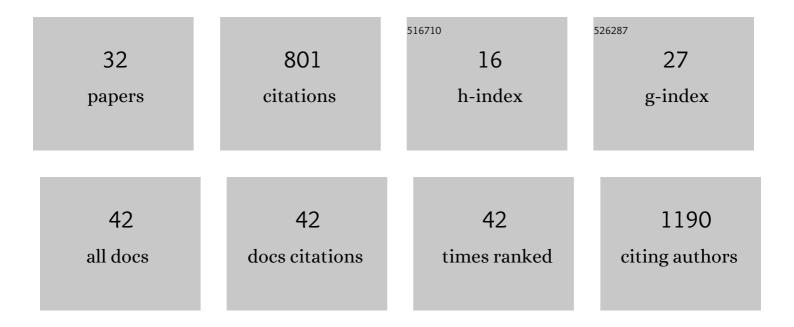
VÃ-tor G Mendes

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Arginine-deprivation–induced oxidative damage sterilizes <i>Mycobacterium tuberculosis</i> . Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 9779-9784.	7.1	97
2	Bacillus foraminis sp. nov., isolated from a non-saline alkaline groundwater. International Journal of Systematic and Evolutionary Microbiology, 2006, 56, 2571-2574.	1.7	55
3	Microcella putealis gen. nov., sp. nov., a Gram-positive alkaliphilic bacterium isolated from a nonsaline alkaline groundwater. Systematic and Applied Microbiology, 2005, 28, 479-487.	2.8	52
4	The molecular biology of mycobacterial trehalose in the quest for advanced tuberculosis therapies. Microbiology (United Kingdom), 2014, 160, 1547-1570.	1.8	50
5	Chimaereicella alkaliphila gen. nov., sp. nov., a Gram-negative alkaliphilic bacterium isolated from a nonsaline alkaline groundwater. Systematic and Applied Microbiology, 2006, 29, 100-108.	2.8	40
6	Structural Biology and the Design of New Therapeutics: From HIV and Cancer to Mycobacterial Infections. Journal of Molecular Biology, 2017, 429, 2677-2693.	4.2	39
7	Organic solutes in Rubrobacter xylanophilus: the first example of di-myo-inositol-phosphate in a thermophile. Extremophiles, 2007, 11, 667-673.	2.3	38
8	Targeting tuberculosis using structure-guided fragment-based drug design. Drug Discovery Today, 2017, 22, 546-554.	6.4	36
9	Identification of the mycobacterial glucosyl-3-phosphoglycerate synthase. FEMS Microbiology Letters, 2008, 280, 195-202.	1.8	33
10	Phenylobacterium falsum sp. nov., an Alphaproteobacterium isolated from a nonsaline alkaline groundwater, and emended description of the genus Phenylobacterium. Systematic and Applied Microbiology, 2005, 28, 295-302.	2.8	32
11	Development of Inhibitors against <i>Mycobacterium abscessus</i> tRNA (m ¹ G37) Methyltransferase (TrmD) Using Fragment-Based Approaches. Journal of Medicinal Chemistry, 2019, 62, 7210-7232.	6.4	32
12	Structure-guided fragment-based drug discovery at the synchrotron: screening binding sites and correlations with hotspot mapping. Philosophical Transactions Series A, Mathematical, Physical, and Engineering Sciences, 2019, 377, 20180422.	3.4	30
13	Biochemical characterization of the maltokinase from Mycobacterium bovis BCG. BMC Biochemistry, 2010, 11, 21.	4.4	29
14	Fragment-Based Design of <i>Mycobacterium tuberculosis</i> InhA Inhibitors. Journal of Medicinal Chemistry, 2020, 63, 4749-4761.	6.4	27
15	Biosynthesis of mycobacterial methylglucose lipopolysaccharides. Natural Product Reports, 2012, 29, 834.	10.3	25
16	Target Identification of Mycobacterium tuberculosis Phenotypic Hits Using a Concerted Chemogenomic, Biophysical, and Structural Approach. Frontiers in Pharmacology, 2017, 8, 681.	3.5	22
17	Fragment-based discovery of a new class of inhibitors targeting mycobacterial tRNA modification. Nucleic Acids Research, 2020, 48, 8099-8112.	14.5	20
18	Two Alternative Pathways for the Synthesis of the Rare Compatible Solute Mannosylglucosylglycerate in <i>Petrotoga mobilis</i> . Journal of Bacteriology, 2010, 192, 1624-1633.	2.2	17

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19	Genome Sequence of Mycobacterium hassiacum DSM 44199, a Rare Source of Heat-Stable Mycobacterial Proteins. Journal of Bacteriology, 2012, 194, 7010-7011.	2.2	17
20	Structural insights into the EthR–DNA interaction using native mass spectrometry. Chemical Communications, 2017, 53, 3527-3530.	4.1	17
21	Mycobacterium tuberculosis Rv2419c, the missing glucosyl-3-phosphoglycerate phosphatase for the second step in methylglucose lipopolysaccharide biosynthesis. Scientific Reports, 2011, 1, 177.	3.3	16
22	A fragment-based approach to assess the ligandability of ArgB, ArgC, ArgD and ArgF in the L-arginine biosynthetic pathway of Mycobacterium tuberculosis. Computational and Structural Biotechnology Journal, 2021, 19, 3491-3506.	4.1	16
23	Fragment Screening against the EthR–DNA Interaction by Native Mass Spectrometry. Angewandte Chemie - International Edition, 2017, 56, 7488-7491.	13.8	12
24	Development of Inhibitors of SAICAR Synthetase (PurC) from <i>Mycobacterium abscessus</i> Using a Fragment-Based Approach. ACS Infectious Diseases, 2022, 8, 296-309.	3.8	10
25	Inhibiting Mycobacterium tuberculosis CoaBC by targeting an allosteric site. Nature Communications, 2021, 12, 143.	12.8	8
26	Mycobacterial OtsA Structures Unveil Substrate Preference Mechanism and Allosteric Regulation by 2-Oxoglutarate and 2-Phosphoglycerate. MBio, 2019, 10, .	4.1	7
27	Crystal structure of <i>Staphylococcus aureus</i> Zn-glyoxalase I: new subfamily of glyoxalase I family. Journal of Biomolecular Structure and Dynamics, 2018, 36, 376-386.	3.5	5
28	Discovery of Novel Inhibitors of Uridine Diphosphate- <i>N</i> -Acetylenolpyruvylglucosamine Reductase (MurB) from <i>Pseudomonas aeruginosa</i> , an Opportunistic Infectious Agent Causing Death in Cystic Fibrosis Patients. Journal of Medicinal Chemistry, 2022, 65, 2149-2173.	6.4	5
29	Structural insights into <i>Escherichia coli</i> phosphopantothenoylcysteine synthetase by native ion mobility–mass spectrometry. Biochemical Journal, 2019, 476, 3125-3139.	3.7	4
30	Structure of Mycobacterium thermoresistibile ClgE defines novel conformational states that contribute to the catalytic mechanism. Scientific Reports, 2015, 5, 17144.	3.3	3
31	Targeting <i>Mycobacterium tuberculosis</i> CoaBC through Chemical Inhibition of 4â€2-Phosphopantothenoyl- <scp>l</scp> -cysteine Synthetase (CoaB) Activity. ACS Infectious Diseases, 2021, 7, 1666-1679.	3.8	3
32	Fragment Screening against the EthR–DNA Interaction by Native Mass Spectrometry. Angewandte Chemie, 2017, 129, 7596-7599.	2.0	2