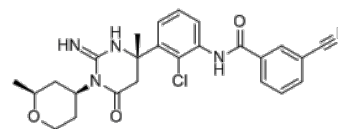


**Product Name** : UCB7362  
**Cat. No.** : PC-49827  
**CAS No.** : 2610631-17-7  
**Molecular Formula** : C<sub>25</sub>H<sub>26</sub>ClN<sub>5</sub>O<sub>3</sub>  
**Molecular Weight** : 479.97  
**Target** : Parasite  
**Solubility** : 10 mM in DMSO



## Biological Activity

UCB7362 (UCB 7362) is a potent, selective inhibitor of malaria parasite **plasmepsin X (PMX)** with IC<sub>50</sub> of 7 nM for *P. falciparum* (Pf PMX) and *Plasmodium vivax* (Pv PMX), SPR K<sub>d</sub> of 5.5 nM, also inhibits PMIX with IC<sub>50</sub> of 142 nM.

UCB7362 displays high selectivity against Cat D and Renin with an IC<sub>50</sub> of 3889 nM and >10,000 nM, respectively.

UCB7362 inhibits parasite growth across all strains tested regardless of geography and resistance profile asexual blood stage (ABS) assay of Pf parasites (3D7 strain, IC<sub>50</sub>=10 nM).

UCB7362 demonstrated no cross-resistance when tested against drug-resistant Plasmodium strains containing mutations generated with other compounds in development.

UCB7362 is estimated to achieve 9 log<sub>10</sub> unit reduction in asexual blood-stage parasites with once-daily dosing of 50 mg for 7 days in PfSCID mouse model.

## References

Martin A Lowe, et al. *J Med Chem.* 2022 Oct 27;65(20):14121-14143.

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

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