

Alexander Tischer

List of Publications by Citations

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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 | 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 |
| 40 | Ionic liquids as refolding additives: variation of the anion. <i>Journal of Biotechnology</i> , 2010 , 150, 64-72 | 3.7 | 82 |
| 39 | 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 |
| 38 | 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 |
| 37 | 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 |
| 36 | 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 |
| 35 | 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 |
| 34 | The linker between the D3 and A1 domains of vWF suppresses A1-GPIb α catch bonds by site-specific binding to the A1 domain. <i>Protein Science</i> , 2013 , 22, 1049-59 | 6.3 | 26 |
| 33 | 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 |
| 32 | 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 |
| 31 | 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 |
| 30 | 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 |
| 29 | 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 |
| 28 | 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 |
| 27 | 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 |
| 26 | Mayo CALR mutation type classification guide using alpha helix propensity. <i>American Journal of Hematology</i> , 2018 , 93, E128-E129 | 7.1 | 11 |
| 25 | 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 |

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| 24 | 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 |
| 23 | 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 |
| 22 | 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 |
| 21 | 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 |
| 20 | 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 |
| 19 | The Chaperonin GroEL: A Versatile Tool for Applied Biotechnology Platforms. <i>Frontiers in Molecular Biosciences</i> , 2018 , 5, 46 | 5.6 | 6 |
| 18 | Sequence Reversal Prevents Chain Collapse and Yields Heat-Sensitive Intrinsic Disorder. <i>Biophysical Journal</i> , 2018 , 115, 328-340 | 2.9 | 5 |
| 17 | Chaperonin-Based Biolayer Interferometry To Assess the Kinetic Stability of Metastable, Aggregation-Prone Proteins. <i>Biochemistry</i> , 2016 , 55, 4885-908 | 3.2 | 5 |
| 16 | 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 |
| 15 | 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 |
| 14 | Measuring Lipid Transfer Protein Activity Using Bicelle-Dilution Model Membranes. <i>Analytical Chemistry</i> , 2020 , 92, 3417-3425 | 7.8 | 3 |
| 13 | 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 |
| 12 | "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 |
| 11 | Functional succinate dehydrogenase deficiency and loss of ascorbic acid transporter SLC23A1 are pathognomonic adverse features of clear cell renal cancer | | 2 |
| 10 | 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 |
| 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 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 |
| 7 | 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 |

- 6 Platelet-type von Willebrand disease: Local disorder of the platelet GPIb α switch drives high-affinity binding to von Willebrand factor. *Journal of Thrombosis and Haemostasis*, **2019**, 17, 2022-2034 ^{15.4} ○
- 5 Quantification of von Willebrand factor and ADAMTS-13 after traumatic injury: a pilot study. *Trauma Surgery and Acute Care Open*, **2021**, 6, e000703 2.4 ○
- 4 Functional Interrogation of Variants of Undetermined Significance of the Isocitrate Dehydrogenase 1 and 2 Genes in Myeloid Neoplasms. *Blood*, **2019**, 134, 1697-1697 2.2
- 3 Misfolding Induced By the Mutations V1314D and F1369I Affects the Function and the Structure of the Von Willebrand A1-Domain. *Blood*, **2014**, 124, 1521-1521 2.2
- 2 The Prognostic Advantage of Calreticulin Mutations in Myelofibrosis Might be Confined to Type 1 or Type 1-like Calreticulin Variants. *Blood*, **2014**, 124, 3166-3166 2.2
- 1 Functional evaluation of isocitrate dehydrogenase 1 and 2 variants of unclear significance in chronic myeloid neoplasms. *Leukemia Research*, **2019**, 87, 106264 2.7