Gregory M Cook

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1604509/publications.pdf

Version: 2024-02-01

66343 91884 6,224 170 42 69 citations h-index g-index papers 185 185 185 5992 docs citations citing authors all docs times ranked

#	Article	IF	CITATIONS
1	Multiplexed transcriptional repression identifies a network of bactericidal interactions between mycobacterial respiratory complexes. IScience, 2022, 25, 103573.	4.1	10
2	An amiloride derivative is active against the F1Fo-ATP synthase and cytochrome bd oxidase of Mycobacterium tuberculosis. Communications Biology, 2022, 5, 166.	4.4	21
3	Deciphering functional redundancy and energetics of malate oxidation in mycobacteria. Journal of Biological Chemistry, 2022, 298, 101859.	3.4	10
4	Potent Bactericidal Antimycobacterials Targeting the Chaperone ClpC1 Based on the Depsipeptide Natural Products Ecumicin and Ohmyungsamycin A. Journal of Medicinal Chemistry, 2022, 65, 4893-4908.	6.4	9
5	Stereochemical Effects on the Antimicrobial Properties of Tetrasubstituted 2,5-Diketopiperazines. ACS Medicinal Chemistry Letters, 2022, 13, 632-640.	2.8	5
6	Rate-limiting transport of positively charged arginine residues through the Sec-machinery is integral to the mechanism of protein secretion. ELife, 2022, 11 , .	6.0	13
7	Antimicrobial tolerance and its role in the development of resistance: Lessons from enterococci. Advances in Microbial Physiology, 2022, , .	2.4	3
8	C-2 derivatized 8-sulfonamidoquinolines as antibacterial compounds. Bioorganic and Medicinal Chemistry, 2021, 29, 115837.	3.0	2
9	Synthesis and Biological Evaluation of (â^') and (+)â€5piroleucettadine and Analogues. ChemMedChem, 2021, 16, 1308-1315.	3.2	1
10	<i>Mycobacterium smegmatis</i> Resists the Bactericidal Activity of Hypochlorous Acid Produced in Neutrophil Phagosomes. Journal of Immunology, 2021, 206, 1901-1912.	0.8	8
11	Growth on Formic Acid Is Dependent on Intracellular pH Homeostasis for the Thermoacidophilic Methanotroph Methylacidiphilum sp. RTK17.1. Frontiers in Microbiology, 2021, 12, 651744.	3.5	12
12	Systematic evaluation of structure–property relationships and pharmacokinetics in 6-(hetero)aryl-substituted matched pair analogs of amiloride and 5-(N,N-hexamethylene)amiloride. Bioorganic and Medicinal Chemistry, 2021, 37, 116116.	3.0	9
13	Using genome comparisons of wild-type and resistant mutants of Methanococcus maripaludis to help understand mechanisms of resistance to methane inhibitors. Access Microbiology, 2021, 3, 000244.	0.5	1
14	Disruption of Metallostasis in the Anaerobic Human Pathogen <i>Fusobacterium nucleatum</i> by the Zinc Ionophore PBT2. ACS Infectious Diseases, 2021, 7, 2285-2298.	3.8	6
15	CRISPR interference identifies vulnerable cellular pathways with bactericidal phenotypes in <i>Mycobacterium tuberculosis</i> . Molecular Microbiology, 2021, 116, 1033-1043.	2.5	17
16	A Concise Synthetic Strategy Towards the Novel Calcium-dependent Lipopeptide Antibiotic, Malacidin A and Analogues. Frontiers in Chemistry, 2021, 9, 687875.	3.6	6
17	Nitric Oxide-Dependent Electron Transport Chain Inhibition by the Cytochrome <i>bc</i> ₁ Inhibitor and Pretomanid Combination Kills <i>Mycobacterium tuberculosis</i> and Chemotherapy, 2021, 65, e0095621.	3.2	9
18	The cryo-EM structure of the bd oxidase from M. tuberculosis reveals a unique structural framework and enables rational drug design to combat TB. Nature Communications, 2021, 12, 5236.	12.8	29

#	Article	IF	CITATIONS
19	Discovery of 5-methylpyrimidopyridone analogues as selective antimycobacterial agents. Bioorganic and Medicinal Chemistry, 2021, 49, 116426.	3.0	1
