

# Discovery of Novel Oral Protein Synthesis Inhibitors of Target Leucyl-tRNA Synthetase

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Discovery of Leucyladenylate Sulfamates as Novel Leucyl-tRNA Synthetase (LRS)-Targeted Mammalian Target of Rapamycin Complex 1 (mTORC1) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 10322-10328.	2.9	15
2	Structure-Based Targeting of Orthologous Pathogen Proteins Accelerates Antiparasitic Drug Discovery. <i>ACS Infectious Diseases</i> , 2017, 3, 281-292.	1.8	13
3	Targeting <i>Toxoplasma gondii</i> CPSF 3 as a new approach to control toxoplasmosis. <i>EMBO Molecular Medicine</i> , 2017, 9, 385-394.	3.3	61
4	Discovery of a Potent and Specific <i>M. tuberculosis</i> Leucyl-tRNA Synthetase Inhibitor: (S)-3-(Aminomethyl)-4-chloro-7-(2-hydroxyethoxy)benzo[c][1,2]oxaborol-1(3H)-ol (GSK656). <i>Journal of Medicinal Chemistry</i> , 2017, 60, 8011-8026.	2.9	118
5	Priming the tuberculosis drug pipeline: new antimycobacterial targets and agents. <i>Current Opinion in Microbiology</i> , 2018, 45, 39-46.	2.3	40
6	In Vitro and In Vivo Activities of DS86760016, a Novel Leucyl-tRNA Synthetase Inhibitor for Gram-Negative Pathogens. <i>Antimicrobial Agents and Chemotherapy</i> , 2018, 62, .	1.4	18
7	Hit Generation in TB Drug Discovery: From Genome to Granuloma. <i>Chemical Reviews</i> , 2018, 118, 1887-1916.	23.0	80
8	The Expanding Diversity of <i>Mycobacterium tuberculosis</i> Drug Targets. <i>ACS Infectious Diseases</i> , 2018, 4, 696-714.	1.8	60
9	New antituberculous drugs derived from natural products: current perspectives and issues in antituberculous drug development. <i>Journal of Antibiotics</i> , 2018, 71, 15-25.	1.0	20
10	A continuous assay for monitoring the synthetic and proofreading activities of multiple aminoacyl-tRNA synthetases for high-throughput drug discovery. <i>RNA Biology</i> , 2018, 15, 659-666.	1.5	11
11	An automated high-throughput system for phenotypic screening of chemical libraries on <i>C. elegans</i> and parasitic nematodes. <i>International Journal for Parasitology: Drugs and Drug Resistance</i> , 2018, 8, 8-21.	1.4	71
12	The 7-phenyl benzoxaborole series is active against <i>Mycobacterium tuberculosis</i> . <i>Tuberculosis</i> , 2018, 108, 96-98.	0.8	22
13	The trypanocidal benzoxaborole AN7973 inhibits trypanosome mRNA processing. <i>PLoS Pathogens</i> , 2018, 14, e1007315.	2.1	53
14	Binding Assessment of Monosaccharide-Boronic Acid Complexes via Tandem Mass Spectrometry. <i>ChemistrySelect</i> , 2018, 3, 8193-8198.	0.7	1
15	The present state of the tuberculosis drug development pipeline. <i>Current Opinion in Pharmacology</i> , 2018, 42, 81-94.	1.7	70
17	Boron in drug design: Recent advances in the development of new therapeutic agents. <i>European Journal of Medicinal Chemistry</i> , 2019, 179, 791-804.	2.6	154
18	Progress and challenges in aminoacyl-tRNA synthetase-based therapeutics. <i>Journal of Biological Chemistry</i> , 2019, 294, 5365-5385.	1.6	103
19	Recent development of leucyl-tRNA synthetase inhibitors as antimicrobial agents. <i>MedChemComm</i> , 2019, 10, 1329-1341.	3.5	25

