

Barbara Campanini

List of Publications by Year in descending order

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81
papers

1,722
citations

236925

25
h-index

330143

37
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83
all docs

83
docs citations

83
times ranked

1599
citing authors

#	ARTICLE	IF	CITATIONS
1	Interaction of serine acetyltransferase with O-acetylserine sulfhydrylase active site: Evidence from fluorescence spectroscopy. <i>Protein Science</i> , 2005, 14, 2115-2124.	7.6	83
2	Design of O-Acetylserine Sulfhydrylase Inhibitors by Mimicking Nature. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 345-356.	6.4	75
3	Dynamics of green fluorescent protein mutant2 in solution, on spin-coated glasses, and encapsulated in wet silica gels. <i>Protein Science</i> , 2002, 11, 1152-1161.	7.6	61
4	Single Amino Acid Replacement Makes <i>Aequorea victoria</i> Fluorescent Proteins Reversibly Photoswitchable. <i>Journal of the American Chemical Society</i> , 2010, 132, 85-95.	13.7	61
5	Green Fluorescent Protein Ground States: The Influence of a Second Protonation Site near the Chromophore. <i>Biochemistry</i> , 2007, 46, 5494-5504.	2.5	60
6	Structure, Mechanism, and Conformational Dynamics of O-Acetylserine Sulfhydrylase from <i>Salmonella typhimurium</i> : Comparison of A and B Isozymes. <i>Biochemistry</i> , 2007, 46, 8315-8330.	2.5	58
7	Kinetics of Acid-Induced Spectral Changes in the GFPmut2 Chromophore. <i>Journal of the American Chemical Society</i> , 2005, 127, 626-635.	13.7	57
8	Unfolding of Green Fluorescent Protein mut2 in wet nanoporous silica gels. <i>Protein Science</i> , 2005, 14, 1125-1133.	7.6	57
9	Isozyme-Specific Ligands for O-acetylserine sulfhydrylase, a Novel Antibiotic Target. <i>PLoS ONE</i> , 2013, 8, e77558.	2.5	43
10	Inhibitors of the Sulfur Assimilation Pathway in Bacterial Pathogens as Enhancers of Antibiotic Therapy. <i>Current Medicinal Chemistry</i> , 2014, 22, 187-213.	2.4	42
11	Iron Metabolism at the Interface between Host and Pathogen: From Nutritional Immunity to Antibacterial Development. <i>International Journal of Molecular Sciences</i> , 2020, 21, 2145.	4.1	42
12	The multifaceted pyridoxal 5-phosphate-dependent O-acetylserine sulfhydrylase. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2011, 1814, 1497-1510.	2.3	39
13	Design and synthesis of trans-2-substituted-cyclopropane-1-carboxylic acids as the first non-natural small molecule inhibitors of O-acetylserine sulfhydrylase. <i>MedChemComm</i> , 2012, 3, 1111.	3.4	36
14	Role of Pyridoxal 5-Phosphate in the Structural Stabilization of O-Acetylserine Sulfhydrylase. <i>Journal of Biological Chemistry</i> , 2000, 275, 40244-40251.	3.4	35
15	A Two-step Process Controls the Formation of the Bifunctional Cysteine Synthase Complex. <i>Journal of Biological Chemistry</i> , 2010, 285, 12813-12822.	3.4	35
16	Fine tuning of the active site modulates specificity in the interaction of O-acetylserine sulfhydrylase isozymes with serine acetyltransferase. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2013, 1834, 169-181.	2.3	35
17	Moonlighting O-acetylserine sulfhydrylase: New functions for an old protein. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2015, 1854, 1184-1193.	2.3	35
18	Singlet oxygen photosensitisation by GFP mutants: oxygen accessibility to the chromophore. <i>Photochemical and Photobiological Sciences</i> , 2010, 9, 1336-1341.	2.9	34

