

# Chad A Dickey

## List of Publications by Year in descending order

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85  
papers

6,642  
citations

61984

43  
h-index

66911

78  
g-index

89  
all docs

89  
docs citations

89  
times ranked

7183  
citing authors

#	ARTICLE	IF	CITATIONS
1	The high-affinity HSP90-CHIP complex recognizes and selectively degrades phosphorylated tau client proteins. <i>Journal of Clinical Investigation</i> , 2007, 117, 648-658.	8.2	545
2	Hsp90-Tau Complex Reveals Molecular Basis for Specificity in Chaperone Action. <i>Cell</i> , 2014, 156, 963-974.	28.9	269
3	C9ORF72 poly(GA) aggregates sequester and impair HR23 and nucleocytoplasmic transport proteins. <i>Nature Neuroscience</i> , 2016, 19, 668-677.	14.8	268
4	Methylthioninium chloride (methylene blue) induces autophagy and attenuates tauopathy in vitro and in vivo. <i>Autophagy</i> , 2012, 8, 609-622.	9.1	260
5	Accelerated neurodegeneration through chaperone-mediated oligomerization of tau. <i>Journal of Clinical Investigation</i> , 2013, 123, 4158-4169.	8.2	246
6	Deletion of the Ubiquitin Ligase CHIP Leads to the Accumulation, But Not the Aggregation, of Both Endogenous Phospho- and Caspase-3-Cleaved Tau Species. <i>Journal of Neuroscience</i> , 2006, 26, 6985-6996.	3.6	234
7	Selectively Reduced Expression of Synaptic Plasticity-Related Genes in Amyloid Precursor Protein + Presenilin-1 Transgenic Mice. <i>Journal of Neuroscience</i> , 2003, 23, 5219-5226.	3.6	223
8	Chemical Manipulation of Hsp70 ATPase Activity Regulates Tau Stability. <i>Journal of Neuroscience</i> , 2009, 29, 12079-12088.	3.6	210
9	Tau Accumulation Activates the Unfolded Protein Response by Impairing Endoplasmic Reticulum-Associated Degradation. <i>Journal of Neuroscience</i> , 2013, 33, 9498-9507.	3.6	204
10	Akt and CHIP coregulate tau degradation through coordinated interactions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 3622-3627.	7.1	203
11	The Hsp90 Cochaperone, FKBP51, Increases Tau Stability and Polymerizes Microtubules. <i>Journal of Neuroscience</i> , 2010, 30, 591-599.	3.6	184
12	Allosteric Drugs: The Interaction of Antitumor Compound MKT-077 with Human Hsp70 Chaperones. <i>Journal of Molecular Biology</i> , 2011, 411, 614-632.	4.2	171
13	Phenothiazine-mediated rescue of cognition in tau transgenic mice requires neuroprotection and reduced soluble tau burden. <i>Molecular Neurodegeneration</i> , 2010, 5, 45.	10.8	160
14	HSP induction mediates selective clearance of tau phosphorylated at proline-directed Ser/Thr sites but not KXGS (MARK) sites. <i>FASEB Journal</i> , 2006, 20, 753-755.	0.5	157
15	DnaJ/Hsc70 chaperone complexes control the extracellular release of neurodegenerative-associated proteins. <i>EMBO Journal</i> , 2016, 35, 1537-1549.	7.8	154
16	Mapping interactions with the chaperone network reveals factors that protect against tau aggregation. <i>Nature Structural and Molecular Biology</i> , 2018, 25, 384-393.	8.2	119
17	Synthesis and Initial Evaluation of YM-08, a Blood-Brain Barrier Permeable Derivative of the Heat Shock Protein 70 (Hsp70) Inhibitor MKT-077, Which Reduces Tau Levels. <i>ACS Chemical Neuroscience</i> , 2013, 4, 930-939.	3.5	109
18	Phosphorylation Dynamics Regulate Hsp27-Mediated Rescue of Neuronal Plasticity Deficits in Tau Transgenic Mice. <i>Journal of Neuroscience</i> , 2010, 30, 15374-15382.	3.6	105

