Avijit Chakrabartty

List of Publications by Year in descending order

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90 papers 7,611 citations

42 h-index 51492 86 g-index

96 all docs 96 docs citations

96 times ranked 8113 citing authors

| # | Article | IF | Citations |
|----|---|------|-----------|
| 1 | Helix propensities of the amino acids measured in alanineâ€based peptides without helixâ€stabilizing sideâ€chain interactions. Protein Science, 1994, 3, 843-852. | 3.1 | 572 |
| 2 | Stability of α-Helices. Advances in Protein Chemistry, 1995, 46, 141-176. | 4.4 | 339 |
| 3 | Aromatic side-chain contribution to far-ultraviolet circular dichroism of helical peptides and its effect on measurement of helix propensities. Biochemistry, 1993, 32, 5560-5565. | 1.2 | 334 |
| 4 | Large differences in the helix propensities of alanine and glycine. Nature, 1991, 351, 586-588. | 13.7 | 325 |
| 5 | Membrane Disruption by Alzheimer \hat{l}^2 -Amyloid Peptides Mediated through Specific Binding to Either Phospholipids or Gangliosides. Journal of Biological Chemistry, 1996, 271, 26482-26489. | 1.6 | 307 |
| 6 | Monomeric Cu,Zn-superoxide Dismutase Is a Common Misfolding Intermediate in the Oxidation Models of Sporadic and Familial Amyotrophic Lateral Sclerosis. Journal of Biological Chemistry, 2004, 279, 15499-15504. | 1.6 | 296 |
| 7 | Oxidation-induced Misfolding and Aggregation of Superoxide Dismutase and Its Implications for Amyotrophic Lateral Sclerosis. Journal of Biological Chemistry, 2002, 277, 47551-47556. | 1.6 | 279 |
| 8 | Helix propagation and Nâ€cap propensities of the amino acids measured in alanineâ€based peptides in 40 volume percent trifluoroethanol. Protein Science, 1996, 5, 2623-2637. | 3.1 | 256 |
| 9 | A prion protein epitope selective for the pathologically misfolded conformation. Nature Medicine, 2003, 9, 893-899. | 15.2 | 252 |
| 10 | Manipulating the Amyloid- \hat{l}^2 Aggregation Pathway with Chemical Chaperones. Journal of Biological Chemistry, 1999, 274, 32970-32974. | 1.6 | 238 |
| 11 | Structural studies of soluble oligomers of the alzheimer β-amyloid peptide. Journal of Molecular Biology, 2000, 297, 73-87. | 2.0 | 217 |
| 12 | An immunological epitope selective for pathological monomer-misfolded SOD1 in ALS. Nature Medicine, 2007, 13, 754-759. | 15.2 | 199 |
| 13 | Characterization of the Interactions of Alzheimer beta-Amyloid Peptides with Phospholipid Membranes. FEBS Journal, 1997, 245, 355-363. | 0.2 | 189 |
| 14 | Determination of Free Energies of N-Capping in .alphaHelixes by Modification of the Lifson-Roig Helix-Coil Theory To Include N- and C-Capping. Biochemistry, 1994, 33, 3396-3403. | 1.2 | 180 |
| 15 | Structure, folding, and misfolding of Cu,Zn superoxide dismutase in amyotrophic lateral sclerosis. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2006, 1762, 1025-1037. | 1.8 | 178 |
| 16 | Lipopeptide detergents designed for the structural study of membrane proteins. Nature Biotechnology, 2003, 21, 171-176. | 9.4 | 174 |
| 17 | Structural Transitions Associated with the Interaction of Alzheimer \hat{l}^2 -Amyloid Peptides with Gangliosides. Journal of Biological Chemistry, 1998, 273, 4506-4515. | 1.6 | 173 |
| 18 | Amyotrophic lateral sclerosis is a non-amyloid disease in which extensive misfolding of SOD1 is unique to the familial form. Acta Neuropathologica, 2010, 119, 335-344. | 3.9 | 171 |

