## **Avijit Chakrabartty**

## List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/958296/avijit-chakrabartty-publications-by-year.pdf

Version: 2024-04-23

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

6,715 81 40 91 h-index g-index citations papers 96 7.8 7,209 5.55 L-index avg, IF ext. papers ext. citations

#	Paper	IF	Citations
91	Physiologically Important Electrolytes as Regulators of TDP-43 Aggregation and Droplet-Phase Behavior. <i>Biochemistry</i> , <b>2019</b> , 58, 590-607	3.2	12
90	Phase to Phase with TDP-43. Biochemistry, 2017, 56, 809-823	3.2	47
89	Transthyretin amyloidosis: an under-recognized neuropathy and cardiomyopathy. <i>Clinical Science</i> , <b>2017</b> , 131, 395-409	6.5	50
88	Interplay of buried histidine protonation and protein stability in prion misfolding. <i>Scientific Reports</i> , <b>2017</b> , 7, 882	4.9	14
87	Quercitrin and quercetin 3-Ed-glucoside as chemical chaperones for the A4V SOD1 ALS-causing mutant. <i>Protein Engineering, Design and Selection</i> , <b>2017</b> , 30, 431-440	1.9	22
86	Simple Elimination of Background Fluorescence in Formalin-Fixed Human Brain Tissue for Immunofluorescence Microscopy. <i>Journal of Visualized Experiments</i> , <b>2017</b> ,	1.6	14
85	Somatostatin binds to the human amyloid [peptide and favors the formation of distinct oligomers. <i>ELife</i> , <b>2017</b> , 6,	8.9	21
84	Cost-effective elimination of lipofuscin fluorescence from formalin-fixed brain tissue by white phosphor light emitting diode array. <i>Biochemistry and Cell Biology</i> , <b>2016</b> , 94, 545-550	3.6	8
83	Structural and functional characterization of KEOPS dimerization by Pcc1 and its role in t6A biosynthesis. <i>Nucleic Acids Research</i> , <b>2016</b> , 44, 6971-80	20.1	17
82	Novel conformation-specific monoclonal antibodies against amyloidogenic forms of transthyretin. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , <b>2016</b> , 23, 86-97	2.7	54
81	Electrostatic Repulsion Governs TDP-43 C-terminal Domain Aggregation. <i>PLoS Biology</i> , <b>2016</b> , 14, e1002	4 <del>9</del> .7⁄	23
80	Substoichiometric inhibition of transthyretin misfolding by immune-targeting sparsely populated misfolding intermediates: a potential diagnostic and therapeutic for TTR amyloidoses. <i>Scientific Reports</i> , <b>2016</b> , 6, 25080	4.9	22
79	Low molecular weight species of TDP-43 generated by abnormal splicing form inclusions in amyotrophic lateral sclerosis and result in motor neuron death. <i>Acta Neuropathologica</i> , <b>2015</b> , 130, 49-6	1 <sup>14.3</sup>	49
78	Determining composition of micron-scale protein deposits in neurodegenerative disease by spatially targeted optical microproteomics. <i>ELife</i> , <b>2015</b> , 4,	8.9	23
77	Wild-type Cu/Zn superoxide dismutase stabilizes mutant variants by heterodimerization. <i>Neurobiology of Disease</i> , <b>2014</b> , 62, 479-88	7.5	15
76	"Structural characterization of the minimal segment of TDP-43 competent for aggregation". <i>Archives of Biochemistry and Biophysics</i> , <b>2014</b> , 545, 53-62	4.1	54
75	Binding of TDP-43 to the 3'UTR of its cognate mRNA enhances its solubility. <i>Biochemistry</i> , <b>2014</b> , 53, 588	35 <u>,9</u> 4	24