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19	A New Anti-Depressive Strategy for the Elderly: Ablation of FKBP5/FKBP51. PLoS ONE, 2011, 6, e24840.	2.5	105
20	Imbalance of Hsp70 family variants fosters tau accumulation. FASEB Journal, 2013, 27, 1450-1459.	0.5	100
21	Allosteric Heat Shock Protein 70 Inhibitors Rapidly Rescue Synaptic Plasticity Deficits by Reducing Aberrant Tau. Biological Psychiatry, 2013, 74, 367-374.	1.3	93
22	Number of A $\beta$ Inoculations in APP+PS1 Transgenic Mice Influences Antibody Titers, Microglial Activation, and Congophilic Plaque Levels. DNA and Cell Biology, 2001, 20, 731-736.	1.9	90
23	Hsp90 activator Aha1 drives production of pathological tau aggregates. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 9707-9712.	7.1	89
24	Targeting Hsp90 and its co-chaperones to treat Alzheimer's disease. Expert Opinion on Therapeutic Targets, 2014, 18, 1219-1232.	3.4	86
25	Cysteine Reactivity Distinguishes Redox Sensing by the Heat-Inducible and Constitutive Forms of Heat Shock Protein 70. Chemistry and Biology, 2012, 19, 1391-1399.	6.0	83
26	The emerging role of peptidylprolyl isomerase chaperones in tau oligomerization, amyloid processing, and Alzheimer's disease. Journal of Neurochemistry, 2015, 133, 1-13.	3.9	81
27	Chaperones in Neurodegeneration. Journal of Neuroscience, 2015, 35, 13853-13859.	3.6	81
28	Amyloid suppresses induction of genes critical for memory consolidation in APP+PS1 transgenic mice. Journal of Neurochemistry, 2004, 88, 434-442.	3.9	80
29	Age-Associated Epigenetic Upregulation of the FKBP5 Gene Selectively Impairs Stress Resiliency. PLoS ONE, 2014, 9, e107241.	2.5	79
30	Development of a High Throughput Drug Screening Assay for the Detection of Changes in Tau Levels - Proof of Concept with HSP90 inhibitors. Current Alzheimer Research, 2005, 2, 231-238.	1.4	77
31	Hsc70 Rapidly Engages Tau after Microtubule Destabilization. Journal of Biological Chemistry, 2010, 285, 16798-16805.	3.4	75
32	Aging Analysis Reveals Slowed Tau Turnover and Enhanced Stress Response in a Mouse Model of Tauopathy. American Journal of Pathology, 2009, 174, 228-238.	3.8	73
33	Analysis of the Tau-Associated Proteome Reveals That Exchange of Hsp70 for Hsp90 Is Involved in Tau Degradation. ACS Chemical Biology, 2012, 7, 1677-1686.	3.4	72
34	Structure and pro-toxic mechanism of the human Hsp90/PPIase/Tau complex. Nature Communications, 2018, 9, 4532.	12.8	68
35	Glucose-regulated Protein 94 Triage of Mutant Myocilin through Endoplasmic Reticulum-associated Degradation Subverts a More Efficient Autophagic Clearance Mechanism. Journal of Biological Chemistry, 2012, 287, 40661-40669.	3.4	66
36	Dysregulation of Na <sup>+</sup> /K <sup>+</sup> ATPase by amyloid in APP+PS1 transgenic mice. BMC Neuroscience, 2005, 6, 7.	1.9	59

