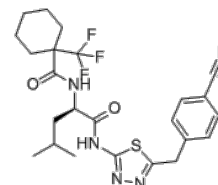


**Product Name** : BAY-805  
**Cat. No.** : PC-49862  
**CAS No.** : 2925481-88-3  
**Molecular Formula** : C<sub>24</sub>H<sub>28</sub>F<sub>3</sub>N<sub>5</sub>O<sub>2</sub>S  
**Molecular Weight** : 507.58  
**Target** : Deubiquitinase (DUB)  
**Solubility** : 10 mM in DMSO



## Biological Activity

BAY-805 (BAY805) is the first highly potent, selective and non-covalent **USP21** inhibitor with IC<sub>50</sub> of 6 nM and 2 nM in HTRF and Ub-rhodamine biochemical assays (hUSP21), respectively.

Cellular inhibition of USP21 with BAY-805 induced cellular NF-κB activation with an EC<sub>50</sub> of 17 nM in the NF-κB reporter assays.

BAY-805 shows the strong competition effect and effectively prevents ubiquitin binding.

BAY-805 exhibits strong ligand-induced protein stabilization resulting in a substantial thermal shift in the melt curve of USP21.

BAY-805 exhibits no antiproliferative effect was observed in Jurkat, Molm-13, A549, MDA-MB-231, and U2OS cell lines at 30 μM.

BAY-805 shows no significant effect on the activity of any of other deubiquitinating enzyme (DUBs) target.

## References

Fabian Göricke, et al. *J Med Chem.* 2023 Feb 20. doi: 10.1021/acs.jmedchem.2c01933.

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

E-mail: tech@probechem.com