Michael D Urbaniak

List of Publications by Year in descending order

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331670 361022 51 1,406 21 35 citations h-index g-index papers 59 59 59 1836 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Chromosome-Scale Assembly of the Complete Genome Sequence of <i>Leishmania </i> (<i>Mundinia </i>) Tj ETQ e0005821.	0.6	34314 rgBT /C 5
2	LGAAP: Leishmaniinae Genome Assembly and Annotation Pipeline. Microbiology Resource Announcements, 2021, 10, e0043921.	0.6	8
3	Chromosome-scale genome sequencing, assembly and annotation of six genomes from subfamily Leishmaniinae. Scientific Data, 2021, 8, 234.	5.3	5
4	Chromosome-Scale Assembly of the Complete Genome Sequence of Leishmania (Mundinia) orientalis, Isolate LSCM4, Strain LV768. Microbiology Resource Announcements, 2021, 10, e0057421.	0.6	5
5	Extensive Translational Regulation through the Proliferative Transition of Trypanosoma cruzi Revealed by Multi-Omics. MSphere, 2021, 6, e0036621.	2.9	10
6	Chromosome-Scale Assembly of the Complete Genome Sequence of Leishmania (Mundinia) enriettii, Isolate CUR178, Strain LV763. Microbiology Resource Announcements, 2021, 10, e0057521.	0.6	2
7	Chromosome-Scale Assembly of the Complete Genome Sequence of Leishmania (Mundinia) sp. Ghana, Isolate GH5, Strain LV757. Microbiology Resource Announcements, 2021, 10, e0059121.	0.6	О
8	Chromosome-Scale Assembly of the Complete Genome Sequence of Porcisia hertigi, Isolate C119, Strain LV43. Microbiology Resource Announcements, 2021, 10, e0065121.	0.6	1
9	A mechanism-inspired UDP- <i>N</i> -acetylglucosamine pyrophosphorylase inhibitor. RSC Chemical Biology, 2020, 1, 13-25.	4.1	20
10	Phosphoproteomic analysis of mammalian infective Trypanosoma brucei subjected to heat shock suggests atypical mechanisms for thermotolerance. Journal of Proteomics, 2020, 219, 103735.	2.4	11
11	Proteome-Wide Quantitative Phosphoproteomic Analysis of Trypanosoma brucei Insect and Mammalian Life Cycle Stages. Methods in Molecular Biology, 2020, 2116, 125-137.	0.9	0
12	Re-evaluation of Diadenosine Tetraphosphate (Ap4A) From a Stress Metabolite to Bona Fide Secondary Messenger. Frontiers in Molecular Biosciences, 2020, 7, 606807.	3 . 5	23
13	Organising the cell cycle in the absence of transcriptional control: Dynamic phosphorylation co-ordinates the Trypanosoma brucei cell cycle post-transcriptionally. PLoS Pathogens, 2019, 15, e1008129.	4.7	33
14	Title is missing!. , 2019, 15, e1008129.		0
15	Title is missing!. , 2019, 15, e1008129.		0
16	Title is missing!. , 2019, 15, e1008129.		0
17	Title is missing!. , 2019, 15, e1008129.		0
18	Dynamic regulation of the Trypanosoma brucei transferrin receptor in response to iron starvation is mediated via the 3'UTR. PLoS ONE, 2018, 13, e0206332.	2.5	9

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19	Development of Chemical Proteomics for the Folateome and Analysis of the Kinetoplastid Folateome. ACS Infectious Diseases, 2018, 4, 1475-1486.	3.8	1
20	Cyclin-dependent kinase 12 is a drug target for visceral leishmaniasis. Nature, 2018, 560, 192-197.	27.8	112
21	Fluorescent mannosides serve as acceptor substrates for glycosyltransferase and sugar-1-phosphate transferase activities in Euglena gracilis membranes. Carbohydrate Research, 2017, 438, 26-38.	2.3	15
22	Cell cycle synchronisation of Trypanosoma brucei by centrifugal counter-flow elutriation reveals the timing of nuclear and kinetoplast DNA replication. Scientific Reports, 2017, 7, 17599.	3.3	25
23	The Dictyostelium prestalk inducer differentiation-inducing factor-1 (DIF-1) triggers unexpectedly complex global phosphorylation changes. Molecular Biology of the Cell, 2015, 26, 805-820.	2.1	15
24	Molecular control of irreversible bistability during trypanosome developmental commitment. Journal of Cell Biology, 2015, 211, 455-468.	5.2	46
25	TrypanoCyc: a community-led biochemical pathways database for Trypanosoma brucei. Nucleic Acids Research, 2015, 43, D637-D644.	14.5	35
26	Genomic and Proteomic Studies on the Mode of Action of Oxaboroles against the African Trypanosome. PLoS Neglected Tropical Diseases, 2015, 9, e0004299.	3.0	34
27	Probing the substrate specificity of <i>Trypanosoma brucei</i> GlcNAc-Pl de- <i>N</i> -acetylase with synthetic substrate analogues. Organic and Biomolecular Chemistry, 2014, 12, 1919-1934.	2.8	6
28	High-Confidence Glycosome Proteome for Procyclic Form <i>Trypanosoma brucei</i> by Epitope-Tag Organelle Enrichment and SILAC Proteomics. Journal of Proteome Research, 2014, 13, 2796-2806.	3.7	92
29	Fragment screening reveals salicylic hydroxamic acid as an inhibitor of Trypanosoma brucei GPI GlcNAc-PI de-N-acetylase. Carbohydrate Research, 2014, 387, 54-58.	2.3	11
30	Global Quantitative SILAC Phosphoproteomics Reveals Differential Phosphorylation Is Widespread between the Procyclic and Bloodstream Form Lifecycle Stages of <i>Trypanosoma brucei</i> . Journal of Proteome Research, 2013, 12, 2233-2244.	3.7	172
31	Genetic and structural validation of <i><scp>A</scp>spergillus fumigatus</i> â€ <scp>UDP</scp> â€xi> <scp>N</scp> âetacetylglucosamine pyrophosphorylase as an antifungal target. Molecular Microbiology, 2013, 89, 479-493.	2.5	29
32	A Novel Allosteric Inhibitor of the Uridine Diphosphate <i>N</i> -Acetylglucosamine Pyrophosphorylase from <i>Trypanosoma brucei</i> . ACS Chemical Biology, 2013, 8, 1981-1987.	3.4	23
33	Chemical Proteomic Analysis Reveals the Drugability of the Kinome of <i>Trypanosoma brucei</i> . ACS Chemical Biology, 2012, 7, 1858-1865.	3.4	53
34	Comparative SILAC Proteomic Analysis of Trypanosoma brucei Bloodstream and Procyclic Lifecycle Stages. PLoS ONE, 2012, 7, e36619.	2.5	147
35	Inhibitors Incorporating Zincâ€Binding Groups Target the GlcNAcâ€PI deâ€ <i>N</i> â€acetylase in <i>Trypanosoma brucei</i> , the Causative Agent of African Sleeping Sickness. Chemical Biology and Drug Design, 2012, 79, 270-278.	3.2	7
36	Synthesis of potential metal-binding group compounds to examine the zinc dependency of the GPI de-N-acetylase metalloenzyme in Trypanosoma brucei. Carbohydrate Research, 2011, 346, 708-714.	2.3	7