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|----|---|-----|-----------|
| 19 | Characterization of Segments from the Central Region of BRCA1: An Intrinsically Disordered Scaffold for Multiple Protein–Protein and Protein–DNA Interactions?. Journal of Molecular Biology, 2005, 345, 275-287. | 2.0 | 157 |
| 20 | Autoinhibition of the Kit Receptor Tyrosine Kinase by the Cytosolic Juxtamembrane Region. Molecular and Cellular Biology, 2003, 23, 3067-3078. | 1.1 | 151 |
| 21 | The effect of enhanced alpha-helicity on the activity of a winter flounder antifreeze polypeptide. FEBS Journal, 1991, 202, 1057-1063. | 0.2 | 138 |
| 22 | Adaptor Protein Self-Assembly Drives the Control of a Cullin-RING Ubiquitin Ligase. Structure, 2012, 20, 1141-1153. | 1.6 | 127 |
| 23 | Prion disease susceptibility is affected by \hat{l}^2 -structure folding propensity and local side-chain interactions in PrP. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 19808-19813. | 3.3 | 119 |
| 24 | The molecular interaction of human salivary histatins with polyphenolic compounds. FEBS Journal, 2001, 268, 4384-4397. | 0.2 | 115 |
| 25 | Alternate Aggregation Pathways of the Alzheimer β-Amyloid Peptide: Aβ Association Kinetics at Endosomal pH. Journal of Molecular Biology, 2003, 325, 743-757. | 2.0 | 97 |
| 26 | Targeting of Monomer/Misfolded SOD1 as a Therapeutic Strategy for Amyotrophic Lateral Sclerosis. Journal of Neuroscience, 2012, 32, 8791-8799. | 1.7 | 87 |
| 27 | Hyperactive Antifreeze Protein from Winter Flounder Is a Very Long Rod-like Dimer of α-Helices*. Journal of Biological Chemistry, 2005, 280, 17920-17929. | 1.6 | 73 |
| 28 | Dimerization of the transmembrane domain of amyloid precursor proteins and familial Alzheimer's disease mutants. BMC Neuroscience, 2008, 9, 17. | 0.8 | 73 |
| 29 | Low molecular weight species of TDP-43 generated by abnormal splicing form inclusions in amyotrophic lateral sclerosis and result in motor neuron death. Acta Neuropathologica, 2015, 130, 49-61. | 3.9 | 71 |
| 30 | Alternate Aggregation Pathways of the Alzheimer \hat{l}^2 -Amyloid Peptide. Journal of Biological Chemistry, 2000, 275, 36436-36440. | 1.6 | 69 |
| 31 | Protein misfolding in the lateâ€onset neurodegenerative diseases: Common themes and the unique case of amyotrophic lateral sclerosis. Proteins: Structure, Function and Bioinformatics, 2013, 81, 1285-1303. | 1.5 | 69 |
| 32 | Novel conformation-specific monoclonal antibodies against amyloidogenic forms of transthyretin. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2016, 23, 86-97. | 1.4 | 69 |
| 33 | Phase to Phase with TDP-43. Biochemistry, 2017, 56, 809-823. | 1.2 | 68 |
| 34 | CCM3/PDCD10 Heterodimerizes with Germinal Center Kinase III (GCKIII) Proteins Using a Mechanism Analogous to CCM3 Homodimerization. Journal of Biological Chemistry, 2011, 286, 25056-25064. | 1.6 | 67 |
| 35 | "Structural characterization of the minimal segment of TDP-43 competent for aggregation― Archives of Biochemistry and Biophysics, 2014, 545, 53-62. | 1.4 | 67 |
| 36 | Transthyretin amyloidosis: an under-recognized neuropathy and cardiomyopathy. Clinical Science, 2017, 131, 395-409. | 1.8 | 66 |