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19	Serine racemase: a key player in neuron activity and in neuropathologies. <i>Frontiers in Bioscience - Landmark</i> , 2013, 18, 1112.	3.0	34
20	<sc>ATP</sc> binding to human serine racemase is cooperative and modulated by glycine. <i>FEBS Journal</i> , 2013, 280, 5853-5863.	4.7	33
21	A novel Bim-BH3-derived Bcl-XL inhibitor: Biochemical characterization, in vitro, in vivo and ex-vivo anti-leukemic activity. <i>Cell Cycle</i> , 2008, 7, 3211-3224.	2.6	32
22	Tracking Unfolding and Refolding of Single GFPmut2 Molecules. <i>Biophysical Journal</i> , 2005, 89, 2033-2045.	0.5	31
23	Rational Design, Synthesis, and Preliminary Structure-Activity Relationships of \pm -Substituted-2-Phenylcyclopropane Carboxylic Acids as Inhibitors of <i>Salmonella typhimurium</i> O-Acetylserine Sulfhydrylase. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 2567-2578.	6.4	28
24	The Energy Landscape of Human Serine Racemase. <i>Frontiers in Molecular Biosciences</i> , 2018, 5, 112.	3.5	28
25	Protonation and Conformational Dynamics of GFP Mutants by Two-Photon Excitation Fluorescence Correlation Spectroscopy. <i>Journal of Physical Chemistry B</i> , 2008, 112, 8806-8814.	2.6	25
26	Identification of the Structural Determinants for the Stability of Substrate and Aminoacrylate External Schiff Bases in O-Acetylserine Sulfhydrylase-A. <i>Biochemistry</i> , 2010, 49, 6093-6103.	2.5	25
27	Human kynurenine aminotransferase reactivity with substrates and inhibitors. <i>FEBS Journal</i> , 2011, 278, 1882-1900.	4.7	25
28	Surface-exposed Tryptophan Residues Are Essential for O-Acetylserine Sulfhydrylase Structure, Function, and Stability. <i>Journal of Biological Chemistry</i> , 2003, 278, 37511-37519.	3.4	24
29	Expanding the chemical space of human serine racemase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 4297-4303.	2.2	22
30	Regulation of human serine racemase activity and dynamics by halides, ATP and malonate. <i>Amino Acids</i> , 2015, 47, 163-173.	2.7	21
31	Cyclopropane-1,2-dicarboxylic acids as new tools for the biophysical investigation of O-acetylserine sulfhydrylases by fluorimetric methods and saturation transfer difference (STD) NMR. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 78-87.	5.2	21
32	Interaction of human hemoglobin and semi-hemoglobins with the <i>Staphylococcus aureus</i> hemophore IsdB: a kinetic and mechanistic insight. <i>Scientific Reports</i> , 2019, 9, 18629.	3.3	21
33	Immobilization of Proteins in Silica Gel: Biochemical and Biophysical Properties. <i>Current Organic Chemistry</i> , 2015, 19, 1653-1668.	1.6	20
34	Study of DNA binding and bending by <i>Bacillus subtilis</i> GabR, a PLP-dependent transcription factor. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2017, 1861, 3474-3489.	2.4	18
35	Magnesium and calcium ions differentially affect human serine racemase activity and modulate its quaternary equilibrium toward a tetrameric form. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2017, 1865, 381-387.	2.3	17
36	Discovery of novel fragments inhibiting O-acetylserine sulphhydrylase by combining scaffold hopping and ligand-based drug design. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018, 33, 1444-1452.	5.2	17