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37	A Multidimensional Strategy to Detect Polypharmacological Targets in the Absence of Structural and Sequence Homology. PLoS Computational Biology, 2010, 6, e1000648.	3.2	72
38	Computer-Aided Identification of <i>Trypanosoma brucei</i> Uridine Diphosphate Galactose 4′-Epimerase Inhibitors: Toward the Development of Novel Therapies for African Sleeping Sickness. Journal of Medicinal Chemistry, 2010, 53, 5025-5032.	6.4	56
39	Chapter 3 The GlcNAcâ€PI deâ€Nâ€acetylase. The Enzymes, 2009, , 49-64.	1.7	4
40	Casein kinase 1 isoform 2 is essential for bloodstream form Trypanosoma brucei. Molecular and Biochemical Parasitology, 2009, 166, 183-185.	1.1	30
41	Synthesis of 1-d-6-O-[2-(N-hydroxyaminocarbonyl)amino-2-deoxy-î±-d-glucopyranosyl]-myo-inositol 1-(n-octadecyl phosphate): a potential metalloenzyme inhibitor of glycosylphosphatidylinositol biosynthesis. Carbohydrate Research, 2008, 343, 1478-1481.	2.3	6
42	Probing <i>Trypanosoma brucei</i> Clycosylphosphatidylinositol Biosynthesis Using Novel Precursorâ€Analogues. Chemical Biology and Drug Design, 2008, 72, 127-132.	3.2	8
43	Probing Enzymes Late in the Trypanosomal Glycosylphosphatidylinositol Biosynthetic Pathway with Synthetic Glycosylphosphatidylinositol Analogues. ACS Chemical Biology, 2008, 3, 625-634.	3.4	26
44	Proteinâ€"Small Molecule Interactions in Neocarzinostatin, the Prototypical Enediyne Chromoprotein Antibiotic. ChemBioChem, 2007, 8, 704-717.	2.6	34
45	Trypanosoma bruceiUDP-galactose-4′-epimerase in ternary complex with NAD+and the substrate analogue UDP-4-deoxy-4-fluoro-α-D-galactose. Acta Crystallographica Section F: Structural Biology Communications, 2006, 62, 829-834.	0.7	16
46	Identification of novel inhibitors of UDP-Glc 4′-epimerase, a validated drug target for african sleeping sickness. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 5744-5747.	2.2	37
47	Galactose Starvation in a Bloodstream Form Trypanosoma brucei UDP-Glucose 4′-Epimerase Conditional Null Mutant. Eukaryotic Cell, 2006, 5, 1906-1913.	3.4	41
48	The N-Acetyl-D-glucosaminylphosphatidylinositol De-N-acetylase of Glycosylphosphatidylinositol Biosynthesis Is a Zinc Metalloenzyme. Journal of Biological Chemistry, 2005, 280, 22831-22838.	3.4	38
49	Design and Synthesis of a Nitrogen Mustard Derivative Stabilized by Apo-neocarzinostatin. Journal of Medicinal Chemistry, 2004, 47, 4710-4715.	6.4	33
50	Chemical synthesis and cytotoxicity of dihydroxylated cyclopentenone analogues of neocarzinostatin chromophore. Bioorganic and Medicinal Chemistry Letters, 2003, 13, 2025-2027.	2.2	5
51	Solution Structure of a Novel Chromoprotein Derived from Apo-Neocarzinostatin and a Synthetic Chromophoreâ€. Biochemistry, 2002, 41, 11731-11739.	2.5	37