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| 37 | Fibrillogenesis of Alzheimer A \hat{l}^2 peptides studied by fluorescence energy transfer. Journal of Molecular Biology, 1997, 269, 214-224. | 2.0 | 64 |
| 38 | Interaction of human and mouse AÎ ² peptides. Journal of Neurochemistry, 2004, 91, 1398-1403. | 2.1 | 51 |
| 39 | Charge substitution shows that repulsive electrostatic interactions impede the oligomerization of Alzheimer amyloid peptides. FEBS Letters, 2005, 579, 3574-3578. | 1.3 | 51 |
| 40 | Alzheimer \hat{l}^2 -amyloid peptides: Structures of amyloid fibrils and alternate aggregation products. Biopolymers, 2001, 60, 381. | 1.2 | 50 |
| 41 | Denaturational Stress Induces Formation of Zinc-Deficient Monomers of Cu,Zn Superoxide Dismutase: Implications for Pathogenesis in Amyotrophic Lateral Sclerosis. Journal of Molecular Biology, 2008, 383, 424-436. | 2.0 | 44 |
| 42 | Early Steps in Oxidation-Induced SOD1 Misfolding: Implications for Non-Amyloid Protein Aggregation in Familial ALS. Journal of Molecular Biology, 2012, 421, 631-652. | 2.0 | 44 |
| 43 | Variants of DsRed fluorescent protein: Development ofÂa copper sensor. Protein Science, 2006, 15, 2442-2447. | 3.1 | 43 |
| 44 | Amyloid \hat{I}^2 -protein (A \hat{I}^2) associated with lipid molecules: immunoreactivity distinct from that of soluble A \hat{I}^2 . FEBS Letters, 1997, 420, 43-46. | 1.3 | 42 |
| 45 | Co-incorporation of A \hat{I}^2 40 and A \hat{I}^2 42 to form mixed pre-fibrillar aggregates. FEBS Journal, 2003, 270, 654-663. | 0.2 | 40 |
| 46 | The PrP-like Protein Doppel Binds Copper. Journal of Biological Chemistry, 2003, 278, 8888-8896. | 1.6 | 39 |
| 47 | ALS-Causing SOD1 Mutations Promote Production of Copper-Deficient Misfolded Species. Journal of Molecular Biology, 2011, 409, 839-852. | 2.0 | 39 |
| 48 | Determining composition of micron-scale protein deposits in neurodegenerative disease by spatially targeted optical microproteomics. ELife, 2015, 4, . | 2.8 | 38 |
| 49 | Somatostatin binds to the human amyloid \hat{l}^2 peptide and favors the formation of distinct oligomers. ELife, 2017, 6, . | 2.8 | 37 |
| 50 | Binding of TDP-43 to the 3′UTR of Its Cognate mRNA Enhances Its Solubility. Biochemistry, 2014, 53, 5885-5894. | 1.2 | 36 |
| 51 | Alzheimer's AÎ ² 40 Studied by NMR at Low pH Reveals That Sodium 4,4-Dimethyl-4-silapentane-1-sulfonate (DSS) Binds and Promotes Î ² -Ball Oligomerization. Journal of Biological Chemistry, 2005, 280, 3675-3685. | 1.6 | 34 |
| 52 | Requirement of aggregation propensity of Alzheimer amyloid peptides for neuronal cell surface binding. BMC Neuroscience, 2007, 8, 29. | 0.8 | 33 |
| 53 | Electrostatic Repulsion Governs TDP-43 C-terminal Domain Aggregation. PLoS Biology, 2016, 14, e1002447. | 2.6 | 33 |
| 54 | Quercitrin and quercetin 3-Î ² -d-glucoside as chemical chaperones for the A4V SOD1 ALS-causing mutant. Protein Engineering, Design and Selection, 2017, 30, 431-440. | 1.0 | 33 |

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| 55 | Reversible assembly of helical filaments by de novo designed minimalist peptides. Biopolymers, 2005, 80, 26-33. | 1.2 | 32 |
| 56 | Primary structures of the alanine-rich antifreeze polypeptides from grubby sculpin, <i>Myoxocephalus aenaeus</i> . Canadian Journal of Zoology, 1988, 66, 403-408. | 0.4 | 26 |
| 57 | Species barriers for chronic wasting disease by in vitro conversion of prion protein. Biochemical and Biophysical Research Communications, 2007, 364, 796-800. | 1.0 | 26 |
| 58 | Substoichiometric inhibition of transthyretin misfolding by immune-targeting sparsely populated misfolding intermediates: a potential diagnostic and therapeutic for TTR amyloidoses. Scientific Reports, 2016, 6, 25080. | 1.6 | 26 |
| 59 | Structural and functional characterization of KEOPS dimerization by Pcc1 and its role in t ⁶ A biosynthesis. Nucleic Acids Research, 2016, 44, 6971-6980. | 6.5 | 26 |
| 60 | Simple Elimination of Background Fluorescence in Formalin-Fixed Human Brain Tissue for Immunofluorescence Microscopy. Journal of Visualized Experiments, 2017, , . | 0.2 | 26 |
| 61 | Physiologically Important Electrolytes as Regulators of TDP-43 Aggregation and Droplet-Phase Behavior. Biochemistry, 2019, 58, 590-607. | 1.2 | 24 |
| 62 | Identification of stable helical bundles from a combinatorial library of amphipathic peptides. Biopolymers, 2004, 76, 244-257. | 1.2 | 22 |
| 63 | Relative and Regional Stabilities of the Hamster, Mouse, Rabbit, and Bovine Prion Proteins toward Urea Unfolding Assessed by Nuclear Magnetic Resonance and Circular Dichroism Spectroscopies. Biochemistry, 2011, 50, 7536-7545. | 1.2 | 22 |
| 64 | Equilibrium folding intermediates of a greek key \hat{l}^2 -barrel protein. Journal of Molecular Biology, 1998, 276, 669-681. | 2.0 | 21 |
| 65 | Two Distinct Conformations of $\hat{Al^2}$ Aggregates on the Surface of Living PC12 Cells. Biophysical Journal, 2009, 96, 4260-4267. | 0.2 | 19 |
| 66 | Calexcitin B Is a New Member of the Sarcoplasmic Calcium-binding Protein Family. Journal of Biological Chemistry, 2001, 276, 22529-22536. | 1.6 | 17 |
| 67 | Interplay of buried histidine protonation and protein stability in prion misfolding. Scientific Reports, 2017, 7, 882. | 1.6 | 17 |
| 68 | N-Terminal Helix-Cap in \hat{l}_{\pm} -Helix 2 Modulates \hat{l}^{2} -State Misfolding in Rabbit and Hamster Prion Proteins. PLoS ONE, 2013, 8, e63047. | 1.1 | 17 |
| 69 | Interactions of Alzheimer amyloid peptides with cultured cells and brain tissue, and their biological consequences. Biopolymers, 2004, 76, 4-14. | 1.2 | 16 |
| 70 | Wild-type Cu/Zn superoxide dismutase stabilizes mutant variants by heterodimerization. Neurobiology of Disease, 2014, 62, 479-488. | 2.1 | 16 |
| 71 | Getting specificity from simplicity in putative proteins from the prebiotic Earth. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 14941-14946. | 3.3 | 15 |
| 72 | Cell Surface Binding and Internalization of Al^2Modulated by Degree of Aggregation. International Journal of Alzheimer's Disease, 2011, 2011, 1-13. | 1.1 | 13 |

