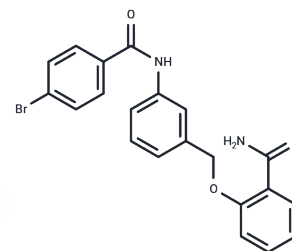


## PARP-1-IN-3

## Chemical Properties

CAS No. :	2976342-33-1
Formula:	C <sub>21</sub> H <sub>17</sub> BrN <sub>2</sub> O <sub>3</sub>
Molecular Weight:	425.28
Storage:	Powder: -20°C for 3 years   In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



## Biological Description

Description	PARP-1-IN-3 is a potent PARP-1 inhibitor that inhibits PARP-1 and PARP-2 with IC <sub>50</sub> values of 0.25 nM and 2.34 nM, respectively. PARP-1-IN-3 has potential anti-inflammatory activity, induces apoptosis, and arrests the cell cycle in the G <sub>2</sub> /M phase. PARP-1-IN-3 may be useful for the study of cancer-related diseases. cancer-related diseases.
Targets(IC <sub>50</sub> )	Apoptosis, Caspase, PARP
In vitro	PARP-1-IN-3 (compound 13f) demonstrates effective anticancer activity with IC <sub>50</sub> values of 0.30 μM, 2.83 μM, 33.69 μM, and 486.87 μM against HCT116, DLD-1, SW480, and NCM460 cells, respectively, following a 48-hour treatment[1]. In HCT116 cells, PARP-1-IN-3 (0.3-3 μM) inhibits colony formation and migration over 24-48 hours[1]. A 48-hour treatment with PARP-1-IN-3 (0.3-3 μM) induces the accumulation of DNA double-strand breaks in HCT116 cells[1]. Treatment with PARP-1-IN-3 (0.3-7.5 μM) for 48-72 hours results in cell cycle arrest at the G <sub>2</sub> /M phase, reduced mitochondrial membrane potential, and apoptosis in HCT116 cells[1].

## Solubility Information

Solubility	DMSO: 100 mg/mL (235.14 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.3514 mL	11.757 mL	23.5139 mL
5 mM	0.4703 mL	2.3514 mL	4.7028 mL
10 mM	0.2351 mL	1.1757 mL	2.3514 mL
50 mM	0.047 mL	0.2351 mL	0.4703 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Lu G, et, al. Discovery of novel benzamide derivatives bearing benzamidophenyl and phenylacetamidophenyl scaffolds as potential antitumor agents via targeting PARP-Eur J Med Chem. 2023 May 5;251:115243.

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