**Proteins** 



## Pifithrin-β

Pathway:

Cat. No.: HY-16702 CAS No.: 60477-34-1 Molecular Formula:  $C_{16}H_{16}N_{2}S$ Molecular Weight: 268.38 Target: MDM-2/p53

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

Analysis.

**Apoptosis** 

## **BIOLOGICAL ACTIVITY**

Description	Pifithrin- $\beta$ (PFT $\beta$ ) is a potent p53 inhibitor with an IC $_{50}$ of 23 $\mu$ M.
IC <sub>50</sub> & Target	IC50: 23 μM (p53) <sup>[1]</sup>
In Vitro	Pifithrin- $\alpha$ , an inhibitor of the p53 protein, is regarded as a lead compound for cancer and neurodegenerative disease therapy. Pifithrin- $\alpha$ is very unstable in culture medium and rapidly converts to its condensation product pifithrin- $\beta$ (PFT $\beta$ ), the N-acetyl derivative <sup>[2]</sup> . After 24 h, the viability assay shows that the pretreatments with 1 and 10 $\mu$ M pifithrin- $\beta$ exerts neuroprotective effects <sup>[3]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

- Cell Commun Signal. 2022 Sep 5;20(1):96.
- Front Immunol. 2020 Feb 20;8:75.
- Cell Biol Toxicol. 2022 Jan 13.
- · Life Sci. 2021 Jun 7;280:119698.
- Front Cell Dev Biol. 2020 Jul 29;8:703.

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## **REFERENCES**

[1]. Christodoulou MS, et al. Synthesis and biological evaluation of imidazolo[2,1-b]benzothiazole derivatives, as potential p53 inhibitors. Bioorg Med Chem. 2011 Mar 1;19(5):1649-57.

[2]. Fernández-Cruz ML, et al. Biological and chemical studies on aryl hydrocarbon receptor induction by the p53 inhibitor pifithrin-α and its condensation product pifithrinβ. Life Sci. 2011 Apr 25;88(17-18):774-83.

[3]. Da Pozzo E, et al. p53 functional inhibitors behaving like pifithrin-β counteract the Alzheimer peptide non-β-amyloid component effects in human SH-SY5Y cells. ACS

Chem Neurosci. 2014 May 21;5(5):390-9.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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