# PRODUCT INFORMATION



## CeMMEC13

Item No. 20224

CAS Registry No.: 1790895-25-8

Formal Name: N-(2,3-dihydro-1,4-benzodioxin-

6-yl)-1,2-dihydro-1-methyl-2-oxo-

4-quinolinecarboxamide

MF:  $C_{19}H_{16}N_2O_4$ FW: 336.3 ≥98% **Purity:** 

UV/Vis.:  $\lambda_{\text{max}}$ : 211, 337 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥2 years

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



CeMMEC13 is supplied as a crystalline solid. A stock solution may be made by dissolving the CeMMEC13 in the solvent of choice, which should be purged with an inert gas. CeMMEC13 is soluble in organic solvents such as DMSO and dimethyl formamide. The solubility of CeMMEC13 in these solvents is approximately

CeMMEC13 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, CeMMEC13 should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. CeMMEC13 has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

CeMMEC13 is an isoquinolinone that selectively inhibits the second bromodomain of TAF1  $(IC_{50} = 2.1 \mu M)$ . It does not bind to bromodomains of BRD4, BRD9, or CREBBP. CeMMEC13 synergizes with (+)-JQ1 (Item No. 11187) to inhibit the proliferation of THP-1 and H23 lung adenocarcinoma cells. 1

### Reference

1. Sdelci, S., Lardeau, C.-H., Tallant, C., et al. Mapping the chemical chromatin reactivation landscape identifies BRD4-TAF1 cross-talk. Nat. Chem. Biol. 12(7), 504-510 (2016).

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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