## (2010-2013)

Structure of a simplified Ehairpin and its ATP complex. <i>Archives of Biochemistry and Biophysics</i> , <b>2013</b> , 537, 62-71	4.1	4
Protein misfolding in the late-onset neurodegenerative diseases: common themes and the unique case of amyotrophic lateral sclerosis. <i>Proteins: Structure, Function and Bioinformatics</i> , <b>2013</b> , 81, 1285-30	)3 <sup>4.2</sup>	62
N-terminal helix-cap in Ehelix 2 modulates Estate misfolding in rabbit and hamster prion proteins. <i>PLoS ONE</i> , <b>2013</b> , 8, e63047	3.7	15
Adaptor protein self-assembly drives the control of a cullin-RING ubiquitin ligase. <i>Structure</i> , <b>2012</b> , 20, 1141-53	5.2	90
Conformation specificity and arene binding in a peptide composed only of Lys, Ile, Ala and Gly. <i>European Biophysics Journal</i> , <b>2012</b> , 41, 63-72	1.9	4
An Arg-rich putative prebiotic protein is as stable as its Lys-rich variant. <i>Archives of Biochemistry and Biophysics</i> , <b>2012</b> , 528, 118-26	4.1	6
Early steps in oxidation-induced SOD1 misfolding: implications for non-amyloid protein aggregation in familial ALS. <i>Journal of Molecular Biology</i> , <b>2012</b> , 421, 631-52	6.5	38
Analyzing complicated protein folding kinetics rapidly by analytical Laplace inversion using a Tikhonov regularization variant. <i>Analytical Biochemistry</i> , <b>2012</b> , 421, 181-90	3.1	7
Targeting of monomer/misfolded SOD1 as a therapeutic strategy for amyotrophic lateral sclerosis. <i>Journal of Neuroscience</i> , <b>2012</b> , 32, 8791-9	6.6	71
Protein Misfolding and Toxicity in Amyotrophic Lateral Sclerosis <b>2012</b> , 257-288		2
ALS-causing SOD1 mutations promote production of copper-deficient misfolded species. <i>Journal of Molecular Biology</i> , <b>2011</b> , 409, 839-52	6.5	33
Cell surface binding and internalization of almodulated by degree of aggregation. <i>International Journal of Alzheimeris Disease</i> , <b>2011</b> , 2011, 962352	3.7	10
Interaction of Alzheimer Amyloid Peptide with Cell Surfaces and Artificial Membranes <b>2011</b> , 231-243		1
Multiphoton ANS fluorescence microscopy as an in vivo sensor for protein misfolding stress. <i>Cell Stress and Chaperones</i> , <b>2011</b> , 16, 549-61	4	8
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. <i>Biochemistry</i> , <b>2011</b> , 50, 7536-45	3.2	21
CCM3/PDCD10 heterodimerizes with germinal center kinase III (GCKIII) proteins using a mechanism analogous to CCM3 homodimerization. <i>Journal of Biological Chemistry</i> , <b>2011</b> , 286, 25056-64	5.4	50
Prion disease susceptibility is affected by beta-structure folding propensity and local side-chain interactions in PrP. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2010</b> , 107, 19808-13	11.5	102
Conversion of Abeta42 into a folded soluble native-like protein using a semi-random library of amphipathic helices. <i>Journal of Molecular Biology</i> , <b>2010</b> , 396, 1284-94	6.5	10
	Protein misfolding in the late-onset neurodegenerative diseases: common themes and the unique case of amyotrophic lateral sclerosis. <i>Proteins: Structure, Function and Bioinformatics</i> , 2013, 81, 1285-30.  N-terminal helix-cap in Bhelix 2 modulates Btate misfolding in rabbit and hamster prion proteins. <i>PLoS ONE</i> , 2013, 8, e63047  Adaptor protein self-assembly drives the control of a cullin-RING ubiquitin ligase. <i>Structure</i> , 2012, 20, 1141-53  Conformation specificity and arene binding in a peptide composed only of Lys, Ile, Ala and Gly. <i>European Biophysics Journal</i> , 2012, 41, 63-72  An Arg-rich putative prebiotic protein is as stable as its Lys-rich variant. <i>Archives of Biochemistry and Biophysics</i> , 2012, 528, 118-26  Early steps in oxidation-induced SOD1 misfolding: implications for non-amyloid protein aggregation in familial ALS. <i>Journal of Molecular Biology</i> , 2012, 421, 631-52  Analyzing complicated protein folding kinetics rapidly by analytical Laplace inversion using a Tikhonov regularization variant. <i>Analytical Biochemistry</i> , 2012, 421, 181-90  Targeting of monomer/misfolded SOD1 as a therapeutic strategy for amyotrophic lateral sclerosis. <i>Journal of Neuroscience</i> , 2012, 32, 8791-9  Protein Misfolding and Toxicity in Amyotrophic Lateral Sclerosis 2012, 257-288  ALS-causing SOD1 mutations promote production of copper-deficient misfolded species. <i>Journal of Molecular Biology</i> , 2011, 409, 839-52  Cell surface binding and internalization of almodulated by degree of aggregation. <i>International Journal of Alzheimers Disease</i> , 2011, 2011, 962352  Interaction of Alzheimer Amyloid Peptide with Cell Surfaces and Artificial Membranes 2011, 231-243  Multiphoton ANS fluorescence microscopy as an in vivo sensor for protein misfolding stress. <i>Cell Stress and Chaperones</i> , 2011, 16, 549-61  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. <i>Biochemistry</i> , 2011, 205, 25056-64	Protein misfolding in the late-onset neurodegenerative diseases: common themes and the unique case of amyotrophic lateral sclerosis. <i>Proteins: Structure, Function and Bioinformatics, 2013, 81, 1285-303<sup>4-2</sup></i> N-terminal helix-cap in thelix 2 modulates listate misfolding in rabbit and hamster prion proteins. <i>PLoS ONE, 2013, 8, e63047</i> Adaptor protein self-assembly drives the control of a cullin-RING ubiquitin ligase. <i>Structure, 2012, 20, 1141-53</i> Conformation specificity and arene binding in a peptide composed only of Lys, Ile, Ala and Gly. <i>European Biophysics Journal, 2012, 41, 63-72</i> An Arg-rich putative prebiotic protein is as stable as its Lys-rich variant. <i>Archives of Biochemistry and Biophysics, 2012, 528, 118-26</i> Early steps in oxidation-induced SOD1 misfolding: implications for non-amyloid protein aggregation in familial ALS. <i>Journal of Molecular Biology, 2012, 421, 631-52</i> Analyzing complicated protein folding kinetics rapidly by analytical Laplace inversion using a Tikhonov regularization variant. <i>Analytical Biochemistry, 2012, 421, 181-90</i> 3.1  Targeting of monomer/misfolded SOD1 as a therapeutic strategy for amyotrophic lateral sclerosis. <i>Journal of Neuroscience, 2012, 32, 8791-9</i> Protein Misfolding and Toxicity in Amyotrophic Lateral Sclerosis 2012, 257-288  ALS-causing SOD1 mutations promote production of copper-deficient misfolded species. <i>Journal of Molecular Biology, 2011, 409, 839-52</i> Cell surface binding and Internalization of alinodulated by degree of aggregation. <i>International Journal of Alzheimens Disease, 2011, 10, 11, 962352</i> Interaction of Alzheimer Amyloid Peptide with Cell Surfaces and Artificial Membranes 2011, 231-243  Multiphoton ANS fluorescence microscopy as an in vivo sensor for protein misfolding stress. <i>Cell Stress and Chaperones, 2011, 16, 549-61</i> 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. <i>Biochemistry, 2011, </i>

