
Product Data Sheet

Product Name: LDC4297 hydrochloride

Cat. No.: GC66446

Chemical Properties

Cas. No. 2319747-14-1

Formula $C_{23}H_{29}ClN_8O$ M.Wt 468.98

Solubility Storage 4°C, away from moisture and light

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure

Background

LDC-4297 is an inhibitor of cyclin-dependent kinase 7 (CDK7; $IC_{50} = <5 \text{ nM}$).¹ It is selective for CDK7 over CDK4, CDK6, and CDK9 ($IC_{50}s = >10, >10, \text{ and } 1.71 \mu\text{M}$), however, it also inhibits CDK2 and CDK1 ($IC_{50}s = 6.4 \text{ and } 53.7 \text{ nM}$, respectively). LDC-4297 (10-100 nM) induces apoptosis in A549, HeLa, and HCT116 cancer cells in a concentration-dependent manner. It inhibits human cytomegalovirus (HCMV) replication in human fibroblasts ($EC_{50} = 24.5 \text{ nM}$).² LDC-4297 also reduces replication of *Herpesviridae*, *Adenoviridae*, *Poxviridae*, *Retroviridae*, and *Orthomyxoviridae* family viruses ($EC_{50}s = 0.02\text{-}1.13 \mu\text{M}$).

1. Kelso, T.W.R., Baumgart, K., Eickhoff, J., et al. Cyclin-dependent kinase 7 controls mRNA synthesis by affecting stability of preinitiation complexes, leading to altered gene expression, cell cycle progression, and survival of tumor cells. *Mol. Cell. Biol.* 34(19):3675-3688(2014)
2. Hutterer, C., Eickhoff, J., Milbradt, J., et al. A novel CDK7 inhibitor of the pyrazolotriazine class exerts broad-spectrum antiviral activity at nanomolar concentrations. *Antimicrob. Agents Chemother.* 59(4):2062-2071(2015)

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

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