**Proteins** 

# **Product** Data Sheet

## **BW-A78U**

Cat. No.: HY-100118 CAS No.: 101155-02-6 Molecular Formula: C<sub>13</sub>H<sub>12</sub>FN<sub>5</sub> Molecular Weight: 257.27

Target: Phosphodiesterase (PDE) Pathway: Metabolic Enzyme/Protease Storage: Powder -20°C 3 years

> 4°C 2 years In solvent -80°C 2 years

-20°C 1 year

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 150 mg/mL (583.05 mM; Need ultrasonic)

| Preparing<br>Stock Solutions | Solvent Mass<br>Concentration | 1 mg      | 5 mg       | 10 mg      |
|------------------------------|-------------------------------|-----------|------------|------------|
|                              | 1 mM                          | 3.8870 mL | 19.4348 mL | 38.8697 mL |
|                              | 5 mM                          | 0.7774 mL | 3.8870 mL  | 7.7739 mL  |
|                              | 10 mM                         | 0.3887 mL | 1.9435 mL  | 3.8870 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 (9.72 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- $\beta$ -CD in saline) Solubility: ≥ 2.5 mg/mL (9.72 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (9.72 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

| Description               | BW-A 78U is a PDE4 inhibitor with an IC $_{50}$ of 3 $\mu$ M.  |  |
|---------------------------|--|--|
| IC <sub>50</sub> & Target | PDE4<br>3 μM (IC <sub>50</sub> )   |  |
| In Vitro                  | BW-A 78U is a PDE4 inhibitor with an IC <sub>50</sub> of 3 $\mu$ M. BW-A 78U fails to significantly inhibit arachidonate release. BW-A 78U is ineffective to inhibit the lipopolysaccharide (LPS)-induced TNF- $\alpha$ release <sup>[1]</sup> . |  |

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

## **PROTOCOL**

Cell Assay [1]

Mononuclear cells are incubated for 30 min with BW-A 78U at the concentration of 10 nM to 10  $\mu$ M. The cells are then stimulated with lipopolysaccharide (10  $\mu$ g/mL) overnight at 37°C in an atmosphere of 5% CO<sub>2</sub> at 100% humidity. Cell-free supernatants are collected, centrifuged (2000 g), and stored frozen at -20°C before TNF- $\alpha$  determination. TNF- $\alpha$  concentrations in cell culture supernatants are determined by specific ELISA using a commercial kit. Sensitivity of the assay is 1 pg/mL. The absorbance at 450 nm is assessed with an ELISA reader<sup>[1]</sup>.

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#### **REFERENCES**

[1]. Boichot E, et al. Anti-inflammatory activities of a new series of selective phosphodiesterase 4 inhibitors derived from 9-benzyladenine. J Pharmacol Exp Ther. 2000 Feb;292(2):647-53.

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

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