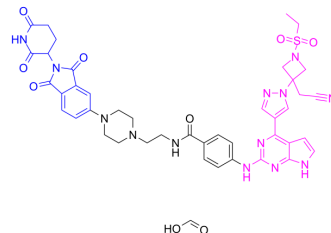


## SJ1008030 formic

<b>Cat. No.:</b>	HY-147330B		
<b>Molecular Formula:</b>	C <sub>43</sub> H <sub>45</sub> N <sub>13</sub> O <sub>9</sub> S		
<b>Molecular Weight:</b>	919.96		
<b>Target:</b>	PROTACs; JAK		
<b>Pathway:</b>	PROTAC; Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt		
<b>Storage:</b>	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 125 mg/mL (135.88 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.0870 mL	5.4350 mL	10.8700 mL
5 mM	0.2174 mL	1.0870 mL	2.1740 mL
10 mM	0.1087 mL	0.5435 mL	1.0870 mL

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

### BIOLOGICAL ACTIVITY

#### Description

SJ1008030 (compound 8) formic is a JAK2 PROTAC which selectively degrades JAK2. SJ1008030 formic inhibits MHH-CALL-4 cell growth with an IC<sub>50</sub> of 5.4 nM. SJ1008030 formic can be used for the research of leukemia<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

JAK2

#### In Vitro

SJ1008030 (compound 8) formic (72 h) shows activity in CRLF2r cell lines (MHH-CALL-4), with an IC<sub>50</sub> of 5.4 nM<sup>[1]</sup>.  
 SJ1008030 (0-4.3 μM; 72 h) formic degrades JAKs, GSPT1, and IKZF1 in a dose-dependent manner in MHH-CALL-4 cells<sup>[1]</sup>.  
 SJ1008030 (0-10 μM; 24h) formic dose-dependently degrades JAK2 and GSPT1 protein in xenograft bone marrow SJBALL021415 cells, indicating the inhibition of JAK-STAT signaling pathway<sup>[1]</sup>.  
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Chang Y, et al. Degradation of Janus kinases in CRLF2-rearranged acute lymphoblastic leukemia. *Blood*. 2021 Dec 9;138(23):2313-2326.

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

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