Sofia Giorgetti

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Hereditary Systemic Amyloidosis Due to Asp76Asn Variant β ₂ -Microglobulin. New England Journal of Medicine, 2012, 366, 2276-2283.	27.0	172
2	The solution structure of human β2â€microglobulin reveals the prodromes of its amyloid transition. Protein Science, 2002, 11, 487-499.	7.6	145
3	A Partially Structured Species of β2-Microglobulin Is Significantly Populated under Physiological Conditions and Involved in Fibrillogenesis. Journal of Biological Chemistry, 2001, 276, 46714-46721.	3.4	137
4	Collagen Plays an Active Role in the Aggregation of β2-Microglobulin under Physiopathological Conditions of Dialysis-related Amyloidosis*. Journal of Biological Chemistry, 2006, 281, 16521-16529.	3.4	128
5	Structure, function and amyloidogenic propensity of apolipoprotein A-I. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2006, 13, 191-205.	3.0	124
6	Detection of two partially structured species in the folding process of the amyloidogenic protein β2-microglobulin. Journal of Molecular Biology, 2001, 307, 379-391.	4.2	115
7	A novel mechanoâ€enzymatic cleavage mechanism underlies transthyretin amyloidogenesis. EMBO Molecular Medicine, 2015, 7, 1337-1349.	6.9	109
8	Heparin Strongly Enhances the Formation of β2-Microglobulin Amyloid Fibrils in the Presence of Type I Collagen. Journal of Biological Chemistry, 2008, 283, 4912-4920.	3.4	108
9	Atomic structure of a nanobody-trapped domain-swapped dimer of an amyloidogenic β2-microglobulin variant. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 1314-1319.	7.1	108
10	Targeting Amyloid Aggregation: An Overview of Strategies and Mechanisms. International Journal of Molecular Sciences, 2018, 19, 2677.	4.1	103
11	Proteolytic cleavage of Ser52Pro variant transthyretin triggers its amyloid fibrillogenesis. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 1539-1544.	7.1	91
12	Effect of Tetracyclines on the Dynamics of Formation and Destructuration of β2-Microglobulin Amyloid Fibrils. Journal of Biological Chemistry, 2011, 286, 2121-2131.	3.4	87
13	Conformational Switching and Fibrillogenesis in the Amyloidogenic Fragment of Apolipoprotein A-I. Journal of Biological Chemistry, 2003, 278, 2444-2451.	3.4	86
14	The Controlling Roles of Trp60 and Trp95 in β2-Microglobulin Function, Folding and Amyloid Aggregation Properties. Journal of Molecular Biology, 2008, 378, 887-897.	4.2	82
15	Structure, Folding Dynamics, and Amyloidogenesis of D76N β2-Microglobulin. Journal of Biological Chemistry, 2013, 288, 30917-30930.	3.4	80
16	Liver biopsy discloses a new apolipoprotein A-I hereditary amyloidosis in several unrelated Italian families. Gastroenterology, 2004, 126, 1416-1422.	1.3	70
17	Plasminogen activation triggers transthyretin amyloidogenesis in vitro. Journal of Biological Chemistry, 2018, 293, 14192-14199.	3.4	68
18	Properties of Some Variants of Human β2-Microglobulin and Amyloidogenesis. Journal of Biological Chemistry, 2004, 279, 9176-9189.	3.4	65

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19	In situ characterization of protein aggregates in human tissues affected by light chain amyloidosis: a FTIR microspectroscopy study. Scientific Reports, 2016, 6, 29096.	3.3	63
20	The workings of the amyloid diseases. Annals of Medicine, 2007, 39, 200-207.	3.8	62
21	Â2-Microglobulin isoforms display an heterogeneous affinity for type I collagen. Protein Science, 2005, 14, 696-702.	7.6	56
22	Native-unlike Long-lived Intermediates along the Folding Pathway of the Amyloidogenic Protein β2-Microglobulin Revealed by Real-time Two-dimensional NMR. Journal of Biological Chemistry, 2010, 285, 5827-5835.	3.4	55
23	Topological investigation of amyloid fibrils obtained from β2-microglobulin. Protein Science, 2009, 11, 2362-2369.	7.6	53