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20	Opportunities for Overcoming Mycobacterium tuberculosis Drug Resistance: Emerging Mycobacterial Targets and Host-Directed Therapy. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2868.	1.8	47
21	First-Time-in-Human Study and Prediction of Early Bactericidal Activity for GSK3036656, a Potent Leucyl-tRNA Synthetase Inhibitor for Tuberculosis Treatment. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 63, .	1.4	50
22	Drug-resistance in <i>Mycobacterium tuberculosis</i> : where we stand. <i>MedChemComm</i> , 2019, 10, 1342-1360.	3.5	57
23	Antibacterial Activity and Mode of Action of a Sulfonamide-Based Class of Oxaborole Leucyl-tRNA-Synthetase Inhibitors. <i>ACS Infectious Diseases</i> , 2019, 5, 1231-1238.	1.8	26
24	Boron-Pleuromutilins as Anti- <i>Wolbachia</i> Agents with Potential for Treatment of Onchocerciasis and Lymphatic Filariasis. <i>Journal of Medicinal Chemistry</i> , 2019, 62, 2521-2540.	2.9	35
25	Optimization of Methionyl tRNA-Synthetase Inhibitors for Treatment of <i>Cryptosporidium</i> Infection. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 63, .	1.4	37
26	New Drugs for the Treatment of Tuberculosis. <i>Clinics in Chest Medicine</i> , 2019, 40, 811-827.	0.8	33
27	Roles of aminoacyl-tRNA synthetases in immune regulation and immune diseases. <i>Cell Death and Disease</i> , 2019, 10, 901.	2.7	58
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29	Impact of Target-Based Drug Design in Anti-bacterial Drug Discovery for the Treatment of Tuberculosis. <i>Challenges and Advances in Computational Chemistry and Physics</i> , 2019, , 307-346.	0.6	3
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35	Aminoacyl-tRNA synthetases as drug targets. <i>The Enzymes</i> , 2020, 48, 321-350.	0.7	7
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37	Synthesis, Characterization, and Biological Evaluation of Novel 7-Oxo-7H-thiazolo[3,2-b]-1,2,4-triazine-2-carboxylic Acid Derivatives. <i>Molecules</i> , 2020, 25, 1307.	1.7	9

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38	Antibiotics in the clinical pipeline in October 2019. <i>Journal of Antibiotics</i> , 2020, 73, 329-364.	1.0	188
39	<i>In Vitro</i> Susceptibility Testing of GSK656 against <i>Mycobacterium</i> Species. <i>Antimicrobial Agents and Chemotherapy</i> , 2020, 64, .	1.4	4
40	Molecular basis of the multifaceted functions of human leucyl-tRNA synthetase in protein synthesis and beyond. <i>Nucleic Acids Research</i> , 2020, 48, 4946-4959.	6.5	11
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44	A Leucyl-tRNA Synthetase Inhibitor with Broad-Spectrum Antimycobacterial Activity. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	23
45	Spontaneous Selection of <i>Cryptosporidium</i> Drug Resistance in a Calf Model of Infection. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	12
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49	Chemical Classes Presenting Novel Antituberculosis Agents Currently in Different Phases of Drug Development: A 2010–2020 Review. <i>Pharmaceuticals</i> , 2021, 14, 461.	1.7	31
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62	Boron-containing small molecules as antiparasitic agents. , 2022, , 155-201.		0
63	Deep learning-driven prediction of drug mechanism of action from large-scale chemical-genetic interaction profiles. <i>Journal of Cheminformatics</i> , 2022, 14, 12.	2.8	9
64	Investigate Natural Product Indolmycin and the Synthetically Improved Analogue Toward Antimycobacterial Agents. <i>ACS Chemical Biology</i> , 2022, 17, 39-53.	1.6	10
65	Identification of novel mycobacterium tuberculosis leucyl-tRNA synthetase inhibitor using a knowledge-based computational screening approach. <i>Journal of King Saud University - Science</i> , 2022, 34, 102032.	1.6	3
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67	A Novel Leucyl-tRNA Synthetase Inhibitor, MRX-6038, Expresses Anti-Mycobacterium abscessus Activity <i>&lt;i&gt;In Vitro&lt;/i&gt;</i> and <i>&lt;i&gt;In Vivo&lt;/i&gt;</i> . <i>Antimicrobial Agents and Chemotherapy</i> , 2022, 66, .	1.4	6
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83	Targeting <i>Mycobacterium tuberculosis</i> iron-scavenging tools: a recent update on siderophores inhibitors. RSC Medicinal Chemistry, 2023, 14, 1885-1913.	1.7	1
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