

# **BAY-598**

## Catalog No: tcsc0015642

Available Sizes

Size: 1mg

Size: 5mg

**Specifications** 

**CAS No:** 1906919-67-2

Formula:

 $C_{22}H_{20}CI_2F_2N_6O_3$ 

**Pathway:** Epigenetics

Target:

Histone Methyltransferase

Purity / Grade:

>98%

#### **Solubility:** 10 mM in DMSO

# **Observed Molecular Weight:** 525.34

### **Product Description**

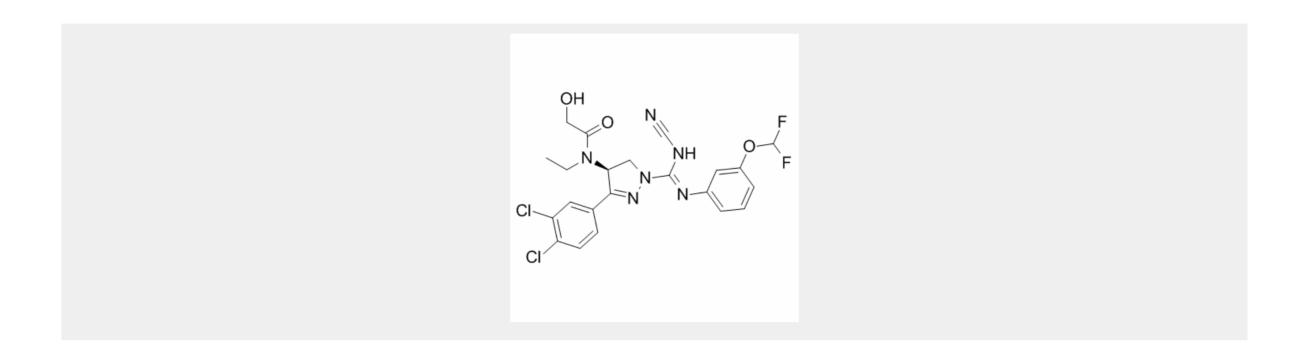
BAY-598 is selective small molecule inhibitor of SMYD2 .

IC50 & Target: SMYD2<sup>[1]</sup>

*In Vitro:* BAY-598 treatment blocks *in vitro* methylation of MAPKAPK3 by SMYD2 but has no activity against the SMYD2-related KMT SMYD3. BAY-598 treatment reduces the growth of *Kras;p53* mutant PDAC cells after 9 d in culture but has little impact on the growth



of Kras;p53;Smyd2 mutant cells<sup>[1]</sup>.



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