

Alexander Tischer

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/2351145/alexander-tischer-publications-by-year.pdf>

Version: 2024-04-27

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.

41
papers

588
citations

12
h-index

23
g-index

44
ext. papers

755
ext. citations

5.1
avg, IF

3.66
L-index

#	Paper	IF	Citations
41	Quantification of von Willebrand factor and ADAMTS-13 after traumatic injury: a pilot study. <i>Trauma Surgery and Acute Care Open</i> , 2021 , 6, e000703	2.4	0
40	MeV-Stealth: A CD46-specific oncolytic measles virus resistant to neutralization by measles-immune human serum. <i>PLoS Pathogens</i> , 2021 , 17, e1009283	7.6	5
39	Functional succinate dehydrogenase deficiency is a common adverse feature of clear cell renal cancer. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021 , 118,	11.5	1
38	Measuring Lipid Transfer Protein Activity Using Bicelle-Dilution Model Membranes. <i>Analytical Chemistry</i> , 2020 , 92, 3417-3425	7.8	3
37	Evidence for the Misfolding of the A1 Domain within Multimeric von Willebrand Factor in Type 2 von Willebrand Disease. <i>Journal of Molecular Biology</i> , 2020 , 432, 305-323	6.5	4
36	Glycosylation sterically inhibits platelet adhesion to von Willebrand factor without altering intrinsic conformational dynamics. <i>Journal of Thrombosis and Haemostasis</i> , 2020 , 18, 79-90	15.4	6
35	Modulating the rate of fibrin formation and clot structure attenuates microvascular thrombosis in systemic inflammation. <i>Blood Advances</i> , 2020 , 4, 1340-1349	7.8	6
34	Platelet-type von Willebrand disease: Local disorder of the platelet GPIb α switch drives high-affinity binding to von Willebrand factor. <i>Journal of Thrombosis and Haemostasis</i> , 2019 , 17, 2022-2034	15.4	0
33	Arabinose Alters Both Local and Distal H-D Exchange Rates in the Escherichia coli AraC Transcriptional Regulator. <i>Biochemistry</i> , 2019 , 58, 2875-2882	3.2	2
32	Ascorbic acid-induced TET activation mitigates adverse hydroxymethylcytosine loss in renal cell carcinoma. <i>Journal of Clinical Investigation</i> , 2019 , 129, 1612-1625	15.9	47
31	Functional Interrogation of Variants of Undetermined Significance of the Isocitrate Dehydrogenase 1 and 2 Genes in Myeloid Neoplasms. <i>Blood</i> , 2019 , 134, 1697-1697	2.2	
30	Functional evaluation of isocitrate dehydrogenase 1 and 2 variants of unclear significance in chronic myeloid neoplasms. <i>Leukemia Research</i> , 2019 , 87, 106264	2.7	
29	The von Willebrand factor Tyr2561 allele is a gain-of-function variant and a risk factor for early myocardial infarction. <i>Blood</i> , 2019 , 133, 356-365	2.2	15
28	"Cooperative collapse" of the denatured state revealed through Clausius-Clapeyron analysis of protein denaturation phase diagrams. <i>Biopolymers</i> , 2018 , 109, e23106	2.2	2
27	Mayo CALR mutation type classification guide using alpha helix propensity. <i>American Journal of Hematology</i> , 2018 , 93, E128-E129	7.1	11
26	Sequence Reversal Prevents Chain Collapse and Yields Heat-Sensitive Intrinsic Disorder. <i>Biophysical Journal</i> , 2018 , 115, 328-340	2.9	5
25	The Von Willebrand Factor Tyr2561 Allele Is a Gain-of-Function Variant and a Potential Risk Factor for Early Myocardial Infarction. <i>Blood</i> , 2018 , 132, 2459-2459	2.2	1

