

Stephen Connelly

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

2,096
citations

471509

17
h-index

677142

22
g-index

23
all docs

23
docs citations

23
times ranked

2567
citing authors

#	ARTICLE	IF	CITATIONS
1	Semi-quantitative models for identifying potent and selective transthyretin amyloidogenesis inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3441-3449.	2.2	8
2	Stabilizing the C _H 2 Domain of an Antibody by Engineering in an Enhanced Aromatic Sequon. <i>ACS Chemical Biology</i> , 2016, 11, 1852-1861.	3.4	40
3	A Fluorogenic Aryl Fluorosulfate for Intraorganellar Transthyretin Imaging in Living Cells and in <i>Caenorhabditis elegans</i> . <i>Journal of the American Chemical Society</i> , 2015, 137, 7404-7414.	13.7	86
4	Iridium-Catalysed C-H Borylation Facilitates a Total Synthesis of the HRV 3C Protease Inhibitor (±)-Thysanone. <i>Synlett</i> , 2014, 25, 556-558.	1.8	7
5	Fluorogenic small molecules requiring reaction with a specific protein to create a fluorescent conjugate for biological imaging—what we know and what we need to learn. <i>Biopolymers</i> , 2014, 101, 484-495.	2.4	8
6	Akt Phosphorylation and Regulation of Transketolase Is a Nodal Point for Amino Acid Control of Purine Synthesis. <i>Molecular Cell</i> , 2014, 55, 264-276.	9.7	70
7	Synthesis and Biological Evaluation of Deoxy Analogues of the Human Rhinovirus 3C Protease Inhibitor Thysanone. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 122-128.	2.4	9
8	Bifunctional coumarin derivatives that inhibit transthyretin amyloidogenesis and serve as fluorescent transthyretin folding sensors. <i>Chemical Communications</i> , 2013, 49, 9188.	4.1	35
9	Aromatic Sulfonyl Fluorides Covalently Kinetically Stabilize Transthyretin to Prevent Amyloidogenesis while Affording a Fluorescent Conjugate. <i>Journal of the American Chemical Society</i> , 2013, 135, 5656-5668.	13.7	142
10	Stilbene Vinyl Sulfonamides as Fluorogenic Sensors of and Traceless Covalent Kinetic Stabilizers of Transthyretin That Prevent Amyloidogenesis. <i>Journal of the American Chemical Society</i> , 2013, 135, 17869-17880.	13.7	33
11	Biological and Structural Evaluation of 10 ^R - and 10 ^S -Methylthio-DDACTHF Reveals a New Role for Sulfur in Inhibition of Glycinamide Ribonucleotide Transformylase. <i>Biochemistry</i> , 2013, 52, 5133-5144.	2.5	7
12	AG10 inhibits amyloidogenesis and cellular toxicity of the familial amyloid cardiomyopathy-associated V122I transthyretin. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 9992-9997.	7.1	120
13	The Transthyretin Amyloidoses: From Delineating the Molecular Mechanism of Aggregation Linked to Pathology to a Regulatory-Agency-Approved Drug. <i>Journal of Molecular Biology</i> , 2012, 421, 185-203.	4.2	267
14	Tafamidis, a potent and selective transthyretin kinetic stabilizer that inhibits the amyloid cascade. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 9629-9634.	7.1	582
15	Potent Kinetic Stabilizers That Prevent Transthyretin-Mediated Cardiomyocyte Proteotoxicity. <i>Science Translational Medicine</i> , 2011, 3, 97ra81.	12.4	61
16	Structure-based design of kinetic stabilizers that ameliorate the transthyretin amyloidoses. <i>Current Opinion in Structural Biology</i> , 2010, 20, 54-62.	5.7	160
17	Chemoselective small molecules that covalently modify one lysine in a non-enzyme protein in plasma. <i>Nature Chemical Biology</i> , 2010, 6, 133-139.	8.0	74
18	A Substructure Combination Strategy To Create Potent and Selective Transthyretin Kinetic Stabilizers That Prevent Amyloidogenesis and Cytotoxicity. <i>Journal of the American Chemical Society</i> , 2010, 132, 1359-1370.	13.7	67

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19	The Phosphatidylinositol 3-Kinase/Akt Cassette Regulates Purine Nucleotide Synthesis. <i>Journal of Biological Chemistry</i> , 2009, 284, 3521-3528.	3.4	53
20	Toward Optimization of the Second Aryl Substructure Common to Transthyretin Amyloidogenesis Inhibitors Using Biochemical and Structural Studies. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 1115-1125.	6.4	66
21	Biochemical and Structural Evaluation of Highly Selective 2-Arylbenzoxazole-Based Transthyretin Amyloidogenesis Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 260-270.	6.4	127
22	Toward Optimization of the Linker Substructure Common to Transthyretin Amyloidogenesis Inhibitors Using Biochemical and Structural Studies. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 6348-6358.	6.4	73