Bedaquiline

Cat. No.:	HY-14881				
CAS No.:	843663-66-1				
Molecular Formula:	C ₃₂ H ₃₁ BrN ₂ O ₂				
Molecular Weight:	555.5				
Target:	Bacterial; Antibiotic				
Pathway:	Anti-infection				
Storage:	Powder	-20°C	3 years		
		4°C	2 years		
	In solvent	-80°C	6 months		
		-20°C	1 month		

SOLVENT & SOLUBILITY

	Preparing Stock Solutions	Mass Solvent Concentration	1 mg	5 mg	10 mg		
		1 mM	1.8002 mL	9.0009 mL	18.0018 mL		
		5 mM	0.3600 mL	1.8002 mL	3.6004 mL		
		10 mM	0.1800 mL	0.9001 mL	1.8002 mL		
	Please refer to the solubility information to select the appropriate solvent.						
Solu 2. Add		. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 0.5 mg/mL (0.90 mM); Clear solution					
		 Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 0.5 mg/mL (0.90 mM); Suspended solution; Need ultrasonic 					
		one by one: 10% DMSO >> 90% corn oil ng/mL (0.90 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	Bedaquiline (TMC207) is a diarylquinoline agent and inhibits Mycobacterium tuberculosis (Mtb) F1FO-ATP synthase through targeting of both the c- and the ε-subunit ^[1] . Bedaquiline has uncoupler activity. Bedaquiline is used for the multi-agent resistant tuberculosis ^[2] .			
IC ₅₀ & Target	Mtb F1FO-ATP synthase ^[1]			
In Vitro	Bedaquiline inhibits the growth of TDR M. tuberculosis strains, with MIC values ranging from 0.125 to 0.5 mg/L ^[2] .			

Product Data Sheet

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Among slowly growing mycobacteria (SGM), bedaquiline exhibits the highest activity against Mycobacterium avium with MIC ₅₀ and MIC₉₀ values of 0.03 and 16 mg/L, respectively. Among rapidly growing mycobacteria (RGM), Mycobacterium abscessus subsp. abscessus (M. abscessus) and Mycobacterium abscessus subsp. massiliense (M. massiliense) seem more susceptible to bedaquiline than Mycobacterium fortuitum, with MIC₅₀ and MIC₉₀ values of 0.13 and >16 mg/L, respectively, for both species. Bedaquiline also shows moderate in vitro activity against NTM species^[3]. Bedaquiline has an excellent in vitro activity against Mycobacterium tuberculosis, including multidrug resistant M tuberculosis^[4].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Cell. 2023 May 11;186(10):2176-2192.e22.
- Nat Commun. 2021 Jun 21;12(1):3816.
- Eur J Med Chem. 6 August 2022, 114639.
- Mbio. 2021 Jun 1;e0108821.
- Front Cell Infect Microbiol. 2016 Nov 8;6:145. eCollection 2016.

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REFERENCES

[1]. Jang JC, et al. Bedaquiline susceptibility test for totally drug-resistant tuberculosis Mycobacterium tuberculosis. J Microbiol. 2017 Apr 20.

[2]. Pang Y, et al. In Vitro Activity of Bedaquiline against Nontuberculous Mycobacteria in China. Antimicrob Agents Chemother. 2017 Apr 24;61(5).

[3]. Chahine EB, et al. Bedaquiline: a novel diarylquinoline for multidrug-resistant tuberculosis. Ann Pharmacother. 2014 Jan;48(1):107-15.

[4]. Sarathy JP, et al. TBAJ-876 displays Bedaquiline-like mycobactericidal potency without retaining the parental drug's uncoupler activity. Antimicrob Agents Chemother. 2019 Nov 11.

Caution: Product has not been fully validated for medical applications. For research use only.

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