



Pemetrexed (disodium)

Catalog No: tcsc1298



Available Sizes

Size: 50mg

Size: 100mg

Size: 200mg



Specifications

CAS No:

150399-23-8

Formula:

 $C_{20}H_{19}N_5Na_2O_6$

Pathway:

Cell Cycle/DNA Damage; Autophagy

Target:

Antifolate; Autophagy

Purity / Grade:

>98%

Solubility:

H2O: 11.19 mg/mL (23.74 mM; Need ultrasonic and warming)

Alternative Names:

LY231514 disodium

Observed Molecular Weight:

471.37

Product Description





Pemetrexed disodium is a novel **antifolate**, the **K**_i values of the pentaglutamate of LY231514 are 1.3, 7.2, and 65 nM for inhibits thymidylate synthase (**TS**), dihydrofolate reductase (**DHFR**), and glycinamide ribonucleotide formyltransferase (**GARFT**), respectively.

IC50 & Target: Ki: 1.3 nM (TS), 7.2 nM (DHFR), 65 nM (GARFT)[1]

In Vitro: Pemetrexed (LY231514) disodium is a novel classical antifolate, the antitumor activity of which may result from simultaneous and multiple inhibition of several key folate-requiring enzymes via its polyglutamated metabolites. Pemetrexed (LY231514) is one of the best substrates that is known for the enzyme FPGS ($K_m = 1.6 \mu M$ and $V_{max}/K_m = 621$). It is likely that polyglutamation and the polyglutamated metabolites of LY231514 play profound roles in determining both the selectivity and the antitumor activity of this novel agent. Whereas LY23I5I4 only moderately inhibits TS ($K_i = 340 \mu M$, recombinant mouse), the pentaglutamate of LY23I5I4 is 100-fold more potent ($K_i = 3.4 \mu M$), making LY231514 one of the most potent folate-based TS inhibitors [1].

In Vivo: The group of mice treated with PC61 plus Pemetrexed demonstrates statistically longer survival than other groups. In a survival analysis, significantly better survival is observed in the group of mice treated with PC61 plus Pemetrexed compare with those treated with PC61 alone, rat IgG plus Pemetrexed, or no treatment^[2].

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