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**Product Data Sheet**

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Product Name: ATC 0065

Cat. No.: GC10994

**Chemical Properties**

Cas. No. 510732-84-0

Chemical Name N2-((1r,4r)-4-((4-bromo-2-(trifluoromethoxy)phenethyl)amino)cyclohexyl)-N4,N4-dimethylquinazoline-2,4-diamine dihydrochloride

SMILES BrC1=CC=C(C(OC(F)(F)F)=C1)CCN[C@@H](CC2)CC[C@H]2NC3=NC4=CC=CC=C4C(N(C)C)=N3.Cl.ClFormula C<sub>25</sub>H<sub>29</sub>BrF<sub>3</sub>N<sub>5</sub>O.2HCl M.Wt 625.35

Solubility Soluble in DMSO Storage Desiccate at RT

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Background**

ATC0065 is a selective and potent antagonist of the melanin-concentrating hormone receptor 1 (MCH1) with IC<sub>50</sub> of 15.7 nM. This component also displays affinity for 5-HT<sub>1A</sub> and 5-HT<sub>2B</sub> receptors with IC<sub>50</sub> of 62.9nM and 266 nM respectively.

Melanin-concentrating hormone receptor (MCH1), a member of the G protein-coupled receptor family 1, is an integral plasma membrane protein which binds melanin-concentrating hormone. The protein can inhibit cAMP accumulation and stimulate intracellular calcium flux, and is probably involved in the neuronal regulation of food consumption.

In vitro assay, ATC0065 bounds with high affinity to the MCH-R1 with IC<sub>50</sub> value of 16 nM and showed good metabolic stability in liver microsomes isolated from human and rat 1.

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

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In rodents, ATC0065 is potent and orally active MCHR1 antagonists with anxiolytic and antidepressant activity. ATC0065 treatment significantly reversed swim stress-induced anxiety in the stress-induced hyperthermia in mice and elevated plus-maze test in rats<sup>2</sup>. However, ATC0065 did not affect spontaneous locomotor activity or rotarod performance in rats<sup>2</sup>.

### References:

1. Kanuma K, Omodera K, Nishiguchi M, et al. Identification of 4-amino-2-cyclohexylaminoquinazolines as metabolically stable melanin-concentrating hormone receptor 1 antagonists. *Bioorganic & medicinal chemistry*. 2006;14(10):3307-3319.
2. Chaki S, Funakoshi T, Hirota-Okuno S, et al. Anxiolytic- and antidepressant-like profile of ATC0065 and ATC0175: nonpeptidic and orally active melanin-concentrating hormone receptor 1 antagonists. *The Journal of pharmacology and experimental therapeutics*. 2005;313(2):831-839.

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