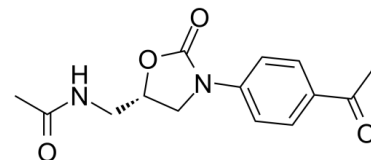


## Data Sheet

<b>Product Name:</b>	Dup-721
<b>Cat. No.:</b>	CS-0226367
<b>CAS No.:</b>	104421-21-8
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>16</sub> N <sub>2</sub> O <sub>4</sub>
<b>Molecular Weight:</b>	276.29
<b>Target:</b>	Antibiotic; Bacterial
<b>Pathway:</b>	Anti-infection
<b>Solubility:</b>	DMSO : 100 mg/mL (361.94 mM; Need ultrasonic)



### BIOLOGICAL ACTIVITY:

DuP-721 is a broad spectrum and orally active **antibacterial agent** against a variety of clinically susceptible and resistant bacteria, especially *M. tuberculosis*<sup>[1]</sup>. **In Vitro:** DuP-721 (1.5-4 µg/ml) inhibits equally the strains of *Mycobacterium tuberculosis* susceptible and resistant to conventional antituberculosis drug. And it does not show cross resistance to any of the anti-tuberculosis drugs tested [1].

DuP-721 is inactive against *M. avium* and *M. intracellulare* at 250 µg/ml. It inhibits *M. gordonae* and *M. fortuitum* at 3.9 µg/ml and *M. kansasii* and *M. scrofulaceum* at 1.95 µg/ml and 15.6 µg/ml, respectively<sup>[1]</sup>.

**In Vivo:** DuP-721 (oral gavage; 50-160 mg/kg) is protective against *M. tuberculosis* infection in mice. DuP-721 protects 100% of the infected animals at 50 mg/kg p.o. dose when administered daily for 17 days, and the same effect is observed at 160 mg/kg dose when the drug is administered only on day 11 and 12 post infection<sup>[1]</sup>.

### References:

[1]. Affiliatio, et al. Antimycobacterial activities of oxazolidinones: a review. *Infect Disord Drug Targets*. 2006 Dec;6(4):343-54.

### CAIndexNames:

Acetamide, N-[[[(5S)-3-(4-acetylphenyl)-2-oxo-5-oxazolidinyl]methyl]-

### SMILES:

O=C1N(C2=CC=C(C=C2)C(C)=O)C[C@@H](O1)CNC(C)=O

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

Tel: 610-426-3128

Fax: 888-484-5008

E-mail: sales@ChemScene.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA