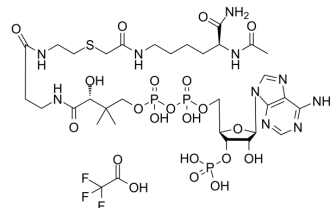


## Lys-CoA TFA

<b>Cat. No.:</b>	HY-131035
<b>Molecular Formula:</b>	C <sub>33</sub> H <sub>54</sub> F <sub>3</sub> N <sub>10</sub> O <sub>21</sub> P <sub>3</sub> S
<b>Molecular Weight:</b>	1108.82
<b>Target:</b>	Histone Acetyltransferase
<b>Pathway:</b>	Epigenetics
<b>Storage:</b>	-20°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

#### In Vitro

H<sub>2</sub>O : 70 mg/mL (63.13 mM; Need ultrasonic)

Concentration	Solvent	Mass	Preparing Stock Solutions		
			1 mg	5 mg	10 mg
1 mM			0.9019 mL	4.5093 mL	9.0186 mL
5 mM			0.1804 mL	0.9019 mL	1.8037 mL
10 mM			0.0902 mL	0.4509 mL	0.9019 mL

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

### BIOLOGICAL ACTIVITY

#### Description

Lys-CoA TFA is a selective p300 histone acetyltransferase (HAT) inhibitor (IC<sub>50</sub>=50-500 nM). Lys-CoA TFA displays >100-fold selectivity for p300 over PCAF (IC<sub>50</sub>=200 μM). Lys-CoA TFA inhibits p300 HAT activity-dependent transcriptional activation<sup>[1]</sup> [2].

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

p300  
50-500 nM (IC<sub>50</sub>)

#### In Vitro

Lys-CoA TFA (0.1 and 1 mM; 60 hours) induces a senescent phenotype in human normal melanocytes<sup>[3]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay<sup>[3]</sup>

Cell Line:	Normal human melanocytes
Concentration:	0.1 and 1 mM
Incubation Time:	60 hours

---

Result:	Almost completely abolished proliferation, induced high levels of SA- $\beta$ -Gal, and inhibited cyclin E expression in a dose-dependent manner.
---------	---

---

---

## REFERENCES

- [1]. Cebrat M, et al. Synthesis and analysis of potential prodrugs of coenzyme A analogues for the inhibition of the histone acetyltransferase p300. *Bioorg Med Chem.* 2003;11(15):3307-3313.
- [2]. Lau OD, et al. HATs off: selective synthetic inhibitors of the histone acetyltransferases p300 and PCAF. *Mol Cell.* 2000;5(3):589-595.
- 

**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