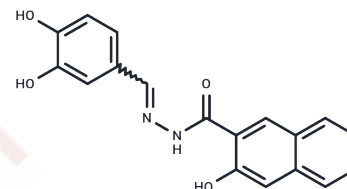


## Dynasore

## Chemical Properties

CAS No. :	304448-55-3
Formula:	C <sub>18</sub> H <sub>14</sub> N <sub>2</sub> O <sub>4</sub>
Molecular Weight:	322.31
Storage:	Powder: -20°C for 3 years   In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



## Biological Description

Description	Dynasore (Dynamamin Inhibitor I) is a highly selective, reversible small-molecule dynamamin inhibitor with an IC <sub>50</sub> of 15 μM. Dynasore also inhibits the mitochondrial dynamamin Drp1 without affecting other small GTPases. By inhibiting the GTPase activity of dynamamin, Dynasore blocks its function in membrane fission, thereby inhibiting clathrin-mediated endocytosis. Dynasore can be used in research on endocytosis, membrane transport, viral entry, and receptor signaling regulation.
Targets(IC50)	Autophagy,HSV,Dynamamin,Virus Protease
In vitro	<p><b>Methods:</b> Mouse bone marrow-derived macrophages were pretreated with 5, 10, 25, or 50 μM Dynasore for 1 hour, then stimulated with 100 ng/mL LPS for 24 hours. Supernatants and cell lysates were collected. Pro-inflammatory cytokine expression levels were detected by real-time quantitative PCR.</p> <p><b>Results:</b> Dynasore dose-dependently inhibited LPS-induced mRNA and protein levels of proinflammatory cytokines (TNF-α, IL-6, IL-12). [1]</p> <p><b>Methods:</b> HCLE cells (Fluo-4 loaded) were pretreated with 40 μM Dynasore or 1 μM YM-58483 for 30 min, followed by induction of oxidative stress with 1 mM tBHP. Cytoplasmic calcium changes were monitored in real-time via fluorescence intensity over 2 hours.</p> <p><b>Results:</b> Dynasore completely blocked tBHP-induced increases in cytoplasmic calcium. [2]</p> <p><b>Methods:</b> Horizontal brainstem slices containing the solitary tract and solitary tract nucleus were prepared from adult rats. Whole-cell patch-clamp recordings were performed on NTS neurons at a clamp voltage of -60 mV. 100 μM Dynasore dissolved in artificial cerebrospinal fluid was continuously administered via a perfusion system. Postsynaptic currents were recorded at different time points (pre-administration, during administration, post-washout).</p> <p><b>Results:</b> Dynasore rapidly increased the frequency of spontaneous excitatory postsynaptic currents (EPSCs). Concurrently, ST-evoked EPSCs progressively failed until completely blocked, with no reversal during the recovery period.[3]</p>
In vivo	<p><b>Methods:</b> An acute lung injury (ALI) model was established in C57BL/6J mice via intratracheal LPS (10 mg/kg) administration. Dynasore (10, 30, 50 mg/kg) was administered as a single intraperitoneal injection 3 hours prior to LPS challenge. Mice were euthanized 24 hours after LPS stimulation.</p> <p><b>Results:</b> Dynasore pretreatment (particularly at 30 and 50 mg/kg) significantly reduced lung tissue histopathology scores and decreased inflammatory mediators. [1]</p>

## A DRUG SCREENING EXPERT

Kinase Assay	RIP1 kinase assay: Phosphorylation of RIP1 requires its kinase activity. Expression constructs of FLAGtagged wild-type (WT) or a kinase-inactive point mutant of RIP1 (K45M) are transfected into 293T cells and RIP1 kinase assay is performed as described in the Methods in the presence of [ $\gamma$ - <sup>32</sup> P]ATP for 30 min at 30°C. Samples are subjected to SDS-PAGE and RIP1 band is visualized by autoradiography. Relative intensities of radioactive bands are quantified and are shown (ratio) in this and all other autoradiographs. In parallel to kinase reactions, a sample of beads is subjected to western blot analysis using anti-RIP1 antibody to ensure equal protein amounts in kinase reactions.
Cell Research	Mouse ventricular myocytes are isolated from male adult C6/Black mouse. Cardiomyocytes subjects to 2 hours of drug treatment followed by oxidative stress (30 $\mu$ M Water <sub>2</sub> for 35 min). For ATP supplement experiments, the cells are treated with 3 mM ATP for 30 min before exposure to Water <sub>2</sub> . Cardiomyocyte survival and viability are analyzed by trypan blue exclusion (TBE) assay[3].

### Solubility Information

Solubility	Ethanol: 1.6 mg/mL (4.96 mM),Sonication is recommended. DMSO: 120 mg/mL (372.31 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+40% PEG300+5% Tween 80+45% Saline: 4 mg/mL (12.41 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>

### Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.1026 mL	15.513 mL	31.026 mL
5 mM	0.6205 mL	3.1026 mL	6.2052 mL
10 mM	0.3103 mL	1.5513 mL	3.1026 mL
50 mM	0.0621 mL	0.3103 mL	0.6205 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

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