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| 73 | Conformational Coupling of Mg2+and Ca2+on the Three-State Folding of Calexcitin Bâ€. Biochemistry, 2003, 42, 5531-5539. | 1.2 | 12 |
| 74 | Cost-effective elimination of lipofuscin fluorescence from formalin-fixed brain tissue by white phosphor light emitting diode array. Biochemistry and Cell Biology, 2016, 94, 545-550. | 0.9 | 12 |
| 75 | Putative Oneâ€Pot Prebiotic Polypeptides with Ribonucleolytic Activity. Chemistry - A European Journal, 2010, 16, 5314-5323. | 1.7 | 11 |
| 76 | All or none fibrillogenesis of a prion peptide. FEBS Journal, 2001, 268, 4885-4891. | 0.2 | 10 |
| 77 | Conversion of A \hat{I}^2 42 into a Folded Soluble Native-like Protein using a Semi-random Library of Amphipathic Helices. Journal of Molecular Biology, 2010, 396, 1284-1294. | 2.0 | 10 |
| 78 | Probing Alzheimer amyloid peptide aggregation using a cell-free fluorescent protein refolding method. Biochemistry and Cell Biology, 2009, 87, 631-639. | 0.9 | 9 |
| 79 | Multiphoton ANS fluorescence microscopy as an in vivo sensor for protein misfolding stress. Cell Stress and Chaperones, 2011, 16, 549-561. | 1.2 | 9 |
| 80 | Progress in transthyretin fibrillogenesis research strengthens the amyloid hypothesis. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 14757-14759. | 3.3 | 7 |
| 81 | Analyzing complicated protein folding kinetics rapidly by analytical Laplace inversion using a Tikhonov regularization variant. Analytical Biochemistry, 2012, 421, 181-190. | 1.1 | 7 |
| 82 | An Arg-rich putative prebiotic protein is as stable as its Lys-rich variant. Archives of Biochemistry and Biophysics, 2012, 528, 118-126. | 1.4 | 6 |
| 83 | Structure of a simplified \hat{l}^2 -hairpin and its ATP complex. Archives of Biochemistry and Biophysics, 2013, 537, 62-71. | 1.4 | 6 |
| 84 | Conformation specificity and arene binding in a peptide composed only of Lys, Ile, Ala and Gly. European Biophysics Journal, 2012, 41, 63-72. | 1.2 | 5 |
| 85 | Nonpolar contributions to conformational specificity in assemblies of designed short helical peptides. Protein Science, 2000, 9, 1011-1023. | 3.1 | 4 |
| 86 | Alternate routes to conformational specificity in a Greek key \hat{l}^2 barrel protein. FEBS Journal, 2001, 268, 4653-4664. | 0.2 | 2 |
| 87 | NMR-driven secondary and tertiary structure model of Ca2+-loaded calexcitin. Biochemical and Biophysical Research Communications, 2006, 343, 520-524. | 1.0 | 2 |
| 88 | Protein Misfolding and Toxicity in Amyotrophic Lateral Sclerosis., 2012,, 257-288. | | 2 |
| 89 | Reply to "Properties of a disease-specific prion probe― Nature Medicine, 2004, 10, 11-12. | 15.2 | 1 |
| 90 | Alzheimer \hat{I}^2 -amyloid peptides: Structures of amyloid fibrils and alternate aggregation products. , 2001, 60, 381. | | 1 |