56	Amyotrophic lateral sclerosis is a non-amyloid disease in which extensive misfolding of SOD1 is unique to the familial form. <i>Acta Neuropathologica</i> , <b>2010</b> , 119, 335-44	14.3	147
55	Putative one-pot prebiotic polypeptides with ribonucleolytic activity. <i>Chemistry - A European Journal</i> , <b>2010</b> , 16, 5314-23	4.8	11
54	Two distinct conformations of Abeta aggregates on the surface of living PC12 cells. <i>Biophysical Journal</i> , <b>2009</b> , 96, 4260-7	2.9	19
53	Probing Alzheimer amyloid peptide aggregation using a cell-free fluorescent protein refolding method. <i>Biochemistry and Cell Biology</i> , <b>2009</b> , 87, 631-9	3.6	7
52	Dimerization of the transmembrane domain of amyloid precursor proteins and familial Alzheimer's disease mutants. <i>BMC Neuroscience</i> , <b>2008</b> , 9, 17	3.2	61
51	Denaturational stress induces formation of zinc-deficient monomers of Cu,Zn superoxide dismutase: implications for pathogenesis in amyotrophic lateral sclerosis. <i>Journal of Molecular Biology</i> , <b>2008</b> , 383, 424-36	6.5	40
50	An immunological epitope selective for pathological monomer-misfolded SOD1 in ALS. <i>Nature Medicine</i> , <b>2007</b> , 13, 754-9	50.5	184
49	Requirement of aggregation propensity of Alzheimer amyloid peptides for neuronal cell surface binding. <i>BMC Neuroscience</i> , <b>2007</b> , 8, 29	3.2	29
48	Getting specificity from simplicity in putative proteins from the prebiotic earth. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2007</b> , 104, 14941-6	11.5	15
47	Species barriers for chronic wasting disease by in vitro conversion of prion protein. <i>Biochemical and Biophysical Research Communications</i> , <b>2007</b> , 364, 796-800	3.4	19
46	Structure, folding, and misfolding of Cu,Zn superoxide dismutase in amyotrophic lateral sclerosis. Biochimica Et Biophysica Acta - Molecular Basis of Disease, <b>2006</b> , 1762, 1025-37	6.9	147
45	NMR-driven secondary and tertiary structure model of Ca2+-loaded calexcitin. <i>Biochemical and Biophysical Research Communications</i> , <b>2006</b> , 343, 520-4	3.4	1
44	Variants of DsRed fluorescent protein: Development of a copper sensor. <i>Protein Science</i> , <b>2006</b> , 15, 2442	. <b>-</b> ₩.3	34
43	Characterization of segments from the central region of BRCA1: an intrinsically disordered scaffold for multiple protein-protein and protein-DNA interactions?. <i>Journal of Molecular Biology</i> , <b>2005</b> , 345, 275	5-8 <b>5</b>	141
42	Charge substitution shows that repulsive electrostatic interactions impede the oligomerization of Alzheimer amyloid peptides. <i>FEBS Letters</i> , <b>2005</b> , 579, 3574-8	3.8	46
41	Reversible assembly of helical filaments by de novo designed minimalist peptides. <i>Biopolymers</i> , <b>2005</b> , 80, 26-33	2.2	30
40	Alzheimer's Abeta40 studied by NMR at low pH reveals that sodium 4,4-dimethyl-4-silapentane-1-sulfonate (DSS) binds and promotes beta-ball oligomerization. <i>Journal of Biological Chemistry</i> , <b>2005</b> , 280, 3675-85	5.4	30
39	Hyperactive antifreeze protein from winter flounder is a very long rod-like dimer of alpha-helices. Journal of Biological Chemistry, <b>2005</b> , 280, 17920-9	5.4	67

