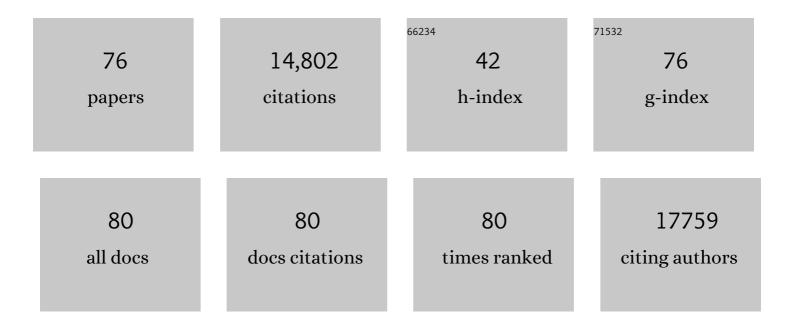
Erich E Wanker

List of Publications by Year in descending order

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#	Article	lF	CITATIONS
1	A Human Protein-Protein Interaction Network: A Resource for Annotating the Proteome. Cell, 2005, 122, 957-968.	13.5	2,169
2	Formation of Neuronal Intranuclear Inclusions Underlies the Neurological Dysfunction in Mice Transgenic for the HD Mutation. Cell, 1997, 90, 537-548.	13.5	2,105
3	EGCG redirects amyloidogenic polypeptides into unstructured, off-pathway oligomers. Nature Structural and Molecular Biology, 2008, 15, 558-566.	3.6	1,249
4	Huntingtin-Encoded Polyglutamine Expansions Form Amyloid-like Protein Aggregates In Vitro and In Vivo. Cell, 1997, 90, 549-558.	13.5	1,224
5	EGCG remodels mature α-synuclein and amyloid-β fibrils and reduces cellular toxicity. Proceedings of the United States of America, 2010, 107, 7710-7715.	3.3	888
6	An empirical framework for binary interactome mapping. Nature Methods, 2009, 6, 83-90.	9.0	800
7	Accumulation of Mutant Huntingtin Fragments in Aggresome-like Inclusion Bodies as a Result of Insufficient Protein Degradation. Molecular Biology of the Cell, 2001, 12, 1393-1407.	0.9	583
8	The hunt for huntingtin function: interaction partners tell many different stories. Trends in Biochemical Sciences, 2003, 28, 425-433.	3.7	456
9	Small-molecule conversion of toxic oligomers to nontoxic β-sheet–rich amyloid fibrils. Nature Chemical Biology, 2012, 8, 93-101.	3.9	400
10	A Protein Interaction Network Links GIT1, an Enhancer of Huntingtin Aggregation, to Huntington's Disease. Molecular Cell, 2004, 15, 853-865.	4.5	398
11	Green tea (â^')-epigallocatechin-gallate modulates early events in huntingtin misfolding and reduces toxicity in Huntington's disease models. Human Molecular Genetics, 2006, 15, 2743-2751.	1.4	357
12	A Directed Protein Interaction Network for Investigating Intracellular Signal Transduction. Science Signaling, 2011, 4, rs8.	1.6	313
13	HIPPIE: Integrating Protein Interaction Networks with Experiment Based Quality Scores. PLoS ONE, 2012, 7, e31826.	1.1	297
14	[24] Membrane filter assay for detection of amyloid-like polyglutamine-containing protein aggregates. Methods in Enzymology, 1999, 309, 375-386.	0.4	217
15	SH3GL3 Associates with the Huntingtin Exon 1 Protein and Promotes the Formation of Polygln-Containing Protein Aggregates. Molecular Cell, 1998, 2, 427-436.	4.5	208
16	Identification of benzothiazoles as potential polyglutamine aggregation inhibitors of Huntington's disease by using an automated filter retardation assay. Proceedings of the National Academy of Sciences of the United States of America, 2002, 99, 16400-16406.	3.3	199
17	The green tea polyphenol (â^')â€epigallocatechin gallate prevents the aggregation of tau protein into toxic oligomers at substoichiometric ratios. FEBS Letters, 2015, 589, 77-83.	1.3	172
18	Evolution and function of CAG/polyglutamine repeats in protein–protein interaction networks. Nucleic Acids Research, 2012, 40, 4273-4287.	6.5	166

