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**Product Data Sheet**

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Product Name: Lagociclovir (MIV-210)

Cat. No.: GC32287

**Chemical Properties**

Cas. No. 92562-88-4

SMILES O=C(N=C(N)N1)C2=C1N([C@@H]3O[C@H](CO)[C@@H](F)C3)C=N2Formula  $C_{10}H_{12}FN_5O_3$  M.Wt 269.23

Solubility Soluble in DMSO Storage Store at -20°C

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**

Lagociclovir (MIV-210) is a prodrug of 3'-fluoro-2',3'-dideoxyguanosine with high oral bioavailability in humans and potent activity against HBV. IC50 value: Target: Anti-HBV compound Oral administration of MIV-210 at 20 or 60 mg/kg of body weight/day induced a rapid virological response in chronically infected woodchucks, reducing serum WHV DNA levels by 4.75 log<sub>10</sub> and 5.72 log<sub>10</sub>, respectively, in 2 weeks. Further, a daily dose of 10 mg/kg decreased the serum WHV load 400-fold after 4 weeks of treatment, and a dose of 5 mg/kg/day was sufficient to maintain this antiviral effect during the following 6-week period. MIV-210 at 20 or 60 mg/kg/day reduced the liver WHV DNA load 200- to 2,500-fold from pretreatment levels and, importantly, led to a 2.0 log<sub>10</sub> drop in the hepatic content of WHV covalently closed circular DNA.

[1]. Michalak TI, et al. Profound antiviral effect of oral administration of MIV-210 on chronic hepadnaviral infection in a woodchuck model of hepatitis B. *Antimicrob Agents Chemother.* 2009 Sep;53(9):3803-14.

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

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