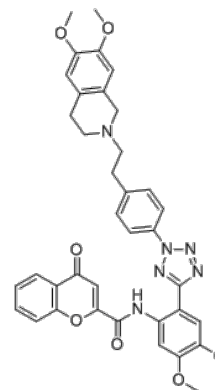


**Product Name** : Encequidar  
**Cat. No.** : PC-73218  
**CAS No.** : 849675-66-7  
**Molecular Formula** : C<sub>38</sub>H<sub>36</sub>N<sub>6</sub>O<sub>7</sub>  
**Molecular Weight** : 688.741  
**Target** : P-glycoprotein (P-gp)  
**Solubility** : 10 mM in DMSO



## Biological Activity

Encequidar (HM30181) is a potent selective inhibitor of **MDR1** (ABCB1, P-gp) with IC<sub>50</sub> of 0.63 nM.

Encequidar (HM30181) effectively blocked transepithelial transport of paclitaxel in MDCK monolayers.

Encequidar (HM30181) does not inhibit MRP1 (ABCC1), MRP2 (ABCC2), and MRP3 (ABCC3), and partially inhibited BCRP (ABCG2) only at very high concentrations.

Oral co-administration of paclitaxel and HM30181 showed a tumor-inhibitory strength equal or superior to that of intravenous paclitaxel in the xenograft model in nude mice.

## References

Kim TE, et al. *Br J Clin Pharmacol*. 2014 Sep;78(3):556-64.

Bauer F, et al. *Eur J Pharmacol*. 2012 Dec 5;696(1-3):18-27.

Kwak JO, et al. *Eur J Pharmacol*. 2010 Feb 10;627(1-3):92-8.

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

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