BAM 15

Cat. No.:	HY-110284		
CAS No.:	210302-17-3		
Molecular Formula:	$C_{16}H_{10}F_{2}N_{6}O$		
Molecular Weight:	340.29		
Target:	Mitochondrial Metabolism		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

SOLVENT & SOLUBILITY

Preparing Stock Solutions		Solvent Mass Concentration	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.9387 mL	14.6933 mL	29.3867 mL
	Stock Solutions	5 mM	0.5877 mL	2.9387 mL	5.8773 mL
		10 mM	0.2939 mL	1.4693 mL	2.9387 mL
	Please refer to the so	10 mM		1.4693 mL	2

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Description	BAM 15 is a mitochondrial protonophore uncoupler. BAM 15 is an oxidative phosphorylation (OXPHOS) uncoupler ^[1] .	
In Vitro	BAM 15 is able to increase O ₂ consumption across a broad dosing range without increasing ROS. BAM 15 and FCCP are structurally unrelated and it is observed that low doses of BAM 15 from 100 nM to 1 μM increase cellular O ₂ consumption rate (OCR) to a similar degree as FCCP, but higher concentrations from 1 μM to 50 μM reveal that BAM 15 is able to maintain uncoupled respiration at a high rate in a range of cell lines. BAM 15 is fully capable of increasing mitochondrial respiration in the presence of oligomycin and does so across a broader concentration range than FCCP in both myoblasts and hepatocytes. BAM 15 induces mitochondrial swelling, demonstrating that BAM 15 is a protonophore. BAM15-treated cells are more viable than FCCP-treated cells when administered across a broad dosing range up to 50 μM ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	



Product Data Sheet

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In Vivo	Compare to vehicle-treated mice, animals that receive BAM 15 are protected from kidney injury as indicated by lower	
	plasma creatinine levels at 24 and 48 h post-ischemia, reduced tubular necrosis, less depletion of brush border villi, less	
	obstruction of proximal tubules, and less immune cell infiltration ^[1] .	
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PROTOCOL	
FROTOCOL	
Kinase Assay ^[1]	Electron flow assays are performed. Briefly, 5 μg of mitochondrial protein in MAS is loaded into a 24-well tissue culture plate and centrifuged at 2000×g for 15 min at 4°C. Prior to the assay, mitochondria are incubated at 37°C for 10 mins in MAS containing 10 mM pyruvate, 2 mM malate, and 5 μM BAM 15 or FCCP. Rotenone (2 μM), succinate (10 mM), antimycin A (4 μ M), and N,N,N',N'-tetramethyl-p-phenylenediamine (TMPD, 100 μM) plus ascorbate (10 mM) are added sequentially as indicated in the figure. N=3 wells/plate of a representative of 3 plates ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
Cell Assay ^[1]	L6 cells are incubated with the fluorescent indicator of mitochondrial membrane potential tetramethylrhodamine (TMRM, 125 nM) or DMSO (1%) control for 30 min. The cells are then centrifuged for 5 min at 700×g and resuspended in unbuffered DMEM at a concentration of 1×10 ⁵ cells/mL. The cells are then treated with BAM 15 or DMSO (0.1%) for 10 min prior to flow cytometric analysis ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
Animal Administration ^[1]	Male mice (8-week old, C57BL/6) are used. Mice are i.p. injected with BAM 15 at 1 or 5 mg/kg, 1 h before kidney IR. Vehicle mice are also injected with the same solution BAM 15 is prepared with (3% DMSO in 50% PEG400) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Kenwood BM, et al. Identification of a novel mitochondrial uncoupler that does not depolarize the plasma membrane. Mol Metab. 2013 Nov 28;3(2):114-23.

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

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