



## Product Specification Sheet

**Product Name:** ITD-1

**Catalog Number:** C4830

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**Technical information:**

Chemical Formula:  $C_{27}H_{29}NO_3$

CAS #: N/A

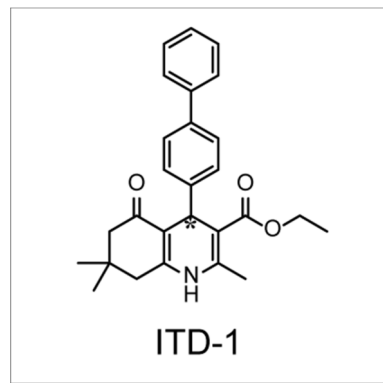
Molecular Weight: 415.52

Purity: >98%

Appearance: Yellow powder

Solubility: Soluble in DMSO up to 50 mM

Chemical Name: ethyl 4-([1,1'-biphenyl]-4-yl)-2,7,7-trimethyl-5-oxo-1,4,5,6,7,8-hexahydroquinoline-3-carboxylate



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**Storage:** Store solid powder at 4°C desiccated; Store DMSO solution at -20°C.

**Handling:**

- To make 10 mM stock solution, add 0.241 mL of DMSO for each 1mg of ITD-1.
- For DMSO solution, briefly spin the vial at 500 rpm in a 50 mL conical tube to ensure maximum sample recovery.

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**Biological Activity:** ITD-1 is a novel selective TGF- $\beta$  pathway-selective inhibitor that inhibits TGF- $\beta$  activity by selectively degrading the TGF- $\beta$ 2 receptor at the proteasome level. ITD-1 does not block the closely related Activin A signaling pathway. ITD-1 stimulates embryonic stem cells to differentiate into cardiomyocytes (IC<sub>50</sub> 0.4-0.8  $\mu$ M) by degrading the receptor and inhibiting intracellular signaling. ITD-1 has been formulated as a salt to increase stability and improve water solubility (~0.1 mg/mL) for ease of handling. As a salt, ITD-1 is chemically and metabolically stable and is non-cytotoxic. ITD-1 can be used to study a wide range of biological questions in cellular processing and TGF- $\beta$  signaling.

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**Reference:** 1. Willems E, et al. Small molecule-mediated TGF- $\beta$  type II receptor degradation promotes cardiomyogenesis in embryonic stem cells. Cell Stem Cell. 2012;11(2):242-52. PMID: [22862949](https://pubmed.ncbi.nlm.nih.gov/22862949/)

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