

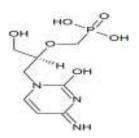
Cidofovir

ALTERNATE NAME: P-[[(1S)-2-(4-amino-2-oxo-1(2H)-pyrimidinyl)-1-(hydroxymethyl)ethoxy]methyl]-phosphonic acid

GS-0504 HPMPC Vistide

CATALOG #: B2682-10 10 mg B2682-50 50 mg

STRUCTURE:



MOLECULAR FORMULA: $C_8H_{14}N_3O_6P$

MOLECULAR WEIGHT: 279.19

CAS NUMBER: 113852-37-2

APPEARANCE: White to off-white solid

PURITY: ≥95%

SOLUBILITY: ~ 2 mg/ml (PBS, pH 7.2)

DESCRIPTION: Cidofovir is a selective inhibitor of viral DNA synthesis through the selective inhibition of viral DNA

polymerase. In vitro, cidofovir is converted to cidofovir diphosphate which incorporates into viral DNA, selectively inhibiting CMV replication and CMV DNA synthesis with IC50 values of 0.1 µM. Cidofovir has significantly long-lasting antiviral action because of the long half-life of its metabolites whose cellular

uptake is slow due to the presence of the negatively charged phosphonate group.

STORAGE TEMPERATURE: -20°C. Protect from moisture

HANDLING: Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of

exposure.

RELATED PRODUCTS:

Fotemustine (B2648) Fludarabine Phosphate (1763) Cytarabine (B2661) Carmofur (B2448)

DISCLAIMER: FOR RESEARCH USE ONLY! Not to be used on humans.