

## Product Data Sheet

Product Name: PRDX1-IN-1

Cat. No.: GC73232

### Chemical Properties

Cas. No.

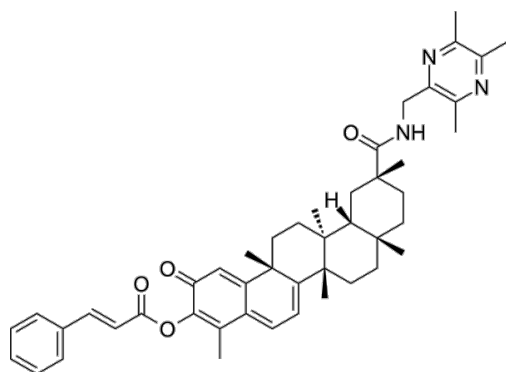
Formula  $C_{46}H_{55}N_3O_4$  M.Wt 713.95

Solubility DMSO : 100 mg/mL (140.07 mM; Storage -20°C  
Need ultrasonic)

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure



### Background

PRDX1-IN-1 is a selective inhibitor of PRDX1 with an IC<sub>50</sub> value of 0.164 μM. PRDX1-IN-1 can be used in researches related to cancer. PRDX1-IN-1 promotes intracellular ROS accumulation, and inhibits the proliferation, invasion and migration of cancer cells besides inducing apoptosis. PRDX1-IN-1 could be used in cancer research.

PRDX1-IN-1 inhibits the proliferation activities of the human lung cancer cells A549, lung cancer cell lines (LTEP-a-2 and H1975), human breast cancer cell line (MDA-MB-231), human hepatoma cell

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

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line (SK-Hep-1) with the IC<sub>50</sub> values of 1.92  $\mu$ M, 2.93  $\mu$ M, 1.99  $\mu$ M, 2.67  $\mu$ M, 2.42 $\mu$ M, respectively[1].PRDX1-IN-1 (compound 7e)(2  $\mu$ M or 4  $\mu$ M, 24 h) promotes intracellular ROS accumulation, and inhibits the invasion and migration of human lung cancer cells A549[1].PRDX1-IN-1 (2  $\mu$ M or 4  $\mu$ M, 24 h) induces the apoptosis of A549 cells[1].PRDX1-IN-1 (2  $\mu$ M or 4  $\mu$ M, 6 h) suppresses the key signaling pathways (AKT and ERK) and promotes the expression of apoptosis-related proteins (cleaved caspase-3/8 and cleaved PARP) in A549 cells[1].

PRDX1-IN-1 (0.5 or 1 mg/kg, intraperitoneal(i.p.), 19 days, every day) inhibited tumor growth in a mouse model of lung cancer[1].

### References:

[1]. Ying Bai, et al. Development of novel celastrol-ligustrazine brids as potent peroxiredoxin 1 inhibitors against lung cancer. Eur J Med Chem. 2023, 259, 115656.

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