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37	The Hsp90 Kinase Co-chaperone Cdc37 Regulates Tau Stability and Phosphorylation Dynamics. <i>Journal of Biological Chemistry</i> , 2011, 286, 16976-16983.	3.4	59
38	Stabilizing the Hsp70-Tau Complex Promotes Turnover in Models of Tauopathy. <i>Cell Chemical Biology</i> , 2016, 23, 992-1001.	5.2	58
39	DnaJ1 Antagonizes Constitutive Hsp70-Mediated Stabilization of Tau. <i>Journal of Molecular Biology</i> , 2012, 421, 653-661.	4.2	56
40	Cellular factors modulating the mechanism of tau protein aggregation. <i>Cellular and Molecular Life Sciences</i> , 2015, 72, 1863-1879.	5.4	55
41	Molecular chaperones and regulation of tau quality control: strategies for drug discovery in tauopathies. <i>Future Medicinal Chemistry</i> , 2011, 3, 1523-1537.	2.3	54
42	Development of Glucose Regulated Protein 94-Selective Inhibitors Based on the Bnlm and Radamide Scaffold. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 3471-3488.	6.4	54
43	MicroRNA-511 Binds to FKBP5 mRNA, Which Encodes a Chaperone Protein, and Regulates Neuronal Differentiation. <i>Journal of Biological Chemistry</i> , 2016, 291, 17897-17906.	3.4	46
44	Duration and Specificity of Humoral Immune Responses in Mice Vaccinated with the Alzheimer's Disease-Associated $\beta$ -Amyloid 1-42 Peptide. <i>DNA and Cell Biology</i> , 2001, 20, 723-729.	1.9	43
45	Human cyclophilin 40 unravels neurotoxic amyloids. <i>PLoS Biology</i> , 2017, 15, e2001336.	5.6	43
46	Targeting the ER-autophagy system in the trabecular meshwork to treat glaucoma. <i>Experimental Eye Research</i> , 2016, 144, 38-45.	2.6	42
47	Rhodacyanine Derivative Selectively Targets Cancer Cells and Overcomes Tamoxifen Resistance. <i>PLoS ONE</i> , 2012, 7, e35566.	2.5	40
48	Hsp70 ATPase Modulators as Therapeutics for Alzheimer's and other Neurodegenerative Diseases. <i>Molecular and Cellular Pharmacology</i> , 2010, 2, 43-46.	1.7	40
49	Isoform-selective Genetic Inhibition of Constitutive Cytosolic Hsp70 Activity Promotes Client Tau Degradation Using an Altered Co-chaperone Complement. <i>Journal of Biological Chemistry</i> , 2015, 290, 13115-13127.	3.4	39
50	Exploiting the interaction between Grp94 and aggregated myocilin to treat glaucoma. <i>Human Molecular Genetics</i> , 2014, 23, 6470-6480.	2.9	38
51	Transformation of the Non-Selective Aminocyclohexanol-Based Hsp90 Inhibitor into a Grp94-Selective Scaffold. <i>ACS Chemical Biology</i> , 2017, 12, 244-253.	3.4	38
52	Pharmacologic reductions of total tau levels; implications for the role of microtubule dynamics in regulating tau expression. <i>Molecular Neurodegeneration</i> , 2006, 1, 6.	10.8	35
53	Identification of dihydropyridines that reduce cellular tau levels. <i>Chemical Communications</i> , 2011, 47, 529-531.	4.1	35
54	The Disease-Associated Chaperone FKBP51 Impairs Cognitive Function by Accelerating AMPA Receptor Recycling. <i>ENeuro</i> , 2019, 6, ENEURO.0242-18.2019.	1.9	35

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55	Bending Tau into Shape: The Emerging Role of Peptidyl-Prolyl Isomerases in Tauopathies. <i>Molecular Neurobiology</i> , 2011, 44, 65-70.	4.0	30
56	Targeting the FKBP51/GR/Hsp90 Complex to Identify Functionally Relevant Treatments for Depression and PTSD. <i>ACS Chemical Biology</i> , 2018, 13, 2288-2299.	3.4	29
57	Spermidine/spermine-N1-acetyltransferase ablation impacts tauopathy-induced polyamine stress response. <i>Alzheimer's Research and Therapy</i> , 2019, 11, 58.	6.2	29
58	The active Hsc70/tau complex can be exploited to enhance tau turnover without damaging microtubule dynamics. <i>Human Molecular Genetics</i> , 2015, 24, 3971-3981.	2.9	28
59	Isoform-selective Hsp90 inhibition rescues model of hereditary open-angle glaucoma. <i>Scientific Reports</i> , 2017, 7, 17951.	3.3	28
60	Early Life Stress and High FKBP5 Interact to Increase Anxiety-Like Symptoms through Altered AKT Signaling in the Dorsal Hippocampus. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2738.	4.1	28
61	Potential synergy between tau aggregation inhibitors and tau chaperone modulators. <i>Alzheimer's Research and Therapy</i> , 2013, 5, 41.	6.2	25
62	Enhanced tau pathology via RanBP9 and Hsp90/Hsc70 chaperone complexes. <i>Human Molecular Genetics</i> , 2017, 26, 3973-3988.	2.9	24
63	Current strategies for the treatment of Alzheimer's disease and other tauopathies. <i>Expert Opinion on Therapeutic Targets</i> , 2006, 10, 665-676.	3.4	22
64	The role of FKBP5 in mood disorders: action of FKBP5 on steroid hormone receptors leads to questions about its evolutionary importance. <i>CNS and Neurological Disorders - Drug Targets</i> , 2013, 12, 1157-62.	1.4	22
65	Reconstructing the Hsp90/Tau Machine. <i>Current Enzyme Inhibition</i> , 2013, 9, 41-45.	0.4	20
66	Aberrant AZIN2 and polyamine metabolism precipitates tau neuropathology. <i>Journal of Clinical Investigation</i> , 2021, 131, .	8.2	20
67	FKBP5 and early life stress affect the hippocampus by an age-dependent mechanism. <i>Brain, Behavior, &amp; Immunity - Health</i> , 2020, 9, 100143.	2.5	19
68	Hippocampal Neurogenesis Is Enhanced in Adult Tau Deficient Mice. <i>Cells</i> , 2020, 9, 210.	4.1	19
69	Small heat shock protein 22 kDa can modulate the aggregation and liquid-liquid phase separation behavior of tau. <i>Protein Science</i> , 2021, 30, 1350-1359.	7.6	19
70	Neurodegeneration and the Heat Shock Protein 70 Machinery: Implications for Therapeutic Development. <i>Current Topics in Medicinal Chemistry</i> , 2016, 16, 2741-2752.	2.1	19
71	Synthesis, Stereochemical Analysis, and Derivatization of Myricanol Provide New Probes That Promote Autophagic Tau Clearance. <i>ACS Chemical Biology</i> , 2015, 10, 1099-1109.	3.4	18
72	The Metamorphic Nature of the Tau Protein: Dynamic Flexibility Comes at a Cost. <i>Frontiers in Neuroscience</i> , 2016, 10, 3.	2.8	18