## (2001-2004)

38	Monomeric Cu,Zn-superoxide dismutase is a common misfolding intermediate in the oxidation models of sporadic and familial amyotrophic lateral sclerosis. <i>Journal of Biological Chemistry</i> , <b>2004</b> , 279, 15499-504	5.4	263
37	Interaction of human and mouse Abeta peptides. Journal of Neurochemistry, 2004, 91, 1398-403	6	39
36	Reply to <b>P</b> roperties of a disease-specific prion probe□ <i>Nature Medicine</i> , <b>2004</b> , 10, 11-12	50.5	1
35	Interactions of Alzheimer amyloid peptides with cultured cells and brain tissue, and their biological consequences. <i>Biopolymers</i> , <b>2004</b> , 76, 4-14	2.2	16
34	Identification of stable helical bundles from a combinatorial library of amphipathic peptides. <i>Biopolymers</i> , <b>2004</b> , 76, 244-57	2.2	21
33	The PrP-like protein Doppel binds copper. <i>Journal of Biological Chemistry</i> , <b>2003</b> , 278, 8888-96	5.4	35
32	Co-incorporation of A beta 40 and A beta 42 to form mixed pre-fibrillar aggregates. <i>FEBS Journal</i> , <b>2003</b> , 270, 654-63		35
31	Lipopeptide detergents designed for the structural study of membrane proteins. <i>Nature Biotechnology</i> , <b>2003</b> , 21, 171-6	44.5	155
30	A prion protein epitope selective for the pathologically misfolded conformation. <i>Nature Medicine</i> , <b>2003</b> , 9, 893-9	50.5	233
29	Conformational coupling of Mg2+ and Ca2+ on the three-state folding of calexcitin B. <i>Biochemistry</i> , <b>2003</b> , 42, 5531-9	3.2	12
28	Alternate aggregation pathways of the Alzheimer beta-amyloid peptide: Abeta association kinetics at endosomal pH. <i>Journal of Molecular Biology</i> , <b>2003</b> , 325, 743-57	6.5	87
27	Autoinhibition of the kit receptor tyrosine kinase by the cytosolic juxtamembrane region. <i>Molecular and Cellular Biology</i> , <b>2003</b> , 23, 3067-78	4.8	133
26	Oxidation-induced misfolding and aggregation of superoxide dismutase and its implications for amyotrophic lateral sclerosis. <i>Journal of Biological Chemistry</i> , <b>2002</b> , 277, 47551-6	5.4	251
25	The molecular interaction of human salivary histatins with polyphenolic compounds. <i>FEBS Journal</i> , <b>2001</b> , 268, 4384-97		102
24	Alternate routes to conformational specificity in a Greek key beta barrel protein. <i>FEBS Journal</i> , <b>2001</b> , 268, 4653-63		2
23	All or none fibrillogenesis of a prion peptide. <i>FEBS Journal</i> , <b>2001</b> , 268, 4885-91		8
22	Alzheimer beta-amyloid peptides: structures of amyloid fibrils and alternate aggregation products. <i>Biopolymers</i> , <b>2001</b> , 60, 381-94	2.2	44
21	Calexcitin B is a new member of the sarcoplasmic calcium-binding protein family. <i>Journal of Biological Chemistry</i> , <b>2001</b> , 276, 22529-36	5.4	15