20	Sterilizing Effects of Novel Regimens Containing TB47, Clofazimine, and Linezolid in a Murine Model of Tuberculosis. Antimicrobial Agents and Chemotherapy, 2021, 65, e0070621.	3.2	10
21	Dual inhibition of the terminal oxidases eradicates antibioticâ€tolerant <i>Mycobacterium tuberculosis</i> . EMBO Molecular Medicine, 2021, 13, e13207.	6.9	47
22	Genomic Profiling of <i>Mycobacterium tuberculosis</i> Strains, Myanmar. Emerging Infectious Diseases, 2021, 27, 2847-2855.	4.3	8
23	Functionalized Dioxonaphthoimidazoliums: A Redox Cycling Chemotype with Potent Bactericidal Activities against <i>Mycobacterium tuberculosis</i> Journal of Medicinal Chemistry, 2021, 64, 15991-16007.	6.4	10
24	Utilization of CRISPR interference to investigate the contribution of genes to pathogenesis in a macrophage model of Mycobacterium tuberculosis infection. Journal of Antimicrobial Chemotherapy, 2021, , .	3.0	9
25	Synthetic Sansanmycin Analogues as Potent <i>Mycobacterium tuberculosis</i> Translocase I Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 17326-17345.	6.4	8
26	Two for the price of one: Attacking the energetic-metabolic hub of mycobacteria to produce new chemotherapeutic agents. Progress in Biophysics and Molecular Biology, 2020, 152, 35-44.	2.9	22
27	Unprecedented Properties of Phenothiazines Unraveled by a NDH-2 Bioelectrochemical Assay Platform. Journal of the American Chemical Society, 2020, 142, 1311-1320.	13.7	18
28	Genomic analysis of Caldalkalibacillus thermarum TA2.A1 reveals aerobic alkaliphilic metabolism and evolutionary hallmarks linking alkaliphilic bacteria and plant life. Extremophiles, 2020, 24, 923-935.	2.3	10
29	Repurposing a neurodegenerative disease drug to treat Gram-negative antibiotic-resistant bacterial sepsis. Science Translational Medicine, 2020, 12, .	12.4	36
30	Microbial energy managementâ€"A product of three broad tradeoffs. Advances in Microbial Physiology, 2020, 77, 139-185.	2.4	12
31	MmpL3 inhibitors as antituberculosis drugs. European Journal of Medicinal Chemistry, 2020, 200, 112390.	5.5	31
32	Transcriptional Inhibition of the F $<$ sub $>$ 1 $<$ /sub $>$ F $<$ sub $>$ 0 $<$ /sub $>$ -Type ATP Synthase Has Bactericidal Consequences on the Viability of Mycobacteria. Antimicrobial Agents and Chemotherapy, 2020, 64, .	3.2	17
33	Synthesis of paenipeptin C′ analogues employing solution-phase CLipPA chemistry. Organic and Biomolecular Chemistry, 2020, 18, 4381-4385.	2.8	7
34	Total Synthesis and Antimycobacterial Activity of Ohmyungsamycinâ€A, Deoxyecumicin, and Ecumicin. Chemistry - A European Journal, 2020, 26, 15200-15205.	3.3	8
35	"CLipPâ€ing on lipids to generate antibacterial lipopeptides. Chemical Science, 2020, 11, 5759-5765.	7.4	15
36	Antitubercular polyhalogenated phenothiazines and phenoselenazine with reduced binding to CNS receptors. European Journal of Medicinal Chemistry, 2020, 201, 112420.	5.5	12

#	Article	IF	Citations
37	Multiple Bactericidal Mechanisms of the Zinc Ionophore PBT2. MSphere, 2020, 5, .	2.9	24
38	Discovery of a Natural Product That Binds to the Mycobacterium tuberculosis Protein Rv1466 Using Native Mass Spectrometry. Molecules, 2020, 25, 2384.	3.8	18
39	Cellular and Structural Basis of Synthesis of the Unique Intermediate Dehydro-F ₄₂₀ -0 in Mycobacteria. MSystems, 2020, 5, .	3.8	9
40	Predicting nitroimidazole antibiotic resistance mutations in Mycobacterium tuberculosis with protein engineering. PLoS Pathogens, 2020, 16, e1008287.	4.7	51
41	Antituberculosis Activity of the Antimalaria Cytochrome <i>bcc</i> Oxidase Inhibitor SCR0911. ACS Infectious Diseases, 2020, 6, 725-737.	3.8	10
