

## Product Data Sheet

Product Name: iMQT\_020

Cat. No.: GC81546

### Chemical Properties

Cas. No. 2463893-46-9

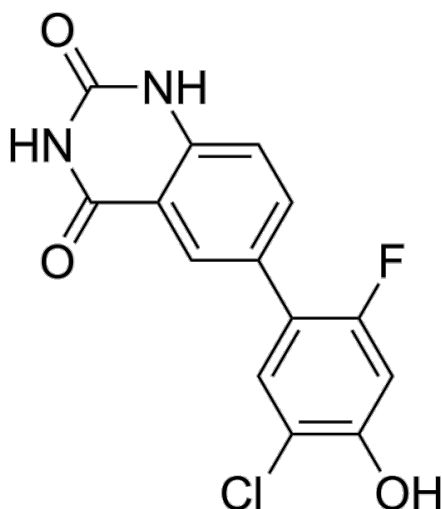
Formula  $C_{14}H_8ClFN_2O_3$  M.Wt 306.68

Solubility DMSO: 100 mg/mL (326.07 mM; Need ultrasonic) Storage Store at -20°C

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

iMQT\_020 is a selective allosteric SLC1A5\_var inhibitor. iMQT\_020 disrupts the trimeric assembly of SLC1A5\_var, causing metabolic crisis in cancer cells and selectively suppressing their growth. iMQT\_020 reduces glutamine anaplerosis and oxidative phosphorylation, resulting in a broad

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

Tel: (909) 407-4943 Fax: (626) 353-8530 E-mail: tech@glpbio.com

Address: 10292 Central Ave. #205, Montclair, CA, USA

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disruption of cancer metabolism. iMQT\_020 reduces GSH levels and increases cellular ROS and mitochondrial ROS. iMQT\_020 induces apoptosis and ferroptosis. iMQT\_020 can epigenetically upregulate PD-L1 expression. iMQT\_020 can be used for the study of pancreatic cancer, lung cancer, and colon cancer [1].

In Vivo, iMQT\_020 (75 mg/kg, i.p., once daily for 35 days) inhibits tumor growth in mice xenografted with MIA PaCa-2 cells by inducing apoptosis and ferroptosis [1]. iMQT\_020 (75 mg/kg, i.p., once daily for 35 days) inhibits tumor growth in a mouse orthotopic transplantation model of MIA PaCa-2 cells [1]. iMQT\_020 (75 mg/kg, i.p., once daily for 35 days) inhibits the growth of xenografted tumors of NCI-H1299 human lung cancer cells and COLO 205 human colon cancer cells in mice [1]. iMQT\_020 (25 mg/kg, i.p., once daily for 21 days), when used in combination with an anti-PD-L1 antibody (aPD-L1), synergistically inhibits the growth of allogeneic tumors formed from KPC, LLC, or MC-38 cells in mice [1].

In Vitro, iMQT\_020 shows the strongest inhibitory effect on mitochondrial glutamine uptake ( $IC_{50} = 6.156 \mu M$ ) at a concentration of  $10 \mu M$  [1]. iMQT\_020 (0-100  $\mu M$ ) directly binds to SLC1A5\_var WT protein ( $K_d = 4.473 \mu M$ ), but does not bind to FIL/AAA mutants [1]. iMQT\_020 (4  $\mu M$ ) significantly reduces the CD signal of the SLC1A5\_var WT protein, indicating a structural change [1]. iMQT\_020 (10  $\mu M$ ) reduces glutamine-derived TCA cycle metabolites (such as glutamate,  $\alpha KG$ , succinic acid, etc.) and downstream products (such as glutathione, proline) in MIA PaCa-2 cells [1]. iMQT\_020 (10  $\mu M$ , 24 h) reduces GSH levels in MIA PaCa-2 cells and increases cellular ROS and mitochondrial ROS [1]. iMQT\_020 (10  $\mu M$ , 24 h) reduces OCR and ECAR in SLC1A5\_var WT-overexpressing MIA PaCa-2 cells, but did not affect FIL/AAA mutants [1]. iMQT\_020 (10  $\mu M$ , 24 h) alters mitochondrial morphology in MIA PaCa-2 cells (reducing fragmentation) and decreases mitochondrial membrane potential (reduced TMRE staining) [1]. iMQT\_020 (48 h) selectively inhibits the viability of cancer cells ( $IC_{50}$  5-40  $\mu M$ ) without affecting normal cells [1]. iMQT\_020 (10  $\mu M$ , 24 h) upregulates PD-L1 mRNA and protein expression in human PDAC cell lines (such as SU.86.86, SW1990) and mouse carcinoma cell lines (KPC, LLC, MC-38) [1].

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[1]. Sung Y, et al. Targeting cancer glutamine dependency with a first-in-class inhibitor of the mitochondrial glutamine transporter SLC1A5\_var. Nat Commun. 2025 Nov 3;16(1):9690.

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