# PRODUCT INFORMATION



# Cedazuridine

Item No. 38767

CAS Registry No.: 1141397-80-9

(4R)-2'-deoxy-2',2'-difluoro-Formal Name:

3,4,5,6-tetrahydro-uridine

Synonym:

MF:  $C_9H_{14}F_2N_2O_5$ 

268.2 FW: **Purity:** ≥95%

Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 vears

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

### **Laboratory Procedures**

Cedazuridine is supplied as a crystalline solid. A stock solution may be made by dissolving the cedazuridine in the solvent of choice, which should be purged with an inert gas. Cedazuridine is soluble in DMSO.

#### Description

Cedazuridine is an inhibitor of cytidine deaminase ( $IC_{50} = 0.4 \mu M$ ).<sup>1</sup> It increases plasma levels of decitabine via inhibition of its metabolism by cytidine deaminase in rhesus monkeys when administered at doses of 1 and 10 mg/kg. Cedazuridine (3 mg/kg) induces decreases in leukemic cell expansion in combination with orally administered 5-azacytidine (Item No. 11164), which is inactive as a single agent, in the bone marrow and spleen in a MOLM-13 acute myeloid leukemia (AML) mouse xenograft model.<sup>2</sup> Formulations containing cedazuridine in combination with decitabine have been used in the treatment of myelodysplastic syndrome (MDS).

# References

- 1. Ferraris, D., Duvall, B., Delahanty, G., et al. Design, synthesis, and pharmacological evaluation of fluorinated tetrahydrouridine derivatives as inhibitors of cytidine deaminase. J. Med. Chem. 57(6), 2582-2588 (2014).
- 2. Ramsey, H.E., Oganesian, A., Gorska, A.E., et al. Oral azacitidine and cedazuridine approximate parenteral azacitidine efficacy in murine model. Target Oncol. 15(2), 231-240 (2020).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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