Inhibitors

# Proteins

# **Product** Data Sheet

# **Brilliant Yellow**

Cat. No.: HY-W110888 CAS No.: 3051-11-4

Molecular Formula:  $C_{26}H_{18}N_4Na_2O_8S_2$ 

Molecular Weight: 624.55 Others Target: Pathway: Others

Storage: 4°C, sealed storage, away from moisture

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 10 mg/mL (16.01 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.6012 mL	8.0058 mL	16.0115 mL
	5 mM	0.3202 mL	1.6012 mL	3.2023 mL
	10 mM	0.1601 mL	0.8006 mL	1.6012 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: ≥ 1 mg/mL (1.60 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1 mg/mL (1.60 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Brilliant Yellow, a diazo-containing sulfonic aci, is also a potent VGLUT-specific inhibitor. Brilliant Yellow is membrane-impermeable. However, there are some Brilliant Yellow analogs with low cytotoxicity and cell penetration. Brilliant Yellow analogs work on glutamatergic transmission in hippocampal neurons <sup>[1]</sup> .
IC <sub>50</sub> & Target	$VGLUT^{[1]}$

### **REFERENCES**

[1]. Kehrl J, et al. Vesicular Glutamate Transporter Inhibitors: Structurally Modified Brilliant Yellow Analogs. Neurochem Res. 2017 Jun;42(6):1823-1832.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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