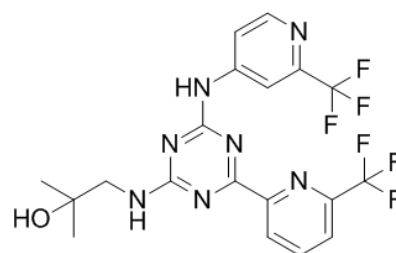


Product Name : Enasidenib
Cat. No. : PC-45529
CAS No. : 1446502-11-9
Molecular Formula : C₁₉H₁₇F₆N₇O
Molecular Weight : 473.375
Target : Isocitrate Dehydrogenase (IDH)
Solubility : 10 mM in DMSO



Biological Activity

Enasidenib (AG-221, CC-90007) is an orally available, selective, potent inhibitor of **mutant IDH2** with IC₅₀s of 100-400 nM for IDH2R140Q homodimer, IDH2R172K homodimer and IDH2WT/R140Q, IDH2WT/R172K.

Enasidenib (AG-221, CC-90007) is less potent for IDH2WT homodimer and IDH1WT homodimer.

Enasidenib (AG-221, CC-90007) suppresses 2HG production and induces cellular differentiation in primary human IDH2 mutation-positive AML cells ex vivo and in xenograft mouse models.

References

Yen K, et al. *Cancer Discov.* 2017 May;7(5):478-493.

Kats LM, et al. *Leukemia.* 2017 Apr 11. doi: 10.1038/leu.2017.84.

Shih AH, et al. *Cancer Discov.* 2017 May;7(5):494-505.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

E-mail: tech@probechem.com