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Product Data Sheet

T0467

Cat. No.:HY-139308CAS No.:859518-94-8Molecular Formula: $C_{24}H_{26}F_3N_5$ Molecular Weight:441.49

Target: Mitochondrial Metabolism; PINK1/Parkin

Pathway: Metabolic Enzyme/Protease; Autophagy; Neuronal Signaling

Storage: Powder -20°C 3 years

In solvent

4°C 2 years
-80°C 6 months
-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 33.33 mg/mL (75.49 mM; Need ultrasonic)

| Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-----------|------------|------------|
| | 1 mM | 2.2651 mL | 11.3253 mL | 22.6506 mL |
| | 5 mM | 0.4530 mL | 2.2651 mL | 4.5301 mL |
| | 10 mM | 0.2265 mL | 1.1325 mL | 2.2651 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: ≥ 2.5 mg/mL (5.66 mM); Clear solution

BIOLOGICAL ACTIVITY

DescriptionT0467 activates parkin mitochondrial translocation in a PINK1-dependent manner in vitro. T0467 do not induce

mitochondrial accumulation of PINK1in dopaminergic neurons. T0467 is a potential compound for PINK1-Parkin signaling

activation, and can be used for parkinson's disease and related disorders research $^{[1]}$.

In Vitro T0467 (2.5-20 μM; 3 hours) stimulates the mitochondrial translocation of GFP-Parkinover 12 μM in HeLa/GFP-Parkin cells^[1].

When HeLa/GFP-Parkin cells are treated with 20 μ M T0467 for 3 h, GTP-Parkin is translocated to the mitochondria in

approximately 21% of cells^[1].

T0467 does not show obvious toxicity in Drosophila at concentrations <50 μ M. All cpds examined mitigated the PINK1 inactivation-mediated larval locomotion defects and mitochondrial morphological defects and reduced ATP production.

T0467 and KTP improved the mitochondrial Ca^{2+} response in Drosophila larval muscles [1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

• Research Square Preprint. 2023 Dec 27

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REFERENCES

[1]. Kahori Shiba-Fukushima, et al. A Cell-Based High-Throughput Screening Identified Two Compounds that Enhance PINK1-Parkin Signaling. iScience. 2020 May 22;23(5):101048.

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

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