20	Progress in transthyretin fibrillogenesis research strengthens the amyloid hypothesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2001</b> , 98, 14757-9	11.5	7
19	Alzheimer Eamyloid peptides: Structures of amyloid fibrils and alternate aggregation products <b>2001</b> , 60, 381		1
18	Nonpolar contributions to conformational specificity in assemblies of designed short helical peptides. <i>Protein Science</i> , <b>2000</b> , 9, 1011-23	6.3	4
17	Alternate aggregation pathways of the Alzheimer beta-amyloid peptide. An in vitro model of preamyloid. <i>Journal of Biological Chemistry</i> , <b>2000</b> , 275, 36436-40	5.4	64
16	Structural studies of soluble oligomers of the Alzheimer beta-amyloid peptide. <i>Journal of Molecular Biology</i> , <b>2000</b> , 297, 73-87	6.5	201
15	Manipulating the amyloid-beta aggregation pathway with chemical chaperones. <i>Journal of Biological Chemistry</i> , <b>1999</b> , 274, 32970-4	5.4	201
14	Equilibrium folding intermediates of a Greek key beta-barrel protein. <i>Journal of Molecular Biology</i> , <b>1998</b> , 276, 669-81	6.5	21
13	Structural transitions associated with the interaction of Alzheimer beta-amyloid peptides with gangliosides. <i>Journal of Biological Chemistry</i> , <b>1998</b> , 273, 4506-15	5.4	142
12	Fibrillogenesis of Alzheimer Abeta peptides studied by fluorescence energy transfer. <i>Journal of Molecular Biology</i> , <b>1997</b> , 269, 214-24	6.5	61
11	Amyloid beta-protein (A beta) associated with lipid molecules: immunoreactivity distinct from that of soluble A beta. <i>FEBS Letters</i> , <b>1997</b> , 420, 43-6	3.8	37
10	Characterization of the interactions of Alzheimer beta-amyloid peptides with phospholipid membranes. <i>FEBS Journal</i> , <b>1997</b> , 245, 355-63		160
9	Membrane disruption by Alzheimer beta-amyloid peptides mediated through specific binding to either phospholipids or gangliosides. Implications for neurotoxicity. <i>Journal of Biological Chemistry</i> , <b>1996</b> , 271, 26482-9	5.4	267
8	Helix propagation and N-cap propensities of the amino acids measured in alanine-based peptides in 40 volume percent trifluoroethanol. <i>Protein Science</i> , <b>1996</b> , 5, 2623-37	6.3	239
7	Stability of EHelices. Advances in Protein Chemistry, 1995, 46, 141-176		301
6	Helix propensities of the amino acids measured in alanine-based peptides without helix-stabilizing side-chain interactions. <i>Protein Science</i> , <b>1994</b> , 3, 843-52	6.3	518
5	Determination of free energies of N-capping in alpha-helices by modification of the Lifson-Roig helix-coil therapy to include N- and C-capping. <i>Biochemistry</i> , <b>1994</b> , 33, 3396-403	3.2	163
4	Aromatic side-chain contribution to far-ultraviolet circular dichroism of helical peptides and its effect on measurement of helix propensities. <i>Biochemistry</i> , <b>1993</b> , 32, 5560-5	3.2	309
3	Large differences in the helix propensities of alanine and glycine. <i>Nature</i> , <b>1991</b> , 351, 586-8	50.4	305

The effect of enhanced alpha-helicity on the activity of a winter flounder antifreeze polypeptide. *FEBS Journal*, **1991**, 202, 1057-63

128

Primary structures of the alanine-rich antifreeze polypeptides from grubby sculpin, Myoxocephalus aenaeus. *Canadian Journal of Zoology*, **1988**, 66, 403-408

1.5 25