## YE6144

Cat. No.:	HY-150095		
Molecular Formula:	C <sub>21</sub> H <sub>27</sub> CIFN <sub>7</sub> O		
Molecular Weight:	447.94		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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## SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (223.24 mM; ultrasonic and warming and heat to 60°C)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	2.2324 mL	11.1622 mL	22.3244 mL	
		5 mM	0.4465 mL	2.2324 mL	4.4649 mL	
		10 mM	0.2232 mL	1.1162 mL	2.2324 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 10 mg/mL (22.32 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 10 mg/mL (22.32 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 10 mg/mL (22.32 mM); Clear solution					

BIOLOGICAL ACTIVITY				
Description	YE6144 is a prototypical interferon regulatory factor 5 (IRF5) inhibitor. YE6144 selectively suppresses IRF5 activity through inhibition of IRF5 phosphorylation <sup>[1]</sup> .			
IC <sub>50</sub> & Target	IRF5 <sup>[1]</sup>			
In Vitro	YE6144 (1 or 3 μM; 30 min) inhibits the phosphorylation of IRF5 in both human PBMCs and mouse splenocytes <sup>[1]</sup> . YE6144 (0-10 μM; 30 min) inhibits the production of type I IFNs with an IC <sub>50</sub> of approximately 0.09 μM in human HC PBMCs <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

## Product Data Sheet

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	Western Blot Analysis <sup>[1]</sup>			
	Cell Line:	Human HC PBMCs and mouse WT splenocytes		
	Concentration:	$1\mu\text{M}$ (PBMCs) and $3\mu\text{M}$ (splenocytes)		
	Incubation Time:	30 min		
	Result:	Inhibited the phosphorylation of IRF5.		
	RT-PCR <sup>[1]</sup>			
	Cell Line:	Mouse WT splenocytes		
	Concentration:	3 μМ		
	Incubation Time:	30 min		
	Result:	Induction of type I IFN genes, Ifnb1 and Ifna stimulated by TLR7 ligands or TLR9 ligands was remarkably weakened.		
In Vivo	YE6144 (40.0 mg/kg; s.c. MCE has not independer	YE6144 (40.0 mg/kg; s.c.; once) suppresses the progression of mouse systemic lupus erythematosus <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	NZB/W F1 mouse model of systemic lupus erythematosus (SLE) <sup>[1]</sup>		
	Dosage:	40.0 mg/kg		
	Administration:	Subcutaneous injection, once		
	Result:	Suppressed the exacerbation of autoantibody production. Splenomegaly and renal dysfunction were also suppressed by the treatment after disease onset.		

## REFERENCES

[1]. Ban T, et al. Genetic and chemical inhibition of IRF5 suppresses pre-existing mouse lupus-like disease. Nat Commun. 2021 Jul 19;12(1):4379.

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

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