MRT68921 hydrochloride

| Cat. No.: | HY-100006B | |
|--------------------|---|--|
| CAS No.: | 2070014-87-6 | |
| Molecular Formula: | C ₂₅ H ₃₅ ClN ₆ O | |
| Molecular Weight: | 471.04 | |
| Target: | ULK | |
| Pathway: | Autophagy | |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. | |

| Description MRT68921 hydrochloride is a potent inhibitor of ULK1 and ULK2, with IC ₅₀ values of 2.9 nM and 1.1 nM, respectively. In Vitro ULK1, a serine/threonine protein kinase, is essential for the initial stages of autophagy. MRT68921 inhibits ULK1 and ULK2 vitro and block autophagy in cells. MRT68921 is the most potent inhibitor of both ULK1 and ULK2, with greater than a 15-reduction in the IC ₅₀ for ULK1 (2.9 nm) and greater than a 30-fold reduction for ULK2 (1.1 nm). Autophagy-inhibiting capae of the compounds is specifically through ULK1. ULK1 inhibition results in accumulation of stalled early autophagosomal structures, indicating a role for ULK1 in the maturation of autophagosomes as well as initiation ^[1] . | | | | | |
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| In Vitro ULK1, a serine/threonine protein kinase, is essential for the initial stages of autophagy. MRT68921 inhibits ULK1 and ULK2 vitro and block autophagy in cells. MRT68921 is the most potent inhibitor of both ULK1 and ULK2, with greater than a 15-reduction in the IC ₅₀ for ULK1 (2.9 nm) and greater than a 30-fold reduction for ULK2 (1.1 nm). Autophagy-inhibiting capa of the compounds is specifically through ULK1. ULK1 inhibition results in accumulation of stalled early autophagosomal structures, indicating a role for ULK1 in the maturation of autophagosomes as well as initiation ^[1] . | | Description | MRT68921 hydrochloride is a potent inhibitor of ULK1 and ULK2, with IC ₅₀ values of 2.9 nM and 1.1 nM, respectively. | | |
| Mice has not independently commed the accuracy of these methods. They are for reference only. | | In Vitro | ULK1, a serine/threonine protein kinase, is essential for the initial stages of autophagy. MRT68921 inhibits ULK1 and ULK2 in vitro and block autophagy in cells. MRT68921 is the most potent inhibitor of both ULK1 and ULK2, with greater than a 15-fold reduction in the IC ₅₀ for ULK1 (2.9 nm) and greater than a 30-fold reduction for ULK2 (1.1 nm). Autophagy-inhibiting capacity of the compounds is specifically through ULK1. ULK1 inhibition results in accumulation of stalled early autophagosomal structures, indicating a role for ULK1 in the maturation of autophagosomes as well as initiation ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | |

CUSTOMER VALIDATION

- Nature. 2022 Oct;610(7931):366-372.
- Cancer Cell. 2021 May 10;39(5):678-693.e11.
- Nature Cancer. 2021 May;2(5):503-514.
- Mol Ther Oncolytics. 28 August 2021.
- J Cell Physiol. 2021 Dec;236(12):8110-8121.

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REFERENCES

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[1]. Petherick KJ, et al. Pharmacological inhibition of ULK1 kinase blocks mammalian target of rapamycin (mTOR)-dependent autophagy. J Biol Chem. 2015 May 1;290(18):11376-83.



Product Data Sheet

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

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