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73	Protein Cross-Linking Capillary Electrophoresis for Protein-Protein Interaction Analysis. <i>Analytical Chemistry</i> , 2016, 88, 8272-8278.	6.5	18
74	Induction of memory-associated immediate early genes by nerve growth factor in rat primary cortical neurons and differentiated mouse Neuro2A cells. <i>Neuroscience Letters</i> , 2004, 366, 10-14.	2.1	17
75	Trifunctional High-Throughput Screen Identifies Promising Scaffold To Inhibit Grp94 and Treat Myocilin-Associated Glaucoma. <i>ACS Chemical Biology</i> , 2018, 13, 933-941.	3.4	17
76	Inhibition of Both Hsp70 Activity and Tau Aggregation <i>in Vitro</i> Best Predicts Tau Lowering Activity of Small Molecules. <i>ACS Chemical Biology</i> , 2016, 11, 2041-2048.	3.4	14
77	Stable calcium-free myocilin olfactomedin domain variants reveal challenges in differentiating between benign and glaucoma-causing mutations. <i>Journal of Biological Chemistry</i> , 2019, 294, 12717-12728.	3.4	13
78	Repeated repeat problems: Combinatorial effect of C9orf72-derived dipeptide repeat proteins. <i>International Journal of Biological Macromolecules</i> , 2019, 127, 136-145.	7.5	13
79	Management of Hsp90-Dependent Protein Folding by Small Molecules Targeting the Aha1 Co-Chaperone. <i>Cell Chemical Biology</i> , 2020, 27, 292-305.e6.	5.2	13
80	Hsp22 with an N-Terminal Domain Truncation Mediates a Reduction in Tau Protein Levels. <i>International Journal of Molecular Sciences</i> , 2020, 21, 5442.	4.1	10
81	FKBP52 overexpression accelerates hippocampal-dependent memory impairments in a tau transgenic mouse model. <i>Npj Aging and Mechanisms of Disease</i> , 2021, 7, 9.	4.5	10
82	The Earliest Tau Dysfunction in Alzheimer's Disease?. <i>American Journal of Pathology</i> , 2011, 179, 2148-2151.	3.8	8
83	Neuronal Life Span Versus Health Span: Principles of Natural Selection at Work in the Degenerating Brain. <i>Journal of Molecular Neuroscience</i> , 2011, 45, 467-72.	2.3	2
84	Commentary on "Cytoskeletal modulators and pleiotropic strategies for Alzheimer drug discovery." The last stand: The dichotomy of chaperone function in Alzheimer's disease. , 2007, 3, 3-6.		1
85	S3-02-01: TARGETING CHAPERONES TO TREAT FTLD. , 2014, 10, P201-P201.		0