## FAPy-adenine

Cat. No.:	HY-113303		
CAS No.:	5122-36-1		
Molecular Formula:	$C_{5}H_{7}N_{5}O$		
Molecular Weight:	153.14		
Target:	Endogenous Metabolite		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

## SOLVENT & SOLUBILITY

In Vitro	DMSO : 17.86 mg/mL (116.63 mM; Need ultrasonic)						
Preparing Stock Solutions	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	6.5300 mL	32.6499 mL	65.2997 mL		
	5 mM	1.3060 mL	6.5300 mL	13.0599 mL			
	10 mM	0.6530 mL	3.2650 mL	6.5300 mL			
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent o Solubility: ≥ 2.08 n	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (13.58 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.79 mg/mL (11.69 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1.79 mg/mL (11.69 mM); Clear solution						

Biologiciterit				
Description	FAPy-adenine is an oxidized DNA base. Fapy-adenine shows an increased trend levels in the Alzheimer's disease brain. Oxidized nucleosides are biochemical markers for tumors, aging, and neurodegenerative diseases <sup>[1][2][3]</sup> .			
IC <sub>50</sub> & Target	Human Endogenous Metabolite			
In Vitro	In the absence of the external field the FAPy-adenine is able to form pairs with all four canonical nucleic acid bases. In contrast, in the presence of the external field the mispairing abilities of FAPy-adenine become insignificant since the most			





	stable dimers are formed with thymine <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	The nuclear DNA damage by oxygen-derived radicals is increased in Alzheimer's disease and support the concept that the brain is under increased oxidative stress in Alzheimer's disease <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Gabbita SP, et al. Increased nuclear DNA oxidation in the brain in Alzheimer's disease.

[2]. Cysewski P, et al. Theoretical description of the coding potential of diamino-5-formamidopyrimidines. Z Naturforsch C J Biosci. 1999 Mar-Apr;54(3-4):239-45.

[3]. Lee SH, et al. A rapid and sensitive method for quantitation of nucleosides in human urine using liquid chromatography/mass spectrometry with direct urine injection. Rapid Commun Mass Spectrom. 2004;18(9):973-7.

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

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