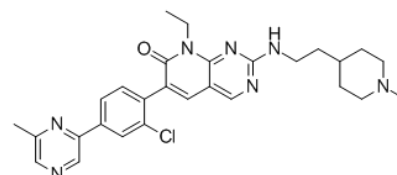


**Product Name** : FRAX-1036  
**Cat. No.** : PC-45644  
**CAS No.** : 1432908-05-8  
**Molecular Formula** : C<sub>28</sub>H<sub>32</sub>ClN<sub>7</sub>O  
**Molecular Weight** : 518.053  
**Target** : p21-activated Kinase (PAK)  
**Solubility** : DMSO: 5.3 mg/mL (Need warming)



## Biological Activity

FRAX1036 is a potent and selective inhibitor of **group I PAKs** (PAK1, Ki=23.3 nM; PAK2, Ki=72.4 nM).

FRAX1036 displays weak affinity on PAK4 (Ki = 2.4 μM).

FRAX1036 inhibits phosphorylation of MEK1-S298 and CRAF-S338 at 2.5-5 μM in MDA-MB-175 cells.

FRAX1036 inhibits MPNST cell growth in vitro and dramatically decreases local and metastatic MPNST growth in animal models combined with PD0325901.

## References

Ong CC, et al. *Breast Cancer Res.* 2015 Apr 23;17:59.

Mortazavi F, et al. *BMC Cancer.* 2015 May 9;15:381.

Semenova G, et al. *Oncogene.* 2017 May 22. doi: 10.1038/onc.2017.143.

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

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