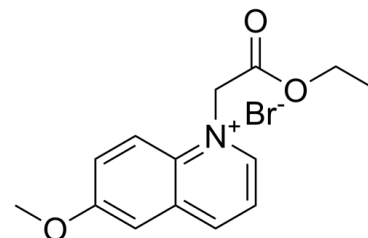


## Data Sheet

<b>Product Name:</b>	MQAE
<b>Cat. No.:</b>	CS-6018
<b>CAS No.:</b>	162558-52-3
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>16</sub> BrNO <sub>3</sub>
<b>Molecular Weight:</b>	326.19
<b>Target:</b>	Fluorescent Dye
<b>Pathway:</b>	Others
<b>Solubility:</b>	DMSO : ≥ 35 mg/mL; H <sub>2</sub> O : 100 mg/mL (ultrasonic)



### BIOLOGICAL ACTIVITY:

MQAE is a fluorescently-labeled deoxyglucose analog that is used primarily to directly monitor glucose uptake by living cells and tissues. It is also used as a topical contrast reagent for the detection of neoplasia. MQAE can be used in real-time confocal, high-resolution, or wide-field fluorescence microscopy as well as in flow cytometry. The probe can be excited by the Argon laser at 488 nm to give the environment-sensitive fluorescence. It has lower photostability than the rhodamine-based fluorescent probes. *In Vitro*: General Protocol

#### 1 Preparation of MQAE working solution

##### 1.1 Preparation of the stock solution

Dissolve 1 mg MQAE in 0.3066 mL DMSO to obtain 10 mM of MQAE.

Note: It is recommended to store the stock solution at -20°C -80°C away from light and avoid repetitive freeze-thaw cycles.

##### 1.2 Preparation of MQAE working solution

Dilute the stock solution in Krebs-hepes buffer (20 mM HEPES, 128 mM NaCl, 2.5 mM KCl, 2.7 mM CaCl<sub>2</sub>, 1 mM MgCl<sub>2</sub>, 16 mM glucose, pH 7.4) to obtain 5-10 mM of MQAE working solution.

Note: Please adjust the concentration of MQAE working solution according to the actual situation.

#### 2 Cell staining

2.1 For suspension cells: Centrifuge at 1000 g at 4°C for 3-5 minutes and then discard the supernatant. Wash twice with PBS, 5 minutes each time.

For adherent cells: Discard the cell culture medium, and add trypsin to dissociate cells to make a single-cell suspension. Centrifuge at 1000 g at 4°C for 3-5 minutes and then discard the supernatant. Wash twice with PBS, 5 minutes each time.

2.2 Add 1 mL of MQAE working solution, and then incubate at room temperature for 30 minutes.

2.3 Centrifuge at 400 g at 4°C for 3-4 minutes and then discard the supernatant.

2.4 Wash twice with PBS, 5 minutes each time.

2.5 Resuspend cells with serum-free cell culture medium or PBS.

#### 3 Storage

-20°C, 1 year. Protect from light.

#### 4 Precautions

4.1 It is recommended to store the stock solution at -20°C or -80°C away from light and avoid repetitive freeze-thaw cycles.

4.2 Please adjust the concentration of MQAE working solution according to the actual situation.

4.3 This product is for R&D use only, not for drug, household, or other uses.

4.4 For your safety and health, please wear a lab coat and disposable gloves to operate.

### References:

[1]. Andersson C, et al. Determination of chloride efflux by X-ray microanalysis versus MQAE-fluorescence. Microsc Res Tech. 2

[2]. Koncz C, et al. Use of MQAE for measurement of intracellular [Cl-] in cultured aortic smooth muscle cells. Am J Physiol. 1994 Dec;267(6 Pt 2):H2114-23.

[3]. Kovalchuk Y, et al. Two-photon chloride imaging using MQAE in vitro and in vivo. Cold Spring Harb Protoc. 2012 Jul 1;2012(7):778-85.

### CAIndexNames:

Quinolinium, 1-(2-ethoxy-2-oxoethyl)-6-methoxy-, bromide (1:1)

### SMILES:

COC1=CC2=CC=C[N+](CC(OCC)=O)=C2C=C1.[Br-]

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

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