42	Discovery of Cephalosporin-3′-Diazeniumdiolates That Show Dual Antibacterial and Antibiofilm Effects against <i>Pseudomonas aeruginosa</i> Clinical Cystic Fibrosis Isolates and Efficacy in a Murine Respiratory Infection Model. ACS Infectious Diseases, 2020, 6, 1460-1479.	3.8	18
43	Synthesis of Functionalised Chromonylâ€pyrimidines and Their Potential as Antimycobacterial Agents. ChemistrySelect, 2020, 5, 4347-4355.	1.5	5
44	Substituted sulfonamide bioisosteres of 8-hydroxyquinoline as zinc-dependent antibacterial compounds. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127110.	2.2	6
45	Predicting nitroimidazole antibiotic resistance mutations in Mycobacterium tuberculosis with protein engineering. FASEB Journal, 2020, 34, 1-1.	0.5	0
46	Title is missing!. , 2020, 16, e1008287.		0
47	Title is missing!. , 2020, 16, e1008287.		0
48	Title is missing!. , 2020, 16, e1008287.		0
49	Title is missing!. , 2020, 16, e1008287.		0
50	Diverse hydrogen production and consumption pathways influence methane production in ruminants. ISME Journal, 2019, 13, 2617-2632.	9.8	132
51	Two uptake hydrogenases differentially interact with the aerobic respiratory chain during mycobacterial growth and persistence. Journal of Biological Chemistry, 2019, 294, 18980-18991.	3.4	28
52	6-Substituted amiloride derivatives as inhibitors of the urokinase-type plasminogen activator for use in metastatic disease. Bioorganic and Medicinal Chemistry Letters, 2019, 29, 126753.	2.2	21
53	Disrupting coupling within mycobacterial F-ATP synthases subunit $\hat{l}\mu$ causes dysregulated energy production and cell wall biosynthesis. Scientific Reports, 2019, 9, 16759.	3.3	29
54	Derailing the aspartate pathway of Mycobacterium tuberculosis to eradicate persistent infection. Nature Communications, 2019, 10, 4215.	12.8	48

#	Article	IF	Citations
55	Microtiter Screening Reveals Oxygen-Dependent Antimicrobial Activity of Natural Products Against Mastitis-Causing Bacteria. Frontiers in Microbiology, 2019, 10, 1995.	3.5	2
56	Inhalable Dry Powder of Bedaquiline for Pulmonary Tuberculosis: In Vitro Physicochemical Characterization, Antimicrobial Activity and Safety Studies. Pharmaceutics, 2019, 11, 502.	4.5	24
57	Tackling tuberculosis in the indigenous people of New Zealand. Lancet Public Health, The, 2019, 4, e496.	10.0	5
58	Alternate quinone coupling in a new class of succinate dehydrogenase may potentiate mycobacterial respiratory control. FEBS Letters, 2019, 593, 475-486.	2.8	17
59	Synthesis and Investigation of Phthalazinones as Antitubercular Agents. Chemistry - an Asian Journal, 2019, 14, 1278-1285.	3.3	9
60	The structure of the catalytic domain of the ATP synthase from <i>Mycobacterium smegmatis</i> is a target for developing antitubercular drugs. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 4206-4211.	7.1	38
61	Structure of F 1 -ATPase from the obligate anaerobe Fusobacterium nucleatum. Open Biology, 2019, 9, 190066.	3.6	3
62	The synthesis and evaluation of quinolinequinones as anti-mycobacterial agents. Bioorganic and Medicinal Chemistry, 2019, 27, 3532-3545.	3.0	19
63	Utilization of CRISPR Interference To Validate MmpL3 as a Drug Target in <i>Mycobacterium tuberculosis</i> . Antimicrobial Agents and Chemotherapy, 2019, 63, .	3.2	37
64	Genomewide Profiling of the Enterococcus faecalis Transcriptional Response to Teixobactin Reveals CroRS as an Essential Regulator of Antimicrobial Tolerance. MSphere, 2019, 4, .	2.9	8
