



## **Sulfanilamide**

Catalog No: tcsc2221



## **Available Sizes**

**Size:** 500g

**Size:** 1000g



## **Specifications**

CAS No:

63-74-1

Formula:

 $C_6H_8N_2O_2S$ 

**Pathway:** 

Anti-infection

**Target:** 

Bacterial

**Purity / Grade:** 

>98%

**Solubility:** 

DMSO :  $\geq$  30 mg/mL (174.22 mM)

**Alternative Names:** 

Sulphanilamide

**Observed Molecular Weight:** 

172.2

## **Product Description**

Sulfanilamide is a competitive inhibitor for bacterial enzyme dihydropteroate synthetase with IC50 of 320  $\mu M$ .





Target: dihydropteroate synthetase; Antibacterial

Sulfanilamide containing the sulfonamide functional group displays inhibitory activity for dihydropteroate synthetase partially purified from Escherichia coli which normally uses para-aminobenzoic acid (PABA) for synthesizing the necessary folic acid acting as a coenzyme in the synthesis of purine, pyrimidine and other amino acids, exhibiting an IC 50 of 320  $\mu$ M for dihydropteroate synthetasea and Km of 2.5  $\mu$ M for PABA [1]. Sulfanilamide shows IC50 of 286.8  $\mu$ g/mL for recombinant S. cerevisiae strains with wild-type FOL1 genes, but the single mutation 55Trp to 55Ala or 57Pro to 57Ser within the putative active site of the fungal DHPS confers resistance to Sulfanilamide with IC50 of >800  $\mu$ g/mL [2]. Administration of Sulfanilamide with the dosage of 100 mg/kg/day is effective in the prevention of P. carinii infection in the immunosuppressed rat model. When the dosage of sulfaguanidine and Sulfanilamide reduced to 10 mg/kg/day, breakthrough P. carinii infection occurs in the rats [3].

$$O_{N} - NH_{2}$$
 $H_{2}N$ 

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