

Martino Calamai

List of Publications by Year in descending order

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34
papers

1,514
citations

471509

17
h-index

454955

30
g-index

38
all docs

38
docs citations

38
times ranked

2274
citing authors

#	ARTICLE	IF	CITATIONS
1	Membrane Phase Drives the Assembly of Gold Nanoparticles on Biomimetic Lipid Bilayers. <i>Journal of Physical Chemistry C</i> , 2022, 126, 4483-4494.	3.1	15
2	3D Printing Silk-Based Bioresorbable Piezoelectric Self-Adhesive Holey Structures for <i>In Vivo</i> Monitoring on Soft Tissues. <i>ACS Applied Materials & Interfaces</i> , 2022, 14, 19253-19264.	8.0	15
3	Full-length TDP-43 and its C-terminal domain form filaments <i>in vitro</i> having non-amyloid properties. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2021, 28, 56-65.	3.0	6
4	Gold Nanostars Bioconjugation for Selective Targeting and SERS Detection of Biofluids. <i>Nanomaterials</i> , 2021, 11, 665.	4.1	11
5	Carbon Nanotubes/Regenerated Silk Composite as a Three-Dimensional Printable Bio-Adhesive Ink with Self-Powering Properties. <i>ACS Applied Materials & Interfaces</i> , 2021, 13, 21007-21017.	8.0	17
6	Quantitative Measurement of the Affinity of Toxic and Nontoxic Misfolded Protein Oligomers for Lipid Bilayers and of its Modulation by Lipid Composition and Trodusquemine. <i>ACS Chemical Neuroscience</i> , 2021, 12, 3189-3202.	3.5	13
7	Plasma Membrane Dynamics and Proteolytic Processing of APP from a Single Molecule/Single Cell Perspective. <i>Biophysical Journal</i> , 2020, 118, 454a.	0.5	0
8	Making biological membrane resistant to the toxicity of misfolded protein oligomers: a lesson from trodusquemine. <i>Nanoscale</i> , 2020, 12, 22596-22614.	5.6	16
9	Design of Biocompatible Liquid Cristal Elastomers Reproducing the Mechanical Properties of Human Cardiac Muscle. <i>Biophysical Journal</i> , 2019, 116, 264a.	0.5	0
10	Development of Light-Responsive Liquid Crystalline Elastomers to Assist Cardiac Contraction. <i>Circulation Research</i> , 2019, 124, e44-e54.	4.5	44
11	Pre-diagnosing and managing patients with GM1 gangliosidosis and related disorders by the evaluation of GM1 ganglioside content. <i>Scientific Reports</i> , 2019, 9, 17684.	3.3	11
12	Three-Dimensional Tracking of Quantum Dot-Conjugated Molecules in Living Cells. <i>Methods in Molecular Biology</i> , 2018, 1814, 425-448.	0.9	9
13	Quantifying the Proteolytic Cleavage of Plasma Membrane Proteins in Living Cells. <i>Current Protocols in Cell Biology</i> , 2018, 81, e58.	2.3	1
14	Direct imaging of APP proteolysis in living cells. <i>PeerJ</i> , 2017, 5, e3086.	2.0	7
15	Single molecule experiments emphasize GM1 as a key player of the different cytotoxicity of structurally distinct A β 42 oligomers. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2016, 1858, 386-392.	2.6	22
16	Molecular insights into cell toxicity of a novel familial amyloidogenic variant of β 2-microglobulin. <i>Journal of Cellular and Molecular Medicine</i> , 2016, 20, 1443-1456.	3.6	23
17	GM1 and GM2 gangliosides: recent developments. <i>Biomolecular Concepts</i> , 2014, 5, 87-93.	2.2	22
18	Partitioning and confinement of GM1 ganglioside induced by amyloid aggregates. <i>FEBS Letters</i> , 2013, 587, 1385-1391.	2.8	38

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19	IDOL Stimulates Clathrin-Independent Endocytosis and Multivesicular Body-Mediated Lysosomal Degradation of the Low-Density Lipoprotein Receptor. <i>Molecular and Cellular Biology</i> , 2013, 33, 1503-1514.	2.3	68
20	Decoupling Polarization of the Golgi Apparatus and GM1 in the Plasma Membrane. <i>PLoS ONE</i> , 2013, 8, e80446.	2.5	15
21	Toxic effects of amyloid fibrils on cell membranes: the importance of ganglioside GM1. <i>FASEB Journal</i> , 2012, 26, 818-831.	0.5	118
22	Amyloid Aggregates Alter the Membrane Mobility of GM1 Gangliosides. <i>Biophysical Journal</i> , 2011, 100, 539a.	0.5	0
23	Single Molecule Tracking Analysis Reveals That the Surface Mobility of Amyloid Oligomers Is Driven by Their Conformational Structure. <i>Journal of the American Chemical Society</i> , 2011, 133, 12001-12008.	13.7	32
24	Formation and Stability of Synaptic Receptor Domains. <i>Physical Review Letters</i> , 2011, 106, 238104.	7.8	35
25	Single particle tracking of amyloid oligomers on the plasma membrane of living cells. , 2011, , .		0
26	Mutational Analysis of the Aggregation-Prone and Disaggregation-Prone Regions of Acylphosphatase. <i>Journal of Molecular Biology</i> , 2009, 387, 965-974.	4.2	12
27	Gephyrin Oligomerization Controls GlyR Mobility and Synaptic Clustering. <i>Journal of Neuroscience</i> , 2009, 29, 7639-7648.	3.6	78
28	Nature and Significance of the Interactions between Amyloid Fibrils and Biological Polyelectrolytes. <i>Biochemistry</i> , 2006, 45, 12806-12815.	2.5	128
29	Reversal of Protein Aggregation Provides Evidence for Multiple Aggregated States. <i>Journal of Molecular Biology</i> , 2005, 346, 603-616.	4.2	86
30	Evidence for a Mechanism of Amyloid Formation Involving Molecular Reorganisation within Native-like Precursor Aggregates. <i>Journal of Molecular Biology</i> , 2005, 351, 910-922.	4.2	129
31	Amyloid Fibril Formation Can Proceed from Different Conformations of a Partially Unfolded Protein. <i>Biophysical Journal</i> , 2005, 89, 4201-4210.	0.5	141
32	Studying the Folding Process of the Acylphosphatase from <i>Sulfolobus solfataricus</i> . A Comparative Analysis with Other Proteins from the Same Superfamily. <i>Biochemistry</i> , 2004, 43, 9116-9126.	2.5	19
33	Relative Influence of Hydrophobicity and Net Charge in the Aggregation of Two Homologous Proteins. <i>Biochemistry</i> , 2003, 42, 15078-15083.	2.5	115
34	Studies of the aggregation of mutant proteins in vitro provide insights into the genetics of amyloid diseases. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002, 99, 16419-16426.	7.1	268