

Cyclo(L-Phe-trans-4-hydroxy-L-Pro)

Chemical Properties

CAS No.:	118477-06-8
Formula:	C ₁₄ H ₁₆ N ₂ O ₃
Molecular Weight:	N/A
Appearance:	N/A
Storage:	0-4°C for short term (days to weeks), or -20°C for long term (months).

Biological Description

Description	Cyclo(L-Phe-trans-4-hydroxy-L-Pro) is a natural product from <i>Phellinus igniarius</i> .
In vitro	We previously identified <i>Streptomyces griseus</i> as an anti-cancer agent (Kim et al., 2014). In this study, we isolated compounds from <i>S. griseus</i> and evaluated their anticancer effect and toxicity in vitro and in vivo. METHODS AND RESULTS: Preparative centrifugal partition chromatography (CPC) was used to obtain three compounds, cyclo(L-[4-hydroxyprolinyl]-L-leucine), Cyclo(L-Phe-trans-4-hydroxy-L-Pro) and phenethyl acetate (PA). We chose PA, which had the highest anticancer activity, as a target compound for further experiments. PA induced the formation of apoptotic bodies, DNA fragmentation, DNA accumulation in G0/G1 phase, and reactive oxygen species (ROS) formation. Furthermore, PA treatment increased Bax/Bcl-xL expression, activated caspase-3, and cleaved poly-ADP-ribose polymerase (PARP) in HL-60 cells. Simultaneous evaluation in vitro and in vivo, revealed that PA exhibited no toxicity in Vero cells and zebrafish embryos. CONCLUSIONS: We revealed, for the first time, that PA generates ROS, and that this ROS accumulation induced the Bcl signaling pathway.

Solubility Information

Solubility	< 1 mg/ml refers to the product slightly soluble or insoluble
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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. The storage conditions and period of the stock solution: - 80 °C for 6 months; - 20 °C for 1 month. Please use it as soon as possible.

Reference

1. Evaluation on Anticancer Effect Against HL-60 Cells and Toxicity in vitro and in vivo of the Phenethyl Acetate Isolated from a Marine Bacterium *Streptomyces griseus* Fisheries & Aquatic Science, 2015 , 18 (1) :35-44.

Inhibitors · Natural Compounds · Compound Libraries

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