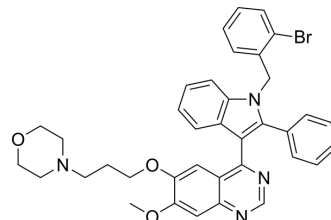


## YS-370

<b>Cat. No.:</b>	HY-132866		
<b>CAS No.:</b>	2470908-79-1		
<b>Molecular Formula:</b>	C <sub>37</sub> H <sub>35</sub> BrN <sub>4</sub> O <sub>3</sub>		
<b>Molecular Weight:</b>	663.6		
<b>Target:</b>	P-glycoprotein		
<b>Pathway:</b>	Membrane Transporter/Ion Channel		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 8.33 mg/mL (12.55 mM; ultrasonic and warming and heat to 80°C)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.5069 mL	7.5347 mL	15.0693 mL
5 mM	0.3014 mL	1.5069 mL	3.0139 mL
10 mM	0.1507 mL	0.7535 mL	1.5069 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

YS-370 (compound 44) is a potent, high selective, and orally active inhibitor of P-glycoprotein (P-gp). YS-370 stimulates the P-gp ATPase activity and has moderate inhibition against CYP3A4. YS-370 effectively reverses multidrug resistance (MDR) to paclitaxel and colchicine in SW620/AD300 and HEK293T-ABCB1 cells. YS-370 in combination with paclitaxel achieves much stronger antitumor activity<sup>[1]</sup>.

### REFERENCES

[1]. Yuan S, et al. Discovery of New 4-Indolyl Quinazoline Derivatives as Highly Potent and Orally Bioavailable P-Glycoprotein Inhibitors. *J Med Chem.* 2021;64(19):14895-14911.

---

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA