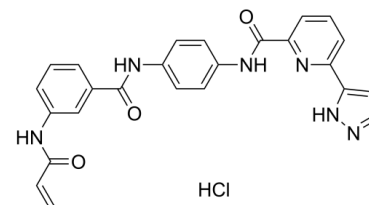


## Data Sheet

<b>Product Name:</b>	JH-X-119-01 (hydrochloride)
<b>Cat. No.:</b>	CS-7638
<b>CAS No.:</b>	2591344-30-6
<b>Molecular Formula:</b>	C <sub>25</sub> H <sub>21</sub> ClN <sub>6</sub> O <sub>3</sub>
<b>Molecular Weight:</b>	488.93
<b>Target:</b>	IRAK
<b>Pathway:</b>	Immunology/Inflammation
<b>Solubility:</b>	DMSO : 50 mg/mL (102.26 mM; Need ultrasonic); H <sub>2</sub> O : < 0.1 mg/mL (ultrasonic;warming;heat to 60°C) (insoluble)



### BIOLOGICAL ACTIVITY:

JH-X-119-01 hydrochloride is a potent and selective interleukin-1 receptor-associated kinases 1 (**IRAK1**) inhibitor. JH-X-119-01 hydrochloride ameliorates LPS-induced sepsis in mice<sup>[1]</sup>. *In Vitro*: Jh-X-119-01 (10 μM) decreases phosphorylation of NF-κB and mRNA levels of IL-6 and TNFα in LPS-treated macrophages in vitro. Jh-X-119-01 selectively inhibits IRAK1 phosphorylation<sup>[1]</sup>. *In Vivo*: Jh-X-119-01 improves survival and decreases immunopathies of LPS-challenged mice. Jh-X-119-01 increases survival of mice at the dose of 5 mg/kg body weight. Survival is further improved when the dose is increased to 10 mg/kg<sup>[1]</sup>.

### References:

[1]. Bin Pan, et al. Selective inhibition of interleukin-1 receptor-associated kinase 1 ameliorates lipopolysaccharide-induced sepsis in mice. *Int Immunopharmacol.* 2020 Aug;85:106597.

### CAIndexNames:

2-Pyridinecarboxamide, N-[4-[[3-[(1-oxo-2-propen-1-yl)amino]benzoyl]amino]phenyl]-6-(1H-pyrazol-3-yl)-, hydrochloride (1:1)

### SMILES:

O=C(C=C)NC1=CC=CC(C(=O)NC2=CC=C(NC(C3=NC(C4=CC=NN4)=CC=C3)=O)C=C2)=O=C1.Cl

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

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