65	Agricultural Origins of a Highly Persistent Lineage of Vancomycin-Resistant <i>Enterococcus faecalis</i> in New Zealand. Applied and Environmental Microbiology, 2019, 85, .	3.1	11
66	Dispersal of Mycobacterium tuberculosis Driven by Historical European Trade in the South Pacific. Frontiers in Microbiology, 2019, 10, 2778.	3.5	28
67	Occurrence and expression of genes encoding methyl-compound production in rumen bacteria. Animal Microbiome, 2019, 1, 15.	3.8	27
68	FAD-sequestering proteins protect mycobacteria against hypoxic and oxidative stress. Journal of Biological Chemistry, 2019, 294, 2903-5814.	3.4	14
69	Pyrazolo[1,5- <i>a</i>]pyridine Inhibitor of the Respiratory Cytochrome <i>bcc</i> Complex for the Treatment of Drug-Resistant Tuberculosis. ACS Infectious Diseases, 2019, 5, 239-249.	3.8	74
70	Functional characterization of BcrR: a one-component transmembrane signal transduction system for bacitracin resistance. Microbiology (United Kingdom), 2019, 165, 475-487.	1.8	7
71	Complete Genome Sequence of a New Zealand Isolate of the Bovine Pathogen Streptococcus uberis. Genome Announcements, 2018, 6, .	0.8	3
72	Structure of the NDH-2 – HQNO inhibited complex provides molecular insight into quinone-binding site inhibitors. Biochimica Et Biophysica Acta - Bioenergetics, 2018, 1859, 482-490.	1.0	20

#	Article	IF	CITATIONS
73	Total Synthesis and Conformational Study of Callyaerinâ€A: Antiâ€Tubercular Cyclic Peptide Bearing a Rare Rigidifying (<i>Z</i>)â€2,3―Diaminoacrylamide Moiety. Angewandte Chemie - International Edition, 2018, 57, 3631-3635.	13.8	24
74	Total Synthesis and Conformational Study of Callyaerinâ€A: Antiâ€Tubercular Cyclic Peptide Bearing a Rare Rigidifying (<i>Z</i>)â€2,3―Diaminoacrylamide Moiety. Angewandte Chemie, 2018, 130, 3693-3697.	2.0	9
75	Innenrù⁄4cktitelbild: Total Synthesis and Conformational Study of Callyaerinâ€A: Antiâ€Tubercular Cyclic Peptide Bearing a Rare Rigidifying (<i>Z</i>)â€2,3―Diaminoacrylamide Moiety (Angew. Chem. 14/2018). Angewandte Chemie, 2018, 130, 3897-3897.	2.0	0
76	Association between anti-tuberculosis drug resistance-conferring mutations and treatment outcomes in Myanmar. Infectious Diseases, 2018, 50, 388-390.	2.8	1
77	Targeting bacterial energetics to produce new antimicrobials. Drug Resistance Updates, 2018, 36, 1-12.	14.4	72
78	Overexpression of a newly identified dâ€amino acid transaminase in <i>Mycobacterium smegmatis</i> complements glutamate racemase deletion. Molecular Microbiology, 2018, 107, 198-213.	2.5	33
79	Chemical Synergy between lonophore PBT2 and Zinc Reverses Antibiotic Resistance. MBio, 2018, 9, .	4.1	56
80	Microbiome dataset from the upper respiratory tract of patients living with HIV, HIV/TB and TB from Myanmar. Data in Brief, 2018, 21, 354-357.	1.0	1
81	Acquired Resistance to Antituberculosis Drugs. Emerging Infectious Diseases, 2018, 24, 2134-2134.	4.3	2
82	â€Tethering' fragment-based drug discovery to identify inhibitors of the essential respiratory membrane protein type II NADH dehydrogenase. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 2239-2243.	2.2	10
83	Ionophoric effects of the antitubercular drug bedaquiline. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, 7326-7331.	7.1	85
84	6-Substituted Hexamethylene Amiloride (HMA) Derivatives as Potent and Selective Inhibitors of the Human Urokinase Plasminogen Activator for Use in Cancer. Journal of Medicinal Chemistry, 2018, 61, 8299-8320.	6.4	56
85	Evaluation of the rapid molecular diagnostic test for the New Zealand Mycobacterium tuberculosis Rangipo strain in a clinical setting. New Zealand Medical Journal, 2018, 131, 70-72.	0.5	0
