

## **Product** Data Sheet

# (E)-4-Hydroxytamoxifen-d<sub>5</sub>

Cat. No.: HY-16950BS Molecular Formula:  $C_{26}H_{24}D_5NO_2$ 

Molecular Weight: 392.54

Target: Estrogen Receptor/ERR; Isotope-Labeled Compounds

Pathway: Vitamin D Related/Nuclear Receptor; Others

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (254.75 mM; Need ultrasonic)

| Preparing<br>Stock Solutions | Solvent Mass<br>Concentration | 1 mg      | 5 mg       | 10 mg      |
|------------------------------|-------------------------------|-----------|------------|------------|
|                              | 1 mM                          | 2.5475 mL | 12.7376 mL | 25.4751 mL |
|                              | 5 mM                          | 0.5095 mL | 2.5475 mL  | 5.0950 mL  |
|                              | 10 mM                         | 0.2548 mL | 1.2738 mL  | 2.5475 mL  |

Please refer to the solubility information to select the appropriate solvent.

### **BIOLOGICAL ACTIVITY**

**Description** (E)-4-Hydroxytamoxifen-d5 ((E)-Afimoxifene-d5) is the deuterium labeled (E)-4-Hydroxytamoxifen. (E)-4-Hydroxytamoxifen ((E)-Afimoxifene), the less active isomer of (Z)-4-hydroxytamoxifen, is an estrogen receptor modulator.

In Vitro Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to

affect the pharmacokinetic and metabolic profiles of drugs[1].

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

### **REFERENCES**

[1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. Ann Pharmacother. 2019;53(2):211-216.

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

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Page 2 of 2 www.MedChemExpress.com