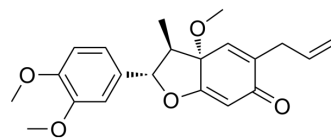


## Kadsurenone

Cat. No.:	HY-N10663
CAS No.:	95851-37-9
Molecular Formula:	C <sub>21</sub> H <sub>24</sub> O <sub>5</sub>
Molecular Weight:	356.41
Target:	Platelet-activating Factor Receptor (PAFR)
Pathway:	GPCR/G Protein
Storage:	-20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



### BIOLOGICAL ACTIVITY

#### In Vivo

Kadsurenone (i.v.; 20 and 30 mg/kg) undergoes hepatobiliary excretion but might not be regulated by P-glycoprotein (P-gp) transported system<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Sprague-Dawley rats <sup>[1]</sup>
Dosage:	0 and 30 mg/kg
Administration:	IV
Result:	Could go through hepatobiliary excretion.

### REFERENCES

[1]. Shu-Pei Huang, et al. Pharmacokinetics of kadsurenone and its interaction with cyclosporin A in rats using a combined HPLC and microdialysis system. J Chromatogr B Analyt Technol Biomed Life Sci. 2009 Jan 15;877(3):247-52.

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

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