86	The mechanism of catalysis by type-II NADH:quinone oxidoreductases. Scientific Reports, 2017, 7, 40165.	3.3	45
87	Microarray dataset on the genome-wide expression profile of an M. smegmatis amtR mutant (JR258) compared to M. smegmatis mc 2 155. Data in Brief, 2017, 10, 38-40.	1.0	2
88	First 2 Extensively Drug-Resistant Tuberculosis Cases From Myanmar Treated With Bedaquiline. Clinical Infectious Diseases, 2017, 65, 531-532.	5.8	5
89	Oxidative Phosphorylation as a Target Space for Tuberculosis: Success, Caution, and Future Directions. Microbiology Spectrum, 2017, 5, .	3.0	89
90	First- and second-line antituberculosis drug resistance patterns among previous treatment failure patients in Myanmar. Journal of Global Antimicrobial Resistance, 2017, 9, 34-35.	2.2	1

#	Article	IF	CITATIONS
91	Rapid molecular diagnosis of the Mycobacterium tuberculosis Rangipo strain responsible for the largest recurring TB cluster in New Zealand. Diagnostic Microbiology and Infectious Disease, 2017, 88, 138-140.	1.8	8
92	Synthesis and biological evaluation of novel teixobactin analogues. Organic and Biomolecular Chemistry, 2017, 15, 8755-8760.	2.8	31
93	Role of Alanine Racemase Mutations in Mycobacterium tuberculosis <code><scp>d</scp></code> -Cycloserine Resistance. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	24
94	Evaluation of the genotype MTBDRsl test for detection of second-line drug resistance in drug-resistant Mycobacterium tuberculosis strains in Myanmar. Infectious Diseases, 2017, 49, 865-866.	2.8	1
95	Mixotrophy drives niche expansion of verrucomicrobial methanotrophs. ISME Journal, 2017, 11, 2599-2610.	9.8	107
96	Exploiting the synthetic lethality between terminal respiratory oxidases to kill <i>Mycobacterium tuberculosis</i> and clear host infection. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, 7426-7431.	7.1	141
97	Genotypic diversity of Mycobacterium tuberculosis strains in Myanmar. Infectious Diseases, 2017, 49, 237-239.	2.8	5
98	Crystal structure of type II NADH:quinone oxidoreductase from ⟨i⟩Caldalkalibacillus thermarum⟨/i⟩ with an improved resolution of 2.15â€Ã Acta Crystallographica Section F, Structural Biology Communications, 2017, 73, 541-549.	0.8	10
99	Draft Genome Sequences of Two Drug-Resistant Mycobacterium tuberculosis Isolates from Myanmar. Genome Announcements, 2016, 4, .	0.8	2
100	Activation of type II NADH dehydrogenase by quinolinequinones mediates antitubercular cell death. Journal of Antimicrobial Chemotherapy, 2016, 71, 2840-2847.	3.0	38
101	A bacterial oxidase like no other?. Science, 2016, 352, 518-519.	12.6	9
102	Drug-resistant tuberculosis among previously treated patients in Yangon, Myanmar. International Journal of Mycobacteriology, 2016, 5, 366-367.	0.6	2
103	Regulation of the thermoalkaliphilic F ₁ -ATPase from <i>Caldalkalibacillus thermarum</i> . Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 10860-10865.	7.1	51
104	CydDC-mediated reductant export in <i>Escherichia coli</i> controls the transcriptional wiring of energy metabolism and combats nitrosative stress. Biochemical Journal, 2016, 473, 693-701.	3.7	36
105	Structure and Function of AmtR in Mycobacterium smegmatis: Implications for Post-Transcriptional Regulation of Urea Metabolism through a Small Antisense RNA. Journal of Molecular Biology, 2016, 428, 4315-4329.	4.2	8
106	Whole-genome sequencing of multidrug-resistant Mycobacterium tuberculosis isolates from Myanmar. Journal of Global Antimicrobial Resistance, 2016, 6, 113-117.	2.2	28
