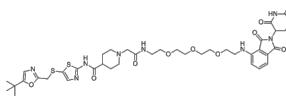


**Product Name** : THAL-SNS-032

**Synonyms** : THAL SNS 032;CDK9 PROTAC degrader

**Cat No.** : M13415

**CAS Number** : 2139287-33-3



**Molecular Formula** : C<sub>40</sub>H<sub>52</sub>N<sub>8</sub>O<sub>10</sub>S<sub>2</sub>

**Formula Weight** : 869.02

**Chemical Name** : N-((5-((5-(tert-butyl)oxazol-2-yl)methyl)thio)thiazol-2-yl)-1-(14-((2-(2,6-dioxopiperidin-3-yl)-1,3-dioxoisooindolin-4-yl)amino)-2-oxo-6,9,12-trioxa-3-azatetradecyl)piperidine-4-carboxamide

**Description** : THAL-SNS-032 is a novel CDK9 degrader PROTAC consisting of a CDK-binding SNS-032 ligand linked to a thalidomide derivative that binds the E3 ubiquitin ligase Cereblon (CRBN); efficiently induces complete CDK9 degradation at 250 nM (6h treatment), inhibits proliferation of MOLT4 cells at lower concentrations (IC<sub>50</sub> = 50 nM) than SNS-032 (IC<sub>50</sub>=173 nM), exhibits more potent inhibition of proliferation than SNS-032 across a panel of 11 different leukemia cancer cell lines; induces rapid degradation of CDK9 without affecting the levels of other SNS-032 targets, and has prolonged cytotoxic effects.

**Pathway** : PROTACs

**Target** : PROTAC

**Receptor** : PROTAC

**Solubility** : —

**SMILES** : O=C(C1CCN(CC(NCCOCCOCCOCNC2=CC=CC(C(N3C(CC4)C(NC4=O)=O)=O)=C2C3=O)=O)CC1)NC5=NC=C(SCC6=NC=C(C(C)(C)C)O6)S5

**Storage** : (-20°C)

**Stability** : ≥ 2 years

**Reference** :

1. Olson CM, et al. Nat Chem Biol. 2017 Dec 18. doi: 10.1038/nchembio.2538.