24	Capillary electrophoresis investigation of a partially unfolded conformation of β2-microglobulin. Electrophoresis, 2002, 23, 918-925.	2.4	52
25	Pharmaceutical Strategies Against Amyloidosis: Old and New Drugs in Targeting a "Protein Misfolding Disease". Current Medicinal Chemistry, 2004, 11, 1065-1084.	2.4	48
26	Effects of the Known Pathogenic Mutations on the Aggregation Pathway of the Amyloidogenic Peptide of Apolipoprotein A-I. Journal of Molecular Biology, 2011, 407, 465-476.	4.2	48
27	Amyloid fibrils derived from the apolipoprotein A1 Leu174Ser variant contain elements of ordered helical structure. Protein Science, 2001, 10, 187-199.	7.6	44
28	β2-Microglobulin H31Y Variant 3D Structure Highlights the Protein Natural Propensity Towards Intermolecular Aggregation. Journal of Molecular Biology, 2004, 335, 1051-1064.	4.2	38
29	Limited proteolysis in the investigation of β2-microglobulin amyloidogenic and fibrillar states. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2005, 1753, 44-50.	2.3	36
30	Proteomics of β2-microglobulin amyloid fibrils. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2005, 1753, 23-33.	2.3	36
31	DE loop mutations affect β2-microglobulin stability and amyloid aggregation. Biochemical and Biophysical Research Communications, 2008, 377, 146-150.	2.1	36
32	Folding and Fibrillogenesis: Clues from β2-Microglobulin. Journal of Molecular Biology, 2010, 401, 286-297.	4.2	35
33	Fibrillar vs Crystalline Full-Length β-2-Microglobulin Studied by High-Resolution Solid-State NMR Spectroscopy. Journal of the American Chemical Society, 2010, 132, 5556-5557.	13.7	32
34	Monitoring the Interaction between β2-Microglobulin and the Molecular Chaperone αB-crystallin by NMR and Mass Spectrometry. Journal of Biological Chemistry, 2013, 288, 17844-17858.	3.4	32
35	Â2-Microglobulin is potentially neurotoxic, but the blood brain barrier is likely to protect the brain from its toxicity. Nephrology Dialysis Transplantation, 2008, 24, 1176-1181.	0.7	31
36	Inhibition of the mechano-enzymatic amyloidogenesis of transthyretin: role of ligand affinity, binding cooperativity and occupancy of the inner channel. Scientific Reports, 2017, 7, 182.	3.3	31

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37	A FTIR microspectroscopy study of the structural and biochemical perturbations induced by natively folded and aggregated transthyretin in HL-1 cardiomyocytes. Scientific Reports, 2018, 8, 12508.	3.3	31
38	Oleuropein aglycone: A polyphenol with different targets against amyloid toxicity. Biochimica Et Biophysica Acta - General Subjects, 2018, 1862, 1432-1442.	2.4	30
39	Human beta-2 microglobulin W60V mutant structure: Implications for stability and amyloid aggregation. Biochemical and Biophysical Research Communications, 2009, 380, 543-547.	2.1	29
40	Co-fibrillogenesis of Wild-type and D76N β2-Microglobulin. Journal of Biological Chemistry, 2016, 291, 9678-9689.	3.4	29
41	Crtap and p3h1 knock out zebrafish support defective collagen chaperoning as the cause of their osteogenesis imperfecta phenotype. Matrix Biology, 2020, 90, 40-60.	3.6	28
42	Citrate-stabilized gold nanoparticles hinder fibrillogenesis of a pathological variant of β ₂ -microglobulin. Nanoscale, 2017, 9, 3941-3951.	5.6	26
43	Solution structure of β2-microglobulin and insights into fibrillogenesis. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2005, 1753, 76-84.	2.3	25
44	Single Point Mutations Induce a Switch in the Molecular Mechanism of the Aggregation of the Alzheimer's Disease Associated Aβ ₄₂ Peptide. ACS Chemical Biology, 2014, 9, 378-382.	3.4	25
45	Amyloid persistence in decellularized liver: biochemical and histopathological characterization. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2016, 23, 1-7.	3.0	25
46	Lysine 58-cleaved beta2-microglobulin is not detectable by 2D electrophoresis in ex vivo amyloid fibrils of two patients affected by dialysis-related amyloidosis. Protein Science, 2006, 16, 343-349.	7.6	24
