

Sonja A Dames

List of Publications by Year in descending order

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29
papers

905
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687363

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times ranked

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citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | NMR- and MD simulation-based structural characterization of the membrane-associating FATC domain of ataxia telangiectasia mutated. <i>Journal of Biological Chemistry</i> , 2019, 294, 7098-7112. | 3.4 | 7 |
| 2 | ¹ H, ¹⁵ N, and ¹³ C chemical shift assignments of the micelle immersed FAT C-terminal (FATC) domains of the human protein kinases ataxia-telangiectasia mutated (ATM) and DNA-dependent protein kinase catalytic subunit (DNA-PKcs) fused to the B1 domain of streptococcal protein G (GB1). <i>Biomolecular NMR Assignments</i> , 2018, 12, 149-154. | 0.8 | 1 |
| 3 | NMR analysis of the backbone dynamics of the small GTPase Rheb and its interaction with the regulatory protein FKBP38. <i>FEBS Letters</i> , 2018, 592, 130-146. | 2.8 | 6 |
| 4 | Target of rapamycin FATC domain as a general membrane anchor: The FKBP12 like domain of FKBP38 as a case study. <i>Protein Science</i> , 2018, 27, 546-560. | 7.6 | 4 |
| 5 | Chemical shift assignment of the intrinsically disordered N-terminus and the rubredoxin domain in the folded metal bound and unfolded oxidized state of mycobacterial protein kinase G. <i>Biomolecular NMR Assignments</i> , 2016, 10, 401-406. | 0.8 | 2 |
| 6 | Oxidative Unfolding of the Rubredoxin Domain and the Natively Disordered N-terminal Region Regulate the Catalytic Activity of Mycobacterium tuberculosis Protein Kinase G. <i>Journal of Biological Chemistry</i> , 2016, 291, 27062-27072. | 3.4 | 12 |
| 7 | A Rigorous and Efficient Method To Reweight Very Large Conformational Ensembles Using Average Experimental Data and To Determine Their Relative Information Content. <i>Journal of Chemical Theory and Computation</i> , 2016, 12, 383-394. | 5.3 | 43 |
| 8 | One short cysteine-rich sequence pattern - two different disulfide-bonded structures - a molecular dynamics simulation study. <i>Journal of Peptide Science</i> , 2015, 21, 480-494. | 1.4 | 1 |
| 9 | Regulation of the Target of Rapamycin and Other Phosphatidylinositol 3-Kinase-Related Kinases by Membrane Targeting. <i>Membranes</i> , 2015, 5, 553-575. | 3.0 | 13 |
| 10 | Expression and purification of the natively disordered and redox sensitive metal binding regions of Mycobacterium tuberculosis protein kinase G. <i>Protein Expression and Purification</i> , 2015, 111, 68-74. | 1.3 | 5 |
| 11 | Subtype-Specific Modulation of Estrogen Receptor Coactivator Interaction by Phosphorylation. <i>ACS Chemical Biology</i> , 2015, 10, 475-484. | 3.4 | 17 |
| 12 | Characterization of residue-dependent differences in the peripheral membrane association of the FATC domain of the kinase target of rapamycin TM by NMR and CD spectroscopy. <i>FEBS Letters</i> , 2014, 588, 1755-1766. | 2.8 | 6 |
| 13 | Characterization of the Immersion Properties of the Peripheral Membrane Anchor of the FATC Domain of the Kinase Target of Rapamycin by NMR, Oriented CD Spectroscopy, and MD Simulations. <i>Journal of Physical Chemistry B</i> , 2014, 118, 4817-4831. | 2.6 | 14 |
| 14 | NMR- and Circular Dichroism-monitored Lipid Binding Studies Suggest a General Role for the FATC Domain as Membrane Anchor of Phosphatidylinositol 3-Kinase-related Kinases (PIKK). <i>Journal of Biological Chemistry</i> , 2013, 288, 20046-20063. | 3.4 | 17 |
| 15 | The FKBP-Rapamycin Binding Domain of Human TOR Undergoes Strong Conformational Changes in the Presence of Membrane Mimetics with and without the Regulator Phosphatidic Acid. <i>Biochemistry</i> , 2012, 51, 4909-4921. | 2.5 | 20 |
| 16 | A fast and simple method for probing the interaction of peptides and proteins with lipids and membrane mimetics using GB1 fusion proteins and NMR spectroscopy. <i>Protein Science</i> , 2012, 21, 1566-1570. | 7.6 | 12 |
| 17 | ¹ H, ¹⁵ N, and ¹³ C assignments of the N-terminal activation domain of Dictyostelium discoideum Formin C. <i>Biomolecular NMR Assignments</i> , 2011, 5, 47-49. | 0.8 | 3 |
| 18 | Structure, Dynamics, Lipid Binding, and Physiological Relevance of the Putative GTPase-binding Domain of Dictyostelium Formin C*. <i>Journal of Biological Chemistry</i> , 2011, 286, 36907-36920. | 3.4 | 19 |

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|----|--|-----|-----------|
| 19 | Structural Basis for the Association of the Redox-sensitive Target of Rapamycin FATC Domain with Membrane-mimetic Micelles. <i>Journal of Biological Chemistry</i> , 2010, 285, 7766-7775. | 3.4 | 20 |
| 20 | A fast and simple method to prepare the FKBP-rapamycin binding domain of human target of rapamycin for NMR binding assays. <i>Protein Expression and Purification</i> , 2008, 59, 31-37. | 1.3 | 3 |
| 21 | Insights into the Low Adhesive Capacity of Human T-cadherin from the NMR Structure of Its N-terminal Extracellular Domain. <i>Journal of Biological Chemistry</i> , 2008, 283, 23485-23495. | 3.4 | 28 |
| 22 | Structure of the Cyclin T binding domain of Hexim1 and molecular basis for its recognition of P-TEFb. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 14312-14317. | 7.1 | 55 |
| 23 | NMR assignment of the Cyclin T-binding domain of human Hexim1. <i>Journal of Biomolecular NMR</i> , 2006, 36, 39-39. | 2.8 | 2 |
| 24 | The Solution Structure of the FATC Domain of the Protein Kinase Target of Rapamycin Suggests a Role for Redox-dependent Structural and Cellular Stability. <i>Journal of Biological Chemistry</i> , 2005, 280, 20558-20564. | 3.4 | 111 |
| 25 | Structural basis for Hif-1 α /CBP recognition in the cellular hypoxic response. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002, 99, 5271-5276. | 7.1 | 376 |
| 26 | Contributions of the ionization states of acidic residues to the stability of the coiled coil domain of matrilin-1. <i>FEBS Letters</i> , 1999, 446, 75-80. | 2.8 | 18 |
| 27 | NMR structure of a parallel homotrimeric coiled coil. <i>Nature Structural and Molecular Biology</i> , 1998, 5, 687-691. | 8.2 | 36 |
| 28 | Heteronuclear NMR assignments and secondary structure of the coiled coil trimerization domain from cartilage matrix protein in oxidized and reduced forms. <i>Protein Science</i> , 1997, 6, 1734-1745. | 7.6 | 40 |
| 29 | A fragment of staphylococcal nuclease with an OB β fold structure shows hydrogen α exchange protection factors in the range reported for α molten globules. <i>Protein Science</i> , 1996, 5, 1942-1946. | 7.6 | 12 |