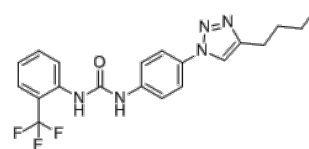


**Product Name** : FHP01  
**Cat. No.** : PC-49243  
**CAS No.** : 1985601-04-4  
**Molecular Formula** : C<sub>20</sub>H<sub>20</sub>F<sub>3</sub>N<sub>5</sub>O  
**Molecular Weight** : 403.409  
**Target** : Other Targets  
**Solubility** : 10 mM in DMSO



## Biological Activity

FHP01 (BA103) is a potent, small molecule inhibitor of **DDX3X helicase** activity with IC<sub>50</sub> of 0.3 μM in in vitro enzyme assays, exhibits very effective antiproliferative and killing activity against different breast cancer cell types (IC<sub>50</sub>=3.058 and 3.21 μM in MDA MB 468 and MDA MB 231, respectively).

FHP01 does not inhibit the ATPase activity of DDX3X and the helicase activity of DDX1 (IC<sub>50</sub>>100 μM).

FHP01 also inhibited WNT signaling, a key tumorigenic pathway already correlated to DDX3X functions in breast cancer model cell lines.

FHP01 inhibits ER+/PR+ (IC<sub>50</sub> = 12.43 and 10.62 μM in MCF7 and T47D cells, respectively) and HER2+ (IC<sub>50</sub> = 13.46 μM in SKBR3) cells, but lower in control MCF10A cells (IC<sub>50</sub> = 28.71 μM).

FHP01 (45 mg/kg, i.p. injection) suppresses growth of MDA MB 231 tumor xenografts in nude mice.

## References

Lisa Gherardini, et al. *Cancers (Basel)*. 2021 Sep 27;13(19):4830.

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

E-mail: tech@probechem.com