47	Benefit of doxycycline treatment on articular disability caused by dialysis related amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2013, 20, 173-178.	3.0	24
48	Dynamic of β2â€Microglobulin Fibril Formation and Reabsorption: The Role of Proteolysis. Seminars in Dialysis, 2001, 14, 117-122.	1.3	23
49	Structural and Folding Dynamic Properties of the T70N Variant of Human Lysozyme. Journal of Biological Chemistry, 2003, 278, 25910-25918.	3.4	23
50	Molecular insights into cell toxicity of a novel familial amyloidogenic variant of β2â€microglobulin. Journal of Cellular and Molecular Medicine, 2016, 20, 1443-1456.	3.6	23
51	Detection of fragments of \hat{l}^2 2-microglobulin in amyloid fibrils. Kidney International, 2000, 57, 349-350.	5.2	22
52	The intracellular quality control system down-regulates the secretion of amyloidogenic apolipoprotein A-l variants: A possible impact on the natural history of the disease. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2011, 1812, 87-93.	3.8	22
53	Class I Major Histocompatibility Complex, the Trojan Horse for Secretion of Amyloidogenic β2-Microglobulin. Journal of Biological Chemistry, 2014, 289, 3318-3327.	3.4	22
54	Decoding the Structural Bases of D76N ß2-Microglobulin High Amyloidogenicity through Crystallography and Asn-Scan Mutagenesis. PLoS ONE, 2015, 10, e0144061.	2.5	22

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55	C. elegans Expressing Human β2-Microglobulin: A Novel Model for Studying the Relationship between the Molecular Assembly and the Toxic Phenotype. PLoS ONE, 2012, 7, e52314.	2.5	21
56	A quantitative and qualitative method for direct 2â€DE analysis of murine cartilage. Proteomics, 2007, 7, 4003-4007.	2.2	20
57	Equilibrium Unfolding Thermodynamics of β2-Microglobulin Analyzed through Native-State H/D Exchange. Biophysical Journal, 2009, 96, 169-179.	0.5	20
58	Variants of β2-microglobulin cleaved at lysine-58 retain the main conformational features of the native protein but are more conformationally heterogeneous and unstable at physiological temperature. FEBS Journal, 2006, 273, 2461-2474.	4.7	19
59	Biochemical and Electrophysiological Modification of Amyloid Transthyretin on Cardiomyocytes. Biophysical Journal, 2016, 111, 2024-2038.	0.5	19
60	Recombinant amyloidogenic domain of ApoA-I: Analysis of its fibrillogenic potential. Biochemical and Biophysical Research Communications, 2006, 351, 223-228.	2.1	18
61	Sulfonated molecules that bind a partially structured species of β ₂ â€microglobulin also influence refolding and fibrillogenesis. Electrophoresis, 2008, 29, 1502-1510.	2.4	18
62	A specific nanobody prevents amyloidogenesis of D76N β2-microglobulin in vitro and modifies its tissue distribution in vivo. Scientific Reports, 2017, 7, 46711.	3.3	18
63	Search of ligands for the amyloidogenic protein β2-microglobulin by capillary electrophoresis and other techniques. Electrophoresis, 2005, 26, 4055-4063.	2.4	17
64	Screening of fibrillogenesis inhibitors of β2-microglobulin: Integrated strategies by mass spectrometry capillary electrophoresis and in silico simulations. Analytica Chimica Acta, 2011, 685, 153-161.	5.4	17
65	Enhanced molecular chaperone activity of the small heatâ€shock protein αBâ€crystallin following covalent immobilization onto a solidâ€phase support. Biopolymers, 2011, 95, 376-389.	2.4	14
66	Increasing the accuracy of proteomic typing by decellularisation of amyloid tissue biopsies. Journal of Proteomics, 2017, 165, 113-118.	2.4	14
67	C. elegans expressing D76N l²2-microglobulin: a model for in vivo screening of drug candidates targeting amyloidosis. Scientific Reports, 2019, 9, 19960.	3.3	14
68	2â€DE and MALDIâ€TOFâ€MS for a comparative analysis of proteins expressed in different cellular models of amyotrophic lateral sclerosis. Electrophoresis, 2007, 28, 4320-4329.	2.4	13
