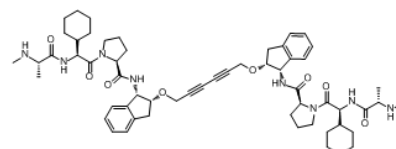


**Product Name** : AZD5582  
**Cat. No.** : PC-43119  
**CAS No.** : 1258392-53-8  
**Molecular Formula** : C<sub>58</sub>H<sub>78</sub>N<sub>8</sub>O<sub>8</sub>  
**Molecular Weight** : 1015.289  
**Target** : IAP  
**Solubility** : DMSO: ≥ 66.25 mg/mL



## Biological Activity

AZD5582 is a dimeric Smac mimetic, potent IAP antagonist that binds potently to the BIR3 domains of cIAP1, cIAP2 and XIAP with IC<sub>50</sub> of 15, 21 and 15 nM, respectively.

AZD5582 causes cIAP1 degradation and induces apoptosis in the MDA-MB-231 breast cancer cell line at subnanomolar concentrations in vitro.

AZD5582 induces cIAP1 degradation and caspase-3 cleavage within tumor cells and causes substantial tumor regressions in MDA-MB-231 xenograft-bearing mice.

## References

Hennessy EJ, et al. *J Med Chem.* 2013 Dec 27;56(24):9897-919.

Moon JH, et al. *Oncotarget.* 2015 Sep 29;6(29):26895-908.

Zhuang J, et al. *Pharmacol Res Perspect.* 2014 Dec;2(6):e00081.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

E-mail: tech@probechem.com