

## **IRAK-1-4 Inhibitor I**

## Catalog No: tcsc0704

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

**Size:** 100mg

Specifications

#### CAS No:

509093-47-4

#### Formula:

 $C_{20}H_{21}N_5O_4$ 

#### Pathway:

Immunology/Inflammation;Protein Tyrosine Kinase/RTK

## Target:

IRAK;IRAK

## Purity / Grade:

>98%

### **Solubility:** 10 mM in DMSO

#### **Alternative Names:**

IRAK-1/4 Inhibitor I

# **Observed Molecular Weight:** 395.41

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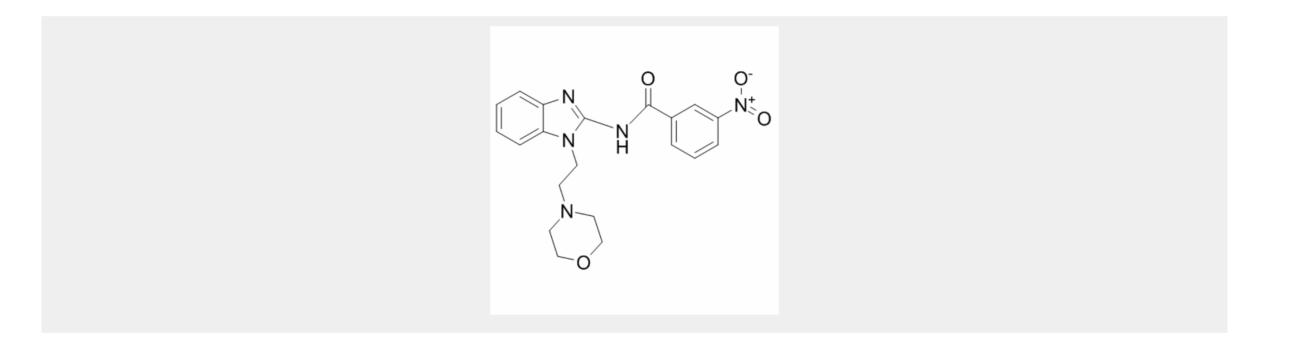


## **Product Description**

IRAK-1-4 Inhibitor I is a dual inhibitor of **IRAK4** and **IRAK1** with **IC**<sub>50</sub> of 0.2  $\mu$ M and 0.3  $\mu$ M, respectively.

IC50 & Target: IC50: 0.2 μM (IRAK-4), 0.3 μM (IRAK-1)<sup>[1]</sup>

*In Vitro:* IRAK-1-4 Inhibitor I has IC<sub>50</sub> greater than the highest concentration tested (10  $\mu$ M) against a panel of 27 other kinases, including the most closely homologous (outside of the IRAK family) Lck and pp60<sup>SRC</sup>. Additionally, IRAK-1-4 Inhibitor I does not show any signs of cytotoxicity in a 72 h proliferation assay in HeLa cells (ED<sub>50</sub>>30  $\mu$ M). Significant inhibition of IRAK-1 is observed with IRAK-1-4 Inhibitor I (IRAK-1 IC<sub>50</sub>=0.3  $\mu$ M)<sup>[1]</sup>. IRAK-1/4 inhibitor eliminates the LPS-induced increases in Bcl10, NF- $\kappa$ B, and IL-8. IRAK-1/4 mediates LPS-induced IL-8 activation and functions upstream of Bcl10. The LPS-induced increase in Bcl10 declines by 73% (from 5.18±0.22 to 2.36±0.08 ng/mL), and the IL-8 response decline by 60% (from 2.64±0.31 to 1.14±0.08 ng/mL)<sup>[2]</sup>.



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