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19	An arginine/lysine-rich motif is crucial for VCP/p97-mediated modulation of ataxin-3 fibrillogenesis. EMBO Journal, 2006, 25, 1547-1558.	3.5	142
20	Protein Aggregation and Pathogenesis of Huntingtons Disease: Mechanisms and Correlations. Biological Chemistry, 2000, 381, 937-942.	1.2	134
21	Human iPSC-Derived Neural Progenitors Are an Effective Drug Discovery Model for Neurological mtDNA Disorders. Cell Stem Cell, 2017, 20, 659-674.e9.	5.2	126
22	Mutant Huntingtin Promotes the Fibrillogenesis of Wild-type Huntingtin. Journal of Biological Chemistry, 2003, 278, 41452-41461.	1.6	107
23	Ataxin-3 is transported into the nucleus and associates with the nuclear matrix. Human Molecular Genetics, 1998, 7, 991-997.	1.4	104
24	The value of high quality protein–protein interaction networks for systems biology. Current Opinion in Chemical Biology, 2006, 10, 551-558.	2.8	100
25	A Y2H-seq approach defines the human protein methyltransferase interactome. Nature Methods, 2013, 10, 339-342.	9.0	99
26	Amyloid-β(1–42) Aggregation Initiates Its Cellular Uptake and Cytotoxicity. Journal of Biological Chemistry, 2016, 291, 19590-19606.	1.6	91
27	Quantitative Interaction Proteomics of Neurodegenerative Disease Proteins. Cell Reports, 2015, 11, 1134-1146.	2.9	88
28	Translation of HTT mRNA with expanded CAG repeats is regulated by the MID1–PP2A protein complex. Nature Communications, 2013, 4, 1511.	5.8	84
29	Epigallocatechin-3-gallate: a useful, effective and safe clinical approach for targeted prevention and individualised treatment of neurological diseases?. EPMA Journal, 2013, 4, 5.	3.3	80
30	The pathobiology of perturbed mutant huntingtin protein–protein interactions in Huntington's disease. Journal of Neurochemistry, 2019, 151, 507-519.	2.1	70
31	Interactome Mapping Provides a Network of Neurodegenerative Disease Proteins and Uncovers Widespread Protein Aggregation in Affected Brains. Cell Reports, 2020, 32, 108050.	2.9	64
32	Metformin reverses early cortical network dysfunction and behavior changes in Huntington's disease. ELife, 2018, 7, .	2.8	64
33	Detection of Alpha-Rod Protein Repeats Using a Neural Network and Application to Huntingtin. PLoS Computational Biology, 2009, 5, e1000304.	1.5	59
34	Modulation of human IAPP fibrillation: cosolutes, crowders and chaperones. Physical Chemistry Chemical Physics, 2015, 17, 8338-8348.	1.3	59
35	UniHl 4: new tools for query, analysis and visualization of the human protein–protein interactome. Nucleic Acids Research, 2009, 37, D657-D660.	6.5	58
36	The palmitoyl acyltransferase HIP14 shares a high proportion of interactors with huntingtin: implications for a role in the pathogenesis of Huntington's disease. Human Molecular Genetics, 2014, 23, 4142-4160.	1.4	58

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37	Maximizing binary interactome mapping with a minimal number of assays. Nature Communications, 2019, 10, 3907.	5.8	57
38	Defective metabolic programming impairs early neuronal morphogenesis in neural cultures and an organoid model of Leigh syndrome. Nature Communications, 2021, 12, 1929.	5.8	55
39	mHTT Seeding Activity: A Marker of Disease Progression and Neurotoxicity in Models of Huntington's Disease. Molecular Cell, 2018, 71, 675-688.e6.	4.5	50
40	Inhibition of Huntingtin Exon-1 Aggregation by the Molecular Tweezer CLR01. Journal of the American Chemical Society, 2017, 139, 5640-5643.	6.6	49
41	Identification of VCP/p97, Carboxyl Terminus of Hsp70-interacting Protein (CHIP), and Amphiphysin II Interaction Partners Using Membrane-based Human Proteome Arrays. Molecular and Cellular Proteomics, 2006, 5, 234-244.	2.5	48
42	Prion-like proteins sequester and suppress the toxicity of huntingtin exon 1. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 12085-12090.	3.3	47
43	Phosphorylation-regulated axonal dependent transport of syntaxin 1 is mediated by a Kinesin-1 adapter. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 5862-5867.	3.3	44
44	An integer programming framework for inferring disease complexes from network data. Bioinformatics, 2016, 32, i271-i277.	1.8	44
45	Multiplex approaches in protein microarray technology. Expert Review of Proteomics, 2005, 2, 499-510.	1.3	39
46	Aggregation of polyQâ€extended proteins is promoted by interaction with their natural coiledâ€coil partners. BioEssays, 2013, 35, 503-507.	1.2	39
47	Mixing Aβ(1–40) and Aβ(1–42) peptides generates unique amyloid fibrils. Chemical Communications, 2020, 56, 8830-8833.	2.2	39
48	Aggregation of Full-length Immunoglobulin Light Chains from Systemic Light Chain Amyloidosis (AL) Patients Is Remodeled by Epigallocatechin-3-gallate. Journal of Biological Chemistry, 2017, 292, 2328-2344.	1.6	37
49	Quantitative interaction mapping reveals an extended UBX domain in ASPL that disrupts functional p97 hexamers. Nature Communications, 2016, 7, 13047.	5.8	35
50	Self-assembly of Mutant Huntingtin Exon-1 Fragments into Large Complex Fibrillar Structures Involves Nucleated Branching. Journal of Molecular Biology, 2018, 430, 1725-1744.	2.0	35
51	Subcellular Localization And Formation Of Huntingtin Aggregates Correlates With Symptom Onset And Progression In A Huntington'S Disease Model. Brain Communications, 2020, 2, fcaa066.	1.5	34
52	Lu <scp>TH</scp> y: a doubleâ€readout bioluminescenceâ€based twoâ€hybrid technology for quantitative mapping of protein–protein interactions in mammalian cells. Molecular Systems Biology, 2018, 14, e8071.	3.2	31
53	Identification of Human Proteins That Modify Misfolding and Proteotoxicity of Pathogenic Ataxin-1. PLoS Genetics, 2012, 8, e1002897.	1.5	29
54	DULIP: A Dual Luminescence-Based Co-Immunoprecipitation Assay for Interactome Mapping in Mammalian Cells. Journal of Molecular Biology, 2015, 427, 3375-3388.	2.0	28

