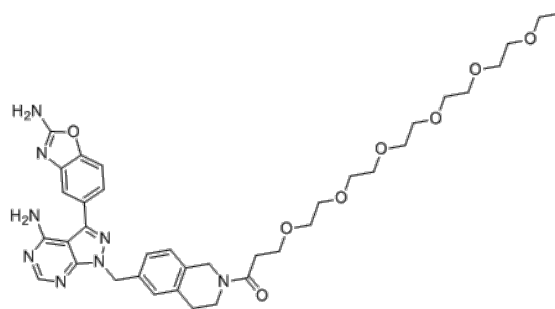


**Product Name** : RMC-5552  
**Cat. No.** : PC-49617  
**CAS No.** : 2382768-62-7  
**Molecular Formula** : C<sub>93</sub>H<sub>136</sub>N<sub>10</sub>O<sub>24</sub>  
**Molecular Weight** : 1778.16  
**Target** : mTOR  
**Solubility** : 10 mM in DMSO



### Biological Activity

RMC-5552 (RMC5552) is a potent, selective bi-steric inhibitor of **mTORC1** with cellular IC<sub>50</sub> of 0.48 nM for p4EBP1 inhibition, 40-fold selectivity over mTORC2.

RMC-5552 displays >53-fold selectivity over other lipid kinases.

Treatment of RMC-5552 resulted in a dose-dependent, prolonged, and profound inhibition of tumor p4EBP1 levels up to 48 h in Michigan Cancer Foundation-7 (MCF-7) breast cancer cell line, which bears an activating mutation in the p110 $\alpha$  catalytic subunit of Phosphoinositide 3-kinase (PIK3CAE545K).

RMC-5552 (10 mg/kg, ip, qw) reduced tumor volume in MCF-7 CDX models.

### References

G Leslie Burnett, et al. *J Med Chem.* 2023 Jan 12;66(1):149-169.

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

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