69	Dâ€strand perturbation and amyloid propensity in betaâ€2 microglobulin. FEBS Journal, 2011, 278, 2349-2358.	4.7	13
70	Comparative study of the stabilities of synthetic in vitro and natural ex vivo transthyretin amyloid fibrils. Journal of Biological Chemistry, 2020, 295, 11379-11387.	3.4	12
71	Interference of citrate-stabilized gold nanoparticles with β2-microglobulin oligomeric association. Chemical Communications, 2018, 54, 5422-5425.	4.1	11
72	Amyloid Formation by Globular Proteins: The Need to Narrow the Gap Between in Vitro and in Vivo Mechanisms. Frontiers in Molecular Biosciences, 2022, 9, 830006.	3.5	11

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73	Reduction of conformational mobility and aggregation in W60G β ₂ â€microglobulin: assessment by ¹⁵ N NMR relaxation. Magnetic Resonance in Chemistry, 2013, 51, 795-807.	1.9	10
74	The effects of an ideal β-turn on β-2 microglobulin fold stability. Journal of Biochemistry, 2011, 150, 39-47.	1.7	9
75	Enhanced toxicity of silver nanoparticles in transgenic <i>Caenorhabditis elegans</i> expressing amyloidogenic proteins. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2015, 22, 221-228.	3.0	9
76	Proteomics in protein misfolding diseases. Clinical Chemistry and Laboratory Medicine, 2009, 47, 627-35.	2.3	8
77	Fibrillogenesis of human <i>β</i> ₂ â€microglobulin in threeâ€dimensional silicon microstructures. Journal of Biophotonics, 2012, 5, 785-792.	2.3	8
78	Misidentification of transthyretin and immunoglobulin variants by proteomics due to methyl lysine formation in formalin-fixed paraffin-embedded amyloid tissue. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 229-237.	3.0	8
79	Purification and Characterization of SolubleCichorium intybusVar.silvestreLipoxygenase. Journal of Agricultural and Food Chemistry, 2005, 53, 6448-6454.	5.2	7
80	The interaction of β2-microglobulin with gold nanoparticles: impact of coating, charge and size. Journal of Materials Chemistry B, 2018, 6, 5964-5974.	5.8	7
81	Insights into a Protein-Nanoparticle System by Paramagnetic Perturbation NMR Spectroscopy. Molecules, 2020, 25, 5187.	3.8	7
82	Capillary electrophoresis analysis of different variants of the amyloidogenic protein β ₂ â€microglobulin as a simple tool for misfolding and stability studies. Electrophoresis, 2015, 36, 2465-2472.	2.4	6
83	Hepatitis C virusâ€associated cryoglobulinaemicglomerulonephritis with delayed appearance ofmonoclonal cryoglobulinaemia. Nephrology Dialysis Transplantation, 2001, 16, 432-434.	0.7	5
84	Clinical, radiological, and biochemical features of a bilateral buttock amyloidoma emerging after 27 years of hemodialysis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2009, 16, 115-121.	3.0	5
85	S-Homocysteinylation effects on transthyretin: worsening of cardiomyopathy onset. Biochimica Et Biophysica Acta - General Subjects, 2020, 1864, 129453.	2.4	5
86	Exploring exchange processes in proteins by paramagnetic perturbation of NMR spectra. Physical Chemistry Chemical Physics, 2020, 22, 6247-6259.	2.8	5
87	The corona of protein–gold nanoparticle systems: the role of ionic strength. Physical Chemistry Chemical Physics, 2022, 24, 1630-1637.	2.8	5
88	Short-Chain Alkanethiol Coating for Small-Size Gold Nanoparticles Supporting Protein Stability. Magnetochemistry, 2017, 3, 40.	2.4	4
89	Clinical ApoAâ€ŧV amyloid is associated with fibrillogenic signal sequence. Journal of Pathology, 2021, 255, 311-318.	4.5	4
90	Characterization of immunoglobulin variable regions of two human pathogenic monoclonal cryocrystalglobulins. Molecular Immunology, 2008, 45, 1519-1524.	2.2	2

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91	Topologically non-trivial metal-organic assemblies inhibit β2-microglobulin amyloidogenesis. Cell Reports Physical Science, 2021, 2, 100477.	5.6	1