107	Biophysical Characterization of a Thermoalkaliphilic Molecular Motor with a High Stepping Torque Gives Insight into Evolutionary ATP Synthase Adaptation. Journal of Biological Chemistry, 2016, 291, 23965-23977.	3.4	21
108	Annotated compound data for modulators of detergent-solubilised or lipid-reconstituted respiratory type II NADH dehydrogenase activity obtained by compound library screening. Data in Brief, 2016, 6, 275-278.	1.0	2

#	Article	IF	CITATIONS
109	Synthesis and activity of a diselenide bond mimetic of the antimicrobial protein caenopore-5. Chemical Science, 2016, 7, 2005-2010.	7.4	21
110	Comparison of lipid and detergent enzyme environments for identifying inhibitors of membrane-bound energy-transducing proteins. Journal of Microbiological Methods, 2016, 120, 41-43.	1.6	6
111	Genomic and metagenomic surveys of hydrogenase distribution indicate H2 is a widely utilised energy source for microbial growth and survival. ISME Journal, 2016, 10, 761-777.	9.8	503
112	Defining the nitrogen regulated transcriptome of Mycobacterium smegmatis using continuous culture. BMC Genomics, 2015, 16, 821.	2.8	29
113	Atmospheric Hydrogen Scavenging: from Enzymes to Ecosystems. Applied and Environmental Microbiology, 2015, 81, 1190-1199.	3.1	81
114	Novel regulatory roles of cAMP receptor proteins in fast-growing environmental mycobacteria. Microbiology (United Kingdom), 2015, 161, 648-661.	1.8	11
115	Bactericidal mode of action of bedaquiline. Journal of Antimicrobial Chemotherapy, 2015, 70, 2028-2037.	3.0	161
116	The cytochrome bd-type quinol oxidase is important for survival of Mycobacterium smegmatis under peroxide and antibiotic-induced stress. Scientific Reports, 2015, 5, 10333.	3.3	101
117	Role of the Transporter-Like Sensor Kinase CbrA in Histidine Uptake and Signal Transduction. Journal of Bacteriology, 2015, 197, 2867-2878.	2.2	22
118	A high-throughput screening assay for identification of inhibitors of the A1AO-ATP synthase of the rumen methanogen Methanobrevibacter ruminantium M1. Journal of Microbiological Methods, 2015, 110, 15-17.	1.6	3
119	Oral Bacitracin: A Consideration for Suppression of Intestinal Vancomycin-Resistant Enterococci (VRE) and for VRE Bacteremia From an Apparent Gastrointestinal Tract Source. Clinical Infectious Diseases, 2015, 60, 1726-1728.	5.8	7
120	Development of a Mycobacterium smegmatis transposon mutant array for characterising the mechanism of action of tuberculosis drugs: Findings with isoniazid and its structural analogues. Tuberculosis, 2015, 95, 432-439.	1.9	10
121	Persistence of the dominant soil phylum <i>Acidobacteria</i> by trace gas scavenging. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 10497-10502.	7.1	117
122	Bacillus subtilis as a Platform for Molecular Characterisation of Regulatory Mechanisms of Enterococcus faecalis Resistance against Cell Wall Antibiotics. PLoS ONE, 2014, 9, e93169.	2.5	9
123	Succinate Dehydrogenase is the Regulator of Respiration in Mycobacterium tuberculosis. PLoS Pathogens, 2014, 10, e1004510.	4.7	87
124	Essentiality of Succinate Dehydrogenase in Mycobacterium smegmatis and Its Role in the Generation of the Membrane Potential Under Hypoxia. MBio, 2014, 5, .	4.1	70
125	An obligately aerobic soil bacterium activates fermentative hydrogen production to survive reductive stress during hypoxia. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 11479-11484.	7.1	117
126	Three different [<scp><scp>NiFe</scp>< scp>] hydrogenases confer metabolic flexibility in the obligate aerobe <scp><i>M</i></scp><i>ycobacterium smegmatis</i>. Environmental Microbiology, 2014, 16, 318-330.</scp>	3.8	63