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37	Inhibition of Nonessential Bacterial Targets: Discovery of a Novel Serine <i>O</i> -Acetyltransferase Inhibitor. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 790-797.	2.8	17
38	Sulfur Mobilization in Cyanobacteria. <i>Journal of Biological Chemistry</i> , 2006, 281, 38769-38780.	3.4	16
39	Evidence of Discrete Substates and Unfolding Pathways in Green Fluorescent Protein. <i>Biophysical Journal</i> , 2007, 92, 1724-1731.	0.5	16
40	Use of Exogenous Enzymes in Human Therapy: Approved Drugs and Potential Applications. <i>Current Medicinal Chemistry</i> , 2022, 29, 411-452.	2.4	16
41	Asymmetry of the Active Site Loop Conformation between Subunits of Glutamate-1-semialdehyde Aminomutase in Solution. <i>BioMed Research International</i> , 2013, 2013, 1-10.	1.9	15
42	Modulation of <i>Escherichia coli</i> serine acetyltransferase catalytic activity in the cysteine synthase complex. <i>FEBS Letters</i> , 2017, 591, 1212-1224.	2.8	15
43	Integration of Enhanced Sampling Methods with Saturation Transfer Difference Experiments to Identify Protein Druggable Pockets. <i>Journal of Chemical Information and Modeling</i> , 2018, 58, 710-723.	5.4	15
44	Stimulated Emission Properties of Fluorophores by CW-STED Single Molecule Spectroscopy. <i>Journal of Physical Chemistry B</i> , 2013, 117, 16405-16415.	2.6	14
45	Investigational Studies on a Hit Compound Cyclopropane- α -Carboxylic Acid Derivative Targeting <i>O</i> -Acetylserine Sulfhydrylase as a Colistin Adjuvant. <i>ACS Infectious Diseases</i> , 2021, 7, 281-292.	3.8	13
46	Pyridoxal 5 α -Phosphate-Dependent Enzymes: Catalysis, Conformation, and Genomics. , 2010, , 273-350.		12
47	Photoinduced Millisecond Switching Kinetics in the GFPmut2 E222Q Mutant. <i>Journal of Physical Chemistry B</i> , 2010, 114, 4664-4677.	2.6	12
48	Structure and single crystal spectroscopy of Green Fluorescent Proteins. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2011, 1814, 824-833.	2.3	12
49	Cyclopropane derivatives as potential human serine racemase inhibitors: unveiling novel insights into a difficult target. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2016, 31, 645-652.	5.2	12
50	Inhibition of <i>O</i> -acetylserine sulfhydrylase by fluoroalanine derivatives. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2018, 33, 1343-1351.	5.2	12
51	Glutamine 89 is a key residue in the allosteric modulation of human serine racemase activity by ATP. <i>Scientific Reports</i> , 2018, 8, 9016.	3.3	12
52	More than a Confinement: α -Soft and α -Hard-Enzyme Entrapment Modulates Biological Catalyst Function. <i>Catalysts</i> , 2019, 9, 1024.	3.5	12
53	Refining the structure-activity relationships of 2-phenylcyclopropane carboxylic acids as inhibitors of <i>O</i> -acetylserine sulfhydrylase isoforms. <i>Journal of Enzyme Inhibition and Medicinal Chemistry</i> , 2019, 34, 31-43.	5.2	12
54	Human serine racemase is allosterically modulated by NADH and reduced nicotinamide derivatives. <i>Biochemical Journal</i> , 2016, 473, 3505-3516.	3.7	11

