

## **Product** Data Sheet

Molecular Weight: 526.62

Target: Free Fatty Acid Receptor

Pathway: GPCR/G Protein

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

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## **BIOLOGICAL ACTIVITY**

Description	TP-051 is a potent FFAR1 agonist with an $K_i$ value of 16 nM for human FFAR1. TP-051 can increase insulin secretion in rat insulinoma cells. TP-051 can be used to research type 2 diabetes <sup>[1]</sup> .
IC <sub>50</sub> & Target	$K_i$ : 16 nM (human FFAR1) $^{[1]}$
In Vitro	TP-051 (compound 31) (0.01-10 $\mu$ M; 2 h; INS-1 cells) augments insulin secretion in a dose-dependent manner in the presence of 11 mM glucose and statistically significant increases in insulin secretion were observed at doses of above 0.1 $\mu$ M of the compound [1].  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Mikami S, et al. Discovery of phenylpropanoic acid derivatives containing polar functionalities as potent and orally bioavailable G protein-coupled receptor 40 agonists for the treatment of type 2 diabetes. J Med Chem. 2012 Apr 26;55(8):3756-76.

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

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Inhibitors