#	Article	IF	CITATIONS
127	Energetics of Respiration and Oxidative Phosphorylation in Mycobacteria. Microbiology Spectrum, 2014, 2, .	3.0	164
128	Structure of the bacterial type <scp>II NADH</scp> dehydrogenase: a monotopic membrane protein with an essential role in energy generation. Molecular Microbiology, 2014, 91, 950-964.	2.5	103
129	A soil actinobacterium scavenges atmospheric H ₂ using two membrane-associated, oxygen-dependent [NiFe] hydrogenases. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 4257-4261.	7.1	116
130	Integration of hydrogenase expression and hydrogen sensing in bacterial cell physiology. Current Opinion in Microbiology, 2014, 18, 30-38.	5.1	49
131	The succinate dehydrogenase assembly factor, SdhE, is required for the flavinylation and activation of fumarate reductase in bacteria. FEBS Letters, 2014, 588, 414-421.	2.8	32
132	Energetics of Pathogenic Bacteria and Opportunities for Drug Development. Advances in Microbial Physiology, 2014, 65, 1-62.	2.4	102
133	Crystal Structure of PhnF, a GntR-Family Transcriptional Regulator of Phosphate Transport in Mycobacterium smegmatis. Journal of Bacteriology, 2014, 196, 3472-3481.	2.2	17
134	Incorporation of triphenylphosphonium functionality improves the inhibitory properties of phenothiazine derivatives in Mycobacterium tuberculosis. Bioorganic and Medicinal Chemistry, 2014, 22, 5320-5328.	3.0	32
135	Investigation of the Essentiality of Glutamate Racemase in Mycobacterium smegmatis. Journal of Bacteriology, 2014, 196, 4239-4244.	2.2	15
136	Hypoxia-Activated Cytochrome <i>bd</i> Expression in Mycobacterium smegmatis Is Cyclic AMP Receptor Protein Dependent. Journal of Bacteriology, 2014, 196, 3091-3097.	2.2	35
137	Editorial overview: Cell regulation: Microbial cell regulation—looking in from the outside. Current Opinion in Microbiology, 2014, 18, v-vii.	5.1	1
138	Characterization of the proline-utilization pathway in <i>Mycobacterium tuberculosis </i> through structural and functional studies. Acta Crystallographica Section D: Biological Crystallography, 2014, 70, 968-980.	2.5	14
139	The Growth and Survival of Mycobacterium smegmatis Is Enhanced by Co-Metabolism of Atmospheric H2. PLoS ONE, 2014, 9, e103034.	2.5	55
140	Ribonucleases in bacterial toxin–antitoxin systems. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2013, 1829, 523-531.	1.9	77
141	A New Type of Na+-Driven ATP Synthase Membrane Rotor with a Two-Carboxylate Ion-Coupling Motif. PLoS Biology, 2013, 11, e1001596.	5.6	61
142	Bridging the Gap Between a TB Drug and Its Target. Science Translational Medicine, 2012, 4, 150fs33.	12.4	3
143	Toxin-Antitoxin Systems of Mycobacterium smegmatis Are Essential for Cell Survival. Journal of Biological Chemistry, 2012, 287, 5340-5356.	3.4	59
144	Regulation of proline metabolism in mycobacteria and its role in carbon metabolism under hypoxia. Molecular Microbiology, 2012, 84, 664-681.	2.5	71

#	Article	IF	CITATIONS
145	Unique Flexibility in Energy Metabolism Allows Mycobacteria to Combat Starvation and Hypoxia. PLoS ONE, 2010, 5, e8614.	2.5	179
146	The vapBC Operon from Mycobacterium smegmatis Is An Autoregulated Toxin–Antitoxin Module That Controls Growth via Inhibition of Translation. Journal of Molecular Biology, 2009, 390, 353-367.	4.2	96
147	Physiology of Mycobacteria. Advances in Microbial Physiology, 2009, 55, 81-319.	2.4	135
148	Unique Rotary ATP Synthase and Its Biological Diversity. Annual Review of Biophysics, 2008, 37, 43-64.	10.0	167
149	The alternative sigma factor SigF of Mycobacterium smegmatis is required for survival of heat shock, acidic pH and oxidative stress. Microbiology (United Kingdom), 2008, 154, 2786-2795.	1.8	42
