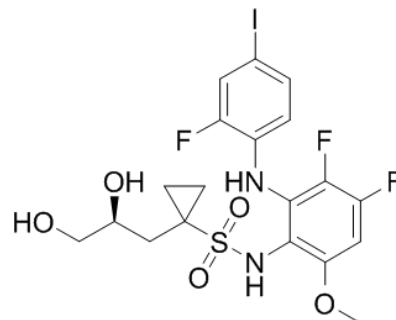


**Product Name** : Refametinib  
**Cat. No.** : PC-44028  
**CAS No.** : 923032-37-5  
**Molecular Formula** : C<sub>19</sub>H<sub>20</sub>F<sub>3</sub>IN<sub>2</sub>O<sub>5</sub>S  
**Molecular Weight** : 572.3372  
**Target** : MEK (MAP2K)  
**Solubility** : DMSO: ≥ 31 mg/mL



## Biological Activity

Refametinib (BAY 86-97661, RDEA-119) is a potent, non-ATP-competitive, highly selective, allosteric inhibitor of **MEK1/2** with IC<sub>50</sub> of 19/47 nM respectively.

Refametinib inhibits anchorage-dependent growth of human cancer cell lines harboring the gain-of-function V600E BRAF mutant with GI<sub>50</sub> of 67-89 nM.

Refametinib exhibits complete suppression of ERK phosphorylation at fully efficacious doses in mice and orally bioactive.

## References

Iverson C, et al. *Cancer Res.* 2009 Sep 1;69(17):6839-47.

Chang Q, et al. *BMC Cancer.* 2010 Sep 28;10:515.

Schmieder R, et al. *Neoplasia.* 2013 Oct;15(10):1161-71.

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

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