

**ND-630** 

Catalog #: 27071-2 Lot #: 170509

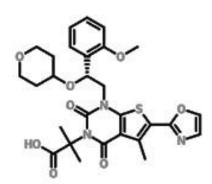
Size: 5 mg

**CAS Registry #:** 1434635-54-7

**Purity:** ≥99%

Chemical Formula: C<sub>28</sub>H<sub>31</sub>N<sub>3</sub>O<sub>8</sub>S

Molecular Weight: 569.63



**Description:** ND-630 (NDI 010976) is a highly specific, reversible inhibitor of ACC1 (IC50 = 2 nM) and ACC2 (IC50 = 6 nM). ND-630 interacts within the ACC phosphopeptide acceptor and dimerization site to prevent dimerization of ACC. Because the dimerization site is not conserved among other mammalian carboxylases, ND-630 lacks the ability to inhibit other mechanistically-related enzymes. In rats with diet-induced obesity, ND-630 reduced hepatic steatosis, improved insulin sensitivity, and reduced weight gain, suggesting ACC inhibition may be useful in treating metabolic disorders, including type 2 diabetes mellitus, and fatty liver disease.

Structure:

Appearance: White powder

HPLC (Area %): 94.7

**Storage/Stability:** Store at -20°C for up to a year. Dissolve in anhydrous DMSO and store DMSO stock solutions at 4°C for a couple of days or store stock solutions at -20°C in aliquots for up to a month. Avoid freeze/thaw cycles of solutions.

## References:

1. Harriman, G., et al., PNAS. 2016; **113 (13)**: E1796-E1805.