150	A Specific Adaptation in the a Subunit of Thermoalkaliphilic F1FO-ATP Synthase Enables ATP Synthesis at High pH but Not at Neutral pH Values. Journal of Biological Chemistry, 2007, 282, 17395-17404.	3.4	48
151	A tridecameric c ring of the adenosine triphosphate (ATP) synthase from the thermoalkaliphilic <i>Bacillus</i> sp. strain TA2.A1 facilitates ATP synthesis at low electrochemical proton potential. Molecular Microbiology, 2007, 65, 1181-1192.	2.5	93
152	Structural Investigations of the Membrane-Embedded Rotor Ring of the F-ATPase from Clostridium paradoxum. Journal of Bacteriology, 2006, 188, 7759-7764.	2.2	59
153	Biochemical and Molecular Characterization of a Na \pm -Translocating F 1 F o -ATPase from the Thermoalkaliphilic Bacterium Clostridium paradoxum. Journal of Bacteriology, 2006, 188, 5045-5054.	2.2	60
154	The Phn system of Mycobacterium smegmatis: a second high-affinity ABC-transporter for phosphate. Microbiology (United Kingdom), 2006, 152, 3453-3465.	1.8	71
155	Inhibition of ATP Hydrolysis by Thermoalkaliphilic F 1 F o -ATP Synthase Is Controlled by the C Terminus of the Îμ Subunit. Journal of Bacteriology, 2006, 188, 3796-3804.	2.2	47
156	Mutants of Mycobacterium smegmatis unable to grow at acidic pH in the presence of the protonophore carbonyl cyanide m-chlorophenylhydrazone. Microbiology (United Kingdom), 2005, 151, 665-672.	1.8	35
157	The F 1 F o -ATP Synthase of Mycobacterium smegmatis Is Essential for Growth. Journal of Bacteriology, 2005, 187, 5023-5028.	2.2	100
158	Bacterial Na+- or H+-coupled ATP Synthases Operating at Low Electrochemical Potential. Advances in Microbial Physiology, 2004, 49, 175-218.	2.4	61
159	Amino acid transport by Sphingomonas sp. strain Ant 17 isolated from oil-contaminated Antarctic soil. Polar Biology, 2003, 26, 560-566.	1.2	5
160	Purification and Biochemical Characterization of the F 1 F o -ATP Synthase from Thermoalkaliphilic Bacillus sp. Strain TA2.A1. Journal of Bacteriology, 2003, 185, 4442-4449.	2.2	59
161	A novel haem compound accumulated in Escherichia coli overexpressing the cydDC operon, encoding an ABC-type transporter required for cytochrome assembly. Archives of Microbiology, 2002, 178, 358-369.	2.2	17
162	Intracellular pH regulation by Mycobacterium smegmatis and Mycobacterium bovis BCG. Microbiology (United Kingdom), 2001, 147, 1017-1024.	1.8	126

#	Article	IF	CITATIONS
163	Roles of respiratory oxidases in protecting Escherichia coli K12 from oxidative stress. Antonie Van Leeuwenhoek, 2000, 78, 23-31.	1.7	100
164	The intracellular pH of the thermophilic bacterium Thermoanaerobacter wiegelii during growth and production of fermentation acids. Extremophiles, 2000, 4, 279-284.	2.3	15
165	Sucrose transport by the alkaliphilic, thermophilic Bacillus sp. strain TA2.A1 is dependent on a sodium gradient. Extremophiles, 2000, 4, 291-296.	2.3	18
166	Oxidase and periplasmic cytochrome assembly in Escherichia coli K-12: CydDC and CcmAB are not required for haem–membrane association. Microbiology (United Kingdom), 2000, 146, 527-536.	1.8	36
167	Survival of Streptococcus pyogenes under stress and starvation. FEMS Microbiology Letters, 1999, 176, 421-428.	1.8	2
168	Oxidative Phosphorylation as a Target Space for Tuberculosis: Success, Caution, and Future Directions., 0,, 295-316.		4
169	Energetics of Respiration and Oxidative Phosphorylation in Mycobacteria., 0,, 389-409.		5
170	Bacterial respiration keeps amazing us in the 21st century. Biochemist, 0, , .	0.5	O