

# Ruben Hartkoorn

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1108021/publications.pdf>

Version: 2024-02-01

44  
papers

3,178  
citations

186265  
28  
h-index

243625  
44  
g-index

44  
all docs

44  
docs citations

44  
times ranked

3980  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pyridylpiperazine-based allosteric inhibitors of RND-type multidrug efflux pumps. <i>Nature Communications</i> , 2022, 13, 115.	12.8	28
2	Rubrolone production by <i>Dactylosporangium vinaceum</i> : biosynthesis, modulation and possible biological function. <i>Applied Microbiology and Biotechnology</i> , 2021, 105, 5541-5551.	3.6	5
3	Intragenic Distribution of IS <i>6110</i> in Clinical <i>Mycobacterium tuberculosis</i> Strains: Bioinformatic Evidence for Gene Disruption Leading to Underdiagnosed Antibiotic Resistance. <i>Microbiology Spectrum</i> , 2021, 9, e0001921.	3.0	9
4	Pyrrrolomycins Are Potent Natural Protonophores. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 63, .	3.2	29
5	Total Synthesis of Ripostatin B and Structure-Activity Relationship Studies on Ripostatin Analogs. <i>Journal of Organic Chemistry</i> , 2018, 83, 7150-7172.	3.2	22
6	MoS <sub>2</sub> /TiO <sub>2</sub> /SiNW surface as an effective substrate for LDI-MS detection of glucose and glutathione in real samples. <i>Talanta</i> , 2017, 171, 101-107.	5.5	24
7	Discovery and Biosynthesis of Gladiolin: A <i>Burkholderia gladioli</i> Antibiotic with Promising Activity against <i>Mycobacterium tuberculosis</i> . <i>Journal of the American Chemical Society</i> , 2017, 139, 7974-7981.	13.7	73
8	The Inosine Monophosphate Dehydrogenase, GuaB2, Is a Vulnerable New Bactericidal Drug Target for Tuberculosis. <i>ACS Infectious Diseases</i> , 2017, 3, 5-17.	3.8	83
9	Identification of aminopyrimidine-sulfonamides as potent modulators of Wag31-mediated cell elongation in mycobacteria. <i>Molecular Microbiology</i> , 2017, 103, 13-25.	2.5	22
10	A druggable secretory protein maturase of <i>Toxoplasma</i> essential for invasion and egress. <i>ELife</i> , 2017, 6, .	6.0	89
11	Micrococcin P1 - A bactericidal thiopeptide active against <i>Mycobacterium tuberculosis</i> . <i>Tuberculosis</i> , 2016, 100, 95-101.	1.9	23
12	The 8-Pyrrole-Benzothiazinones Are Noncovalent Inhibitors of DprE1 from <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 4446-4452.	3.2	85
13	Lead selection and characterization of antitubercular compounds using the Nested Chemical Library. <i>Tuberculosis</i> , 2015, 95, S200-S206.	1.9	26
14	Thiophenecarboxamide Derivatives Activated by EthA Kill <i>Mycobacterium tuberculosis</i> by Inhibiting the CTP Synthetase PyrG. <i>Chemistry and Biology</i> , 2015, 22, 917-927.	6.0	72
15	Whole cell screen based identification of spiropiperidines with potent antitubercular properties. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 3234-3245.	2.2	31
16	Bioluminescence for Assessing Drug Potency against Nonreplicating <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2015, 59, 4012-4019.	3.2	30
17	2-Carboxyquinoxalines Kill <i>Mycobacterium tuberculosis</i> through Noncovalent Inhibition of DprE1. <i>ACS Chemical Biology</i> , 2015, 10, 705-714.	3.4	116
18	Towards a new combination therapy for tuberculosis with next generation benzothiazinones. <i>EMBO Molecular Medicine</i> , 2014, 6, 372-383.	6.9	311

