21	Natural Xanthones from Garcinia mangostana with Multifunctional Activities for the Therapy of Alzheimer's Disease. Neurochemical Research, 2016, 41, 1806-1817.	3.3	59
22	In vivo amyloid aggregation kinetics tracked by timeâ€lapse confocal microscopy in realâ€ŧime. Biotechnology Journal, 2016, 11, 172-177.	3.5	14
23	Ultra rapid in vivo screening for anti-Alzheimer anti-amyloid drugs. Scientific Reports, 2016, 6, 23349.	3.3	37
24	Amyloids in solid-state nuclear magnetic resonance: potential causes of the usually low resolution. International Journal of Nanomedicine, 2015, 10, 6975.	6.7	5
25	Magnetic Nanoparticles Cross the Blood-Brain Barrier: When Physics Rises to a Challenge. Nanomaterials, 2015, 5, 2231-2248.	4.1	67
26	Could <i>α</i> -Synuclein Amyloid-Like Aggregates Trigger a Prionic Neuronal Invasion?. BioMed Research International, 2015, 2015, 1-7.	1.9	10
27	Novel Levetiracetam Derivatives That Are Effective against the Alzheimer-like Phenotype in Mice: Synthesis, in Vitro, ex Vivo, and in Vivo Efficacy Studies. Journal of Medicinal Chemistry, 2015, 58, 6018-6032.	6.4	58
28	Predicting the aggregation propensity of prion sequences. Virus Research, 2015, 207, 127-135.	2.2	7
29	Shogaol–huprine hybrids: Dual antioxidant and anticholinesterase agents with β-amyloid and tau anti-aggregating properties. Bioorganic and Medicinal Chemistry, 2014, 22, 5298-5307.	3.0	37
30	Tetrahydrobenzo[h][1,6]naphthyridine-6-chlorotacrine hybrids as a new family of anti-Alzheimer agents targeting l²-amyloid, tau, and cholinesterase pathologies. European Journal of Medicinal Chemistry, 2014, 84, 107-117.	5.5	57
31	Thioflavin-S Staining of Bacterial Inclusion Bodies for the Fast, Simple, and Inexpensive Screening of Amyloid Aggregation Inhibitors. Current Medicinal Chemistry, 2014, 21, 1152-1159.	2.4	44
32	Screening for Amyloid Aggregation: In-Silico, In-Vitro and In-Vivo Detection. Current Protein and Peptide Science, 2014, 15, 477-489.	1.4	9
33	Thioflavin-S staining coupled to flow cytometry. A screening tool to detect in vivo protein aggregation. Molecular BioSystems, 2012, 8, 2839.	2.9	47
34	Discovery of Novel Inhibitors of Amyloid β-Peptide 1–42 Aggregation. Journal of Medicinal Chemistry, 2012, 55, 9521-9530.	6.4	39
35	Using bacterial inclusion bodies to screen for amyloid aggregation inhibitors. Microbial Cell Factories, 2012, 11, 55.	4.0	33
36	Yeast prions form infectious amyloid inclusion bodies in bacteria. Microbial Cell Factories, 2012, 11, 89.	4.0	26

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#	Article	IF	CITATIONS
37	Temperature Dependence of the Aggregation Kinetics of Sup35 and Ure2p Yeast Prions. Biomacromolecules, 2012, 13, 474-483.	5.4	18
38	Native Structure Protects SUMO Proteins from Aggregation into Amyloid Fibrils. Biomacromolecules, 2012, 13, 1916-1926.	5.4	28
39	Effect of the surface charge of artificial model membranes on the aggregation of amyloid β-peptide. Biochimie, 2012, 94, 1730-1738.	2.6	40
40	Aggregation of the neuroblastoma-associated mutant (S120G) of the human nucleoside diphosphate kinase-A/NM23-H1 into amyloid fibrils. Naunyn-Schmiedeberg's Archives of Pharmacology, 2011, 384, 373-381.	3.0	5
41	Bacterial Inclusion Bodies of Alzheimer's Disease βâ€Amyloid Peptides Can Be Employed To Study Native‣ike Aggregation Intermediate States. ChemBioChem, 2011, 12, 407-423.	2.6	90
42	Deciphering the role of the thermodynamic and kinetic stabilities of SH3 domains on their aggregation inside bacteria. Proteomics, 2010, 10, 4172-4185.	2.2	23
43	The Role of Protein Sequence and Amino Acid Composition in Amyloid Formation: Scrambling and Backward Reading of IAPP Amyloid Fibrils. Journal of Molecular Biology, 2010, 404, 337-352.	4.2	38
44	Energy barriers for HETâ€s prion forming domain amyloid formation. FEBS Journal, 2009, 276, 5053-5064.	4.7	23
45	Characterization of the amyloid bacterial inclusion bodies of the HET-s fungal prion. Microbial Cell Factories, 2009, 8, 56.	4.0	37
46	Study and selection of in vivo protein interactions by coupling bimolecular fluorescence complementation and flow cytometry. Nature Protocols, 2008, 3, 22-33.	12.0	51
47	Kinetic and thermodynamic stability of bacterial intracellular aggregates. FEBS Letters, 2008, 582, 3669-3673.	2.8	24
48	Inclusion bodies: Specificity in their aggregation process and amyloid-like structure. Biochimica Et Biophysica Acta - Molecular Cell Research, 2008, 1783, 1815-1825.	4.1	131
49	Studies on bacterial inclusion bodies. Future Microbiology, 2008, 3, 423-435.	2.0	34
50	The in Vivo and in Vitro Aggregation Properties of Globular Proteins Correlate With Their Conformational Stability: The SH3 Case. Journal of Molecular Biology, 2008, 378, 1116-1131.	4.2	56
51	Detection of transient protein–protein interactions by bimolecular fluorescence complementation: The Abl-SH3 case. Proteomics, 2007, 7, 1023-1036.	2.2	85