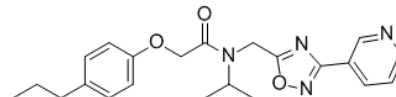


**Product Name** : PI-1840  
**Cat. No.** : PC-42948  
**CAS No.** : 1401223-22-0  
**Molecular Formula** : C<sub>22</sub>H<sub>26</sub>N<sub>4</sub>O<sub>3</sub>  
**Molecular Weight** : 394.4668  
**Target** : Proteasome  
**Solubility** : 10 mM in DMSO



### Biological Activity

PI-1840 is a potent, selective, noncovalent, reversible inhibitor for **proteasome chymotrypsin-like** (CT-L) activity with IC<sub>50</sub> of 27 nM, shows no effect on trypsin-like and peptidylglutamyl peptide hydrolyzing activities (IC<sub>50</sub>>100 μM). PI-1840 also displays 100-fold more selectivity for the constitutive proteasome over the immunoproteasome. PI-1840 induces the accumulation of proteasome substrates p27, Bax, and IκB-α, inhibits survival pathways and viability, and induces apoptosis in intact cancer cells, also sensitizes human cancer cells to the mdm2/p53 disruptor Nutlin. PI-1840 suppresses the growth in nude mice of human breast tumor xenografts.

### References

Kazi A, et al. *J Biol Chem*. 2014 Apr 25;289(17):11906-15.  
Ozcan S, et al. *J Med Chem*. 2013 May 23;56(10):3783-805.

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

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