#	ARTICLE	IF	CITATIONS
19	Assessing the essentiality of the decaprenylphospho- $\beta$ -arabinofuranose pathway in <i>Mycobacterium tuberculosis</i> using conditional mutants. <i>Molecular Microbiology</i> , 2014, 92, 194-211.	2.5	76
20	Anticytolytic Screen Identifies Inhibitors of Mycobacterial Virulence Protein Secretion. <i>Cell Host and Microbe</i> , 2014, 16, 538-548.	11.0	83
21	Mechanism of Action of 5-Nitrothiophenes against <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 2944-2947.	3.2	31
22	Cross-Resistance between Clofazimine and Bedaquiline through Upregulation of MmpL5 in <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2014, 58, 2979-2981.	3.2	376
23	Rv2466c Mediates the Activation of TP053 To Kill Replicating and Non-replicating <i>Mycobacterium tuberculosis</i> . <i>ACS Chemical Biology</i> , 2014, 9, 1567-1575.	3.4	41
24	Synthesis and Antimycobacterial Activity of 2,1-Dihydropyridomycins. <i>ACS Medicinal Chemistry Letters</i> , 2013, 4, 264-268.	2.8	24
25	Streptomycin-Starved <i>Mycobacterium tuberculosis</i> 18b, a Drug Discovery Tool for Latent Tuberculosis. <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 5782-5789.	3.2	88
26	Genome-Wide Definition of the SigF Regulon in <i>Mycobacterium tuberculosis</i> . <i>Journal of Bacteriology</i> , 2012, 194, 2001-2009.	2.2	46
27	In Vitro Combination Studies of Benzothiazinone Lead Compound BTZ043 against <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2012, 56, 5790-5793.	3.2	112
28	Towards a new tuberculosis drug: pyridomycin – nature's isoniazid. <i>EMBO Molecular Medicine</i> , 2012, 4, 1032-1042.	6.9	175
29	Virulence Regulator EspR of <i>Mycobacterium tuberculosis</i> Is a Nucleoid-Associated Protein. <i>PLoS Pathogens</i> , 2012, 8, e1002621.	4.7	115
30	Tuberculosis drugs: new candidates and how to find more. <i>Future Microbiology</i> , 2011, 6, 617-633.	2.0	36
31	HIV protease inhibitors are substrates for OATP1A2, OATP1B1 and OATP1B3 and lopinavir plasma concentrations are influenced by SLCO1B1 polymorphisms. <i>Pharmacogenetics and Genomics</i> , 2010, 20, 112-120.	1.5	160
32	Leads for antitubercular compounds from kinase inhibitor library screens. <i>Tuberculosis</i> , 2010, 90, 354-360.	1.9	92
33	Sigma Factor F Does Not Prevent Rifampin Inhibition of RNA Polymerase or Cause Rifampin Tolerance in <i>Mycobacterium tuberculosis</i> . <i>Journal of Bacteriology</i> , 2010, 192, 5472-5479.	2.2	14
34	Simple Model for Testing Drugs against Nonreplicating <i>Mycobacterium tuberculosis</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2010, 54, 4150-4158.	3.2	117
35	Intracellular accumulation of efavirenz and nevirapine is independent of P-glycoprotein activity in cultured CD4 T cells and primary human lymphocytes. <i>Journal of Antimicrobial Chemotherapy</i> , 2009, 64, 1002-1007.	3.0	41
36	Intracellular boosting of darunavir using known transport inhibitors in primary PBMC. <i>British Journal of Clinical Pharmacology</i> , 2009, 68, 375-380.	2.4	15

#	ARTICLE	IF	CITATIONS
37	Cultured CD4T cells and primary human lymphocytes express hOATPs: intracellular accumulation of saquinavir and lopinavir. <i>British Journal of Pharmacology</i> , 2008, 155, 875-883.	5.4	40
38	Tacrine-induced liver damage: an analysis of 19 candidate genes. <i>Pharmacogenetics and Genomics</i> , 2007, 17, 1091-1100.	1.5	37
39	A rapid and sensitive HPLC-MS method for the detection of plasma and cellular rifampicin. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2007, 857, 76-82.	2.3	34
40	Differential drug susceptibility of intracellular and extracellular tuberculosis, and the impact of P-glycoprotein. <i>Tuberculosis</i> , 2007, 87, 248-255.	1.9	119
41	Low Levels of Pyrazinamide and Ethambutol in Children with Tuberculosis and Impact of Age, Nutritional Status, and Human Immunodeficiency Virus Infection. <i>Antimicrobial Agents and Chemotherapy</i> , 2006, 50, 407-413.	3.2	120
42	Down regulation of multidrug resistance protein-1 expression in patients with early rheumatoid arthritis exposed to methotrexate as a first disease-modifying antirheumatic drug. <i>Annals of the Rheumatic Diseases</i> , 2006, 65, 1390-1393.	0.9	20
43	Modulation of the intracellular accumulation of saquinavir in peripheral blood mononuclear cells by inhibitors of MRP1, MRP2, P-gp and BCRP. <i>Aids</i> , 2005, 19, 2097-2102.	2.2	84
44	Expression of P-glycoprotein, multidrug-resistance proteins 1 and 2 in CEM, CEM(VBL), CEM(E1000), MDCKII(MRP1) and MDCKII(MRP2) cell lines. <i>Aids</i> , 2003, 17, 2276-8.	2.2	4