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## Product Data Sheet

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Product Name: CBB1007 trihydrochloride

Cat. No.: GC35624

**Chemical Properties**

Cas. No. 2070015-03-9

SMILES NC(N1CCN(CC1)CC2=CC(C(OC)=O)=CC(C(N3CCN(CC3)C(C4=CC=C(C=C4)C(N)=N)=O)=O)=C2)=N.[H]Cl.[H]Cl.[H]ClFormula C<sub>27</sub>H<sub>37</sub>Cl<sub>3</sub>N<sub>8</sub>O<sub>4</sub>

M.Wt 643.99

Solubility DMSO: ≥ 54 mg/mL (83.85 mM)

Storage Store at -20°C

General For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath tips for a while. Stock solution can be stored below -20°C for several months.

Shipping Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice Condition upon request.

Structure **Background**

CBB1007 trihydrochloride is a cell-permeable amidino-guanidinium compound that acts as a potent, reversible and substrate competitive LSD1 selective inhibitor (IC<sub>50</sub> = 5.27 μM for hLSD1). IC<sub>50</sub> Value: 5.27 uM Target: hLSD1 CBB1007 efficiently can block LSD1-mediated demethylation of H3K4Me<sub>2</sub> and H3K4Me (IC<sub>50</sub> ≤ 5 μM) with no effect on H3K4Me<sub>3</sub> and H3K9Me<sub>2</sub>, and LSD2 and JARID1A activities. Increases H3K4Me<sub>2</sub> and H3K4Me contents (IC<sub>50</sub> ≤ 5 μM), and causes activation of epigenetically suppressed CHRM4/M4-ArchR and SCN3A genes in F9 cells (IC<sub>50</sub> ≤ 3.74 μM). CBB1007 was Shown to preferentially arrest the growth of pluripotent tumors with minimal effect on non-pluripotent cancer or normal somatic cells (IC<sub>50</sub> ≥ 100 μM).

[1]. Wang J, et al. Novel histone demethylase LSD1 inhibitors selectively target cancer cells with pluripotent stem cell properties. Cancer Res. 2011 Dec 1;71(23):7238-49.

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

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