# Data Sheet (Cat.No.T18605)



## PROTAC ERα Degrader-2

Chemical Proper	ties
CAS No. :	1351169-29-3
Formula:	C42H61N5O8
Molecular Weight:	763.96 (e) <sup></sup> (-) (-) (-) (-) (-) (-) (-) (-) (-) (-)
Appearance:	no data available
Storage:	Powder: -20°C for 3 years   In solvent: -80°C for 1 year

### **Biological Description**

Description	PROTAC ERα Degrader-2 is composed of a cIAP1 ligand binding group, a linker, and an estrogen receptor α (ERα) binding group, serving as an ERα degrader. It achieves maximal ERα degradation in human mammary tumor MCF7 cells at a concentration of 30 μM. Degradation inducers that utilize cIAP1 are referred to as specific and non-genetic IAP-dependent protein erasers (SNIPERs)[1].
Targets(IC50)	Others,PROTACs

Solubility Information	®	
Solubility	DMSO: 200 mg/mL (261.79 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)	

#### Preparing Stock Solutions

.0	1mg	5mg	10mg
1 mM	1.309 mL	6.5448 mL	13.0897 mL
5 mM	0.2618 mL	1.309 mL	2.6179 mL
10 mM	0.1309 mL	0.6545 mL	1.309 mL
50 mM	0.0262 mL	0.1309 mL	0.2618 mL

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.

#### Reference

Scheepstra M, et al. Bivalent Ligands for Protein Degradation in Drug Discovery. Comput Struct Biotechnol J. 2019 Jan 25;17:160-176.

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