24	The Chaperonin GroEL: A Versatile Tool for Applied Biotechnology Platforms. <i>Frontiers in Molecular Biosciences</i> , 2018 , 5, 46	5.6	6
23	A Novel Kleefstra Syndrome-associated Variant That Affects the Conserved TPL Motif within the Ankyrin Repeat of EHMT1 Leads to Abnormal Protein Folding. <i>Journal of Biological Chemistry</i> , 2017 , 292, 3866-3876	5.4	11
22	Enhanced Local Disorder in a Clinically Elusive von Willebrand Factor Provokes High-Affinity Platelet Clumping. <i>Journal of Molecular Biology</i> , 2017 , 429, 2161-2177	6.5	11
21	The Von Willebrand Factor A1-Collagen III Interaction Is Independent of Conformation and Type 2 Von Willebrand Disease Phenotype. <i>Journal of Molecular Biology</i> , 2017 , 429, 32-47	6.5	7
20	Mutational Constraints on Local Unfolding Inhibit the Rheological Adaptation of von Willebrand Factor. <i>Journal of Biological Chemistry</i> , 2016 , 291, 3848-59	5.4	18
19	Data on the purification and crystallization of the loss-of-function von Willebrand disease variant (p.Gly1324Ser) of the von Willebrand factor A1 domain. <i>Data in Brief</i> , 2016 , 7, 1700-1706	1.2	6
18	Chaperonin-Based Biolayer Interferometry To Assess the Kinetic Stability of Metastable, Aggregation-Prone Proteins. <i>Biochemistry</i> , 2016 , 55, 4885-908	3.2	5
17	Structural origins of misfolding propensity in the platelet adhesive von Willebrand factor A1 domain. <i>Biophysical Journal</i> , 2015 , 109, 398-406	2.9	10
16	Thermodynamic and fibril formation studies of full length immunoglobulin light chain AL-09 and its germline protein using scan rate dependent thermal unfolding. <i>Biophysical Chemistry</i> , 2015 , 207, 13-20	3.5	34
15	The effects of N-ethyl-N'-methyl imidazolium chloride on the solubility, stability and aggregation of tc-rPA. <i>FEBS Journal</i> , 2014 , 281, 1738-49	5.7	7
14	Accessibilities of N-terminal myristoyl chain and cysteines in guanylyl cyclase-activating protein-2 (GCAP-2) studied by covalent labeling and mass spectrometry. <i>Rapid Communications in Mass Spectrometry</i> , 2014 , 28, 835-8	2.2	1
13	Kinetic control in protein folding for light chain amyloidosis and the differential effects of somatic mutations. <i>Journal of Molecular Biology</i> , 2014 , 426, 347-61	6.5	41
12	A molten globule intermediate of the von Willebrand factor A1 domain firmly tethers platelets under shear flow. <i>Proteins: Structure, Function and Bioinformatics</i> , 2014 , 82, 867-78	4.2	16
11	Misfolding of vWF to pathologically disordered conformations impacts the severity of von Willebrand disease. <i>Biophysical Journal</i> , 2014 , 107, 1185-1195	2.9	27
10	Alanine and proline content modulate global sensitivity to discrete perturbations in disordered proteins. <i>Proteins: Structure, Function and Bioinformatics</i> , 2014 , 82, 3373-84	4.2	12
9	Oxidative refolding of rPA in l-ArgHCl and in ionic liquids: A correlation between hydrophobicity, salt effects, and refolding yield. <i>Biopolymers</i> , 2014 , 101, 1129-40	2.2	1
8	The prognostic advantage of calreticulin mutations in myelofibrosis might be confined to type 1 or type 1-like CALR variants. <i>Blood</i> , 2014 , 124, 2465-6	2.2	105
7	Misfolding Induced By the Mutations V1314D and F1369I Affects the Function and the Structure of the Von Willebrand A1-Domain. <i>Blood</i> , 2014 , 124, 1521-1521	2.2	

6	The Prognostic Advantage of Calreticulin Mutations in Myelofibrosis Might be Confined to Type 1 or Type 1-like Calreticulin Variants. <i>Blood</i> , 2014 , 124, 3166-3166	2.2	
5	Urea-temperature phase diagrams capture the thermodynamics of denatured state expansion that accompany protein unfolding. <i>Protein Science</i> , 2013 , 22, 1147-60	6.3	13
4	The linker between the D3 and A1 domains of vWF suppresses A1-GPIb catch bonds by site-specific binding to the A1 domain. <i>Protein Science</i> , 2013 , 22, 1049-59	6.3	26
3	Ionic liquids as refolding additives: variation of the anion. <i>Journal of Biotechnology</i> , 2010 , 150, 64-72	3.7	82
2	L-arginine hydrochloride increases the solubility of folded and unfolded recombinant plasminogen activator rPA. <i>Protein Science</i> , 2010 , 19, 1783-95	6.3	39
1	Functional succinate dehydrogenase deficiency and loss of ascorbic acid transporter SLC23A1 are pathognomonic adverse features of clear cell renal cancer		2