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55	Human serine racemase is nitrosylated at multiple sites. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2018, 1866, 813-821.	2.3	11
56	GFP-mut2 Proteins in Trehalose-Water Matrixes: Spatially Heterogeneous Protein-Water-Sugar Structures. <i>Biophysical Journal</i> , 2007, 93, 284-293.	0.5	10
57	Role of histidine 148 in stability and dynamics of a highly fluorescent GFP variant. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2013, 1834, 770-779.	2.3	10
58	Structural insight into the interaction of <i>O</i> -acetylserine sulfhydrylase with competitive, peptidic inhibitors by saturation transfer difference NMR. <i>FEBS Letters</i> , 2016, 590, 943-953.	2.8	10
59	Unfolding of pyridoxal 5-phosphate-dependent <i>O</i> -acetylserine sulfhydrylase probed by time-resolved tryptophan fluorescence. <i>BBA - Proteins and Proteomics</i> , 2002, 1596, 47-54.	2.1	9
60	Enhanced Green Fluorescent Protein (GFP) fluorescence after polyelectrolyte caging. <i>Optics Express</i> , 2006, 14, 9815.	3.4	9
61	Combination of SAXS and Protein Painting Discloses the Three-Dimensional Organization of the Bacterial Cysteine Synthase Complex, a Potential Target for Enhancers of Antibiotic Action. <i>International Journal of Molecular Sciences</i> , 2019, 20, 5219.	4.1	9
62	Exploring <i>O</i> -acetylserine sulfhydrylase-B isoenzyme from <i>Salmonella typhimurium</i> by fluorescence spectroscopy. <i>Archives of Biochemistry and Biophysics</i> , 2011, 505, 178-185.	3.0	8
63	The allosteric interplay between nitrosylation and glycine binding controls the activity of human serine racemase. <i>FEBS Journal</i> , 2021, 288, 3034-3054.	4.7	8
64	Activation of an anti-bacterial toxin by the biosynthetic enzyme CysK: mechanism of binding, interaction specificity and competition with cysteine synthase. <i>Scientific Reports</i> , 2017, 7, 8817.	3.3	7
65	Voltage regulation of single green fluorescent protein mutants. <i>Biophysical Chemistry</i> , 2007, 125, 368-374.	2.8	6
66	Cryo-EM structures of staphylococcal IsdB bound to human hemoglobin reveal the process of heme extraction. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2116708119.	7.1	6
67	Environment effects on the oscillatory unfolding kinetics of GFP. <i>European Biophysics Journal</i> , 2007, 36, 795-803.	2.2	5
68	Off to a slow start: Analyzing lag phases and accelerating rates in steady-state enzyme kinetics. <i>Analytical Biochemistry</i> , 2020, 593, 113595.	2.4	5
69	Discovery of Substituted (2-Aminooxazol-4-yl)isoxazole-3-carboxylic Acids as Inhibitors of Bacterial Serine Acetyltransferase in the Quest for Novel Potential Antibacterial Adjuvants. <i>Pharmaceuticals</i> , 2021, 14, 174.	3.8	5
70	A Novel Assay for Phosphoserine Phosphatase Exploiting Serine Acetyltransferase as the Coupling Enzyme. <i>Life</i> , 2021, 11, 485.	2.4	5
71	Structural stability of green fluorescent proteins entrapped in polyelectrolyte nanocapsules. <i>Journal of Biophotonics</i> , 2008, 1, 310-319.	2.3	4
72	A Competitive <i>O</i> -Acetylserine Sulfhydrylase Inhibitor Modulates the Formation of Cysteine Synthase Complex. <i>Catalysts</i> , 2021, 11, 700.	3.5	4

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73	Effect of the point mutation H148G on GFPmut2 unfolding kinetics by fluorescence spectroscopy. <i>Biophysical Chemistry</i> , 2011, 157, 24-32.	2.8	3
74	Insight into GFPmut2 pH Dependence by Single Crystal Microspectrophotometry and X-ray Crystallography. <i>Journal of Physical Chemistry B</i> , 2018, 122, 11326-11337.	2.6	3
75	Birth of a pathway for sulfur metabolism in early amniote evolution. <i>Nature Ecology and Evolution</i> , 2020, 4, 1239-1246.	7.8	3
76	Human serine racemase is inhibited by glyceraldehyde 3-phosphate, but not by glyceraldehyde 3-phosphate dehydrogenase. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2021, 1869, 140544.	2.3	3
77	Revealing the Dynamic Allosteric Changes Required for Formation of the Cysteine Synthase Complex by Hydrogen-Deuterium Exchange MS. <i>Molecular and Cellular Proteomics</i> , 2021, 20, 100098.	3.8	1
78	Human Serine Racemase Weakly Binds the Third PDZ Domain of PSD-95. <i>International Journal of Molecular Sciences</i> , 2022, 23, 4959.	4.1	1
79	Inhibitors of O-Acetylserine Sulfhydrylase with a Cyclopropane-Carboxylic Acid Scaffold Are Effective Colistin Adjuvants in Gram Negative Bacteria. <i>Pharmaceuticals</i> , 2022, 15, 766.	3.8	1
80	Green Fluorescent Protein Photodynamics as a Tool for Fluorescence Correlative Studies and Applications. <i>Springer Series on Fluorescence</i> , 2011, , 35-55.	0.8	0
81	Exploring the chemical space around N-(5-nitrothiazol-2-yl)-1,2,3-thiadiazole-4-carboxamide, a hit compound with serine acetyltransferase (SAT) inhibitory properties. <i>Results in Chemistry</i> , 2022, 4, 100443.	2.0	0