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产品名称: **MUT056399**
产品别名: **Fab-001**

生物活性:				
Description	MUT056399 (Fab-001) is a highly potent inhibitor of the FabI enzyme of both <i>S. aureus</i> and <i>E. coli</i> with 50% inhibitory concentration IC ₅₀ s of 12 nM and 58 nM, respectively.			
IC ₅₀ & Target	IC ₅₀ value: 12 nM (for <i>S. aureus</i>), 58 nM (for <i>E. coli</i>)[1]			
In Vitro	MUT056399 (Fab-001) is a highly potent new inhibitor of the FabI enzyme of both <i>Staphylococcus aureus</i> and <i>Escherichia coli</i> . MUT056399 is very active against <i>S. aureus</i> strains, including methicillin-susceptible <i>S. aureus</i> (MSSA), methicillin-resistant <i>S. aureus</i> (MRSA), linezolid-resistant, and multidrug-resistant strains, with MIC ₉₀ s between 0.03 and 0.12 µg/ml. MUT056399 is also active against coagulase-negative staphylococci, with MIC ₉₀ s between 0.12 and 4 µg/ml. MUT056399 is very active against the 118 <i>S. aureus</i> strains tested, including MSSA and MRSA isolates and linezolid-resistant and multidrug-resistant strains, with MIC ₉₀ s between ≤0.03 and 0.12 µg/ml.			
In Vivo	MUT056399 (Fab-001), administered subcutaneously, protected mice from a lethal systemic infection induced by MSSA, MRSA, and vancomycin-intermediate <i>S. aureus</i> strains (50% effective doses ranging from 19.3 mg/kg/day to 49.6 mg/kg/day). In the nonneutropenic murine thigh infection model, the same treatment with MUT056399 reduced the bacterial multiplication of MSSA and MRSA in the thighs of immunocompetent mice.			
Solvent&Solubility	In Vitro: DMSO: ≥ 31 mg/mL (105.70 mM) * "≥" means soluble, but saturation unknown.			
	Preparing Stock Solutions	Solvent / Mass / Concentration	1 mg	5 mg
		1 mM	3.4098 mL	17.0491 mL
		5 mM	0.6820 mL	3.4098 mL
		10 mM	0.3410 mL	1.7049 mL
	*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液。一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限: -80°C, 6 months; -20°C, 1 month。 -80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。			
References	[1]. Escaich S, et al. The MUT056399 inhibitor of FabI is a new antistaphylococcal compound. <i>Antimicrob Agents Chemother</i> . 2011 Oct;55(10):4692-7. [2]. Schiebel J, et al. An ordered water channel in <i>Staphylococcus aureus</i> FabI: unraveling the mechanism of substrate recognition and reduction. <i>Biochemistry</i> . 2015 Mar 17;54(10):1943-55.			