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55	Current Approaches Toward Quantitative Mapping of the Interactome. Frontiers in Genetics, 2016, 7, 74.	1.1	28
56	Interleukinâ€12/23 deficiency differentially affects pathology in male and female Alzheimer's diseaseâ€like mice. EMBO Reports, 2020, 21, e48530.	2.0	24
57	The E3 Ubiquitin Ligase MID1 Catalyzes Ubiquitination and Cleavage of Fu. Journal of Biological Chemistry, 2014, 289, 31805-31817.	1.6	23
58	Development and application of a DNA microarray-based yeast two-hybrid system. Nucleic Acids Research, 2013, 41, 1496-1507.	6.5	19
59	Identification of the Mitochondrial MSRB2 as a Binding Partner of LG72. Cellular and Molecular Neurobiology, 2014, 34, 1123-1130.	1.7	18
60	DCAF8, a novel MuRF1 interaction partner, promotes muscle atrophy. Journal of Cell Science, 2019, 132, .	1.2	17
61	Shedding a new light on Huntington's disease: how blood can both propagate and ameliorate disease pathology. Molecular Psychiatry, 2021, 26, 5441-5463.	4.1	16
62	Dynamics of huntingtin protein interactions in the striatum identifies candidate modifiers of Huntington disease. Cell Systems, 2022, 13, 304-320.e5.	2.9	15
63	Pathogenic Polyglutamine Tracts Are Potent Inducers of Spontaneous Sup35 and Rnq1 Amyloidogenesis. PLoS ONE, 2010, 5, e9642.	1.1	14
64	Identification of an RNA Polymerase III Regulator Linked to Disease-Associated Protein Aggregation. Molecular Cell, 2017, 65, 1096-1108.e6.	4.5	14
65	Sclerotiorin Stabilizes the Assembly of Nonfibrillar Abeta42 Oligomers with Low Toxicity, Seeding Activity, and Beta-sheet Content. Journal of Molecular Biology, 2020, 432, 2080-2098.	2.0	12
66	Spontaneous self-assembly of pathogenic huntingtin exon 1 protein into amyloid structures. Essays in Biochemistry, 2014, 56, 167-180.	2.1	12
67	FEZ1 Forms Complexes with CRMP1 and DCC to Regulate Axon and Dendrite Development. ENeuro, 2021, 8, ENEURO.0193-20.2021.	0.9	11
68	A functionally defined high-density NRF2 interactome reveals new conditional regulators of ARE transactivation. Redox Biology, 2020, 37, 101686.	3.9	10
69	The Anti-amyloid Compound DO1 Decreases Plaque Pathology and Neuroinflammation-Related Expression Changes in 5xFAD Transgenic Mice. Cell Chemical Biology, 2019, 26, 109-120.e7.	2.5	8
70	Small, Seeding-Competent Huntingtin Fibrils Are Prominent Aggregate Species in Brains of zQ175 Huntington's Disease Knock-in Mice. Frontiers in Neuroscience, 2021, 15, 682172.	1.4	7
71	Assessment of Ethanol-Induced Toxicity on iPSC-Derived Human Neurons Using a Novel High-Throughput Mitochondrial Neuronal Health (MNH) Assay. Frontiers in Cell and Developmental Biology, 2020, 8, 590540.	1.8	6
72	Schizophrenia risk candidate protein ZNF804A interacts with STAT2 and influences interferon-mediated gene transcription in mammalian cells. Journal of Molecular Biology, 2021, 433, 167184.	2.0	6

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73	A Filter Retardation Assay Facilitates the Detection and Quantification of Heat-Stable, Amyloidogenic Mutant Huntingtin Aggregates in Complex Biosamples. Methods in Molecular Biology, 2018, 1780, 31-40.	0.4	5
74	Flexible web-based integration of distributed large-scale human protein interaction maps. Journal of Integrative Bioinformatics, 2007, 4, 40-50.	1.0	3
75	CellFIE: CRISPR- and Cell Fusion-based Two-hybrid Interaction Mapping of Endogenous Proteins. Journal of Molecular Biology, 2021, 433, 167305.	2.0	Ο
76	Klinische Proteomik. , 2008, , 297-313.		0