

Enasidenib, Mutant isocitrate dehydrogenase 2 (IDH2) inhibitor

Catalog	Unit
TBI4660-5MG	5 mg
TBI4660-25MG	25 mg

Product Details

Formal Name: 2-Methyl-1-[[4-[6(trifluoromethyl)-2-pyridinyl]-6-[[2-(trifluoromethyl)-4-pyridinyl]amino]-1,3,5-

triazin-2-yl]amino]-2-propanol **Alternate Names:** AG-221 **Molecular Formula:** C₁₉H₁₇F₆N₇O

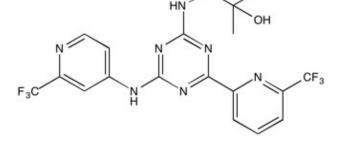
Formula Weight: 473.38 **CAS Number:** 1446502-11-9

Purity: >98%

Formulation: powder

Solubility: Soluble in DMSO (up to 25 mg/ml)

Storage: -20° C Stability: ≥ 1 year.



Applications

Mutant isocitrate dehydrogenase 2 (IDH2) inhibitor

Functions

Enasidenib is a potent (IC50's = 100 nM IDH2R140Q homodimer, 30 nM IDH2R140Q/WT heterodimer and 10 nM IDH2R172K/WT heterodimer) and selective inhibitor of mutant isocitrate dehydrogenase 2 (IDH2). It suppressed the production of the oncometabolite (R)-2-Hydroxyglutarate (a competitive inhibitor of α KG-dependent dioxygenases which leads to epigenetic dysregulation) and induced cellular differentiation in primary human IDH2 mutation-positive acute myeloid leukemia cells. Recently approved for clinical use by the FDA.

Application Procedures

First dissolved in DMSO (up to 25 mg/ml), then diluted to aqueous buffer. Solutions in DMSO may be stored at -20°C for up to 3 months.

For research use only.