

# DMAT

**Catalog No: tcsc1289** 

Available Sizes

Size: 10mg

Size: 50mg

🗐 Sp

**Specifications** 

**CAS No:** 749234-11-5

## Formula:

 $C_9H_7Br_4N_3$ 

Pathway: Stem Cell/Wnt;Cell Cycle/DNA Damage

## **Target:**

Casein Kinase; Casein Kinase

#### Purity / Grade:

>98%

## Solubility: DMSO : 50 mg/mL (104.87 mM; Need ultrasonic)

#### **Alternative Names:**

Casein kinase II Inhibitor;CK2 Inhibitor

## **Observed Molecular Weight:**

476.79

## **Product Description**

DMAT is a potent and specific **CK2** inhibitor with an  $IC_{50}$  value of 130 nM.

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IC50 & Target: IC50: 130 nM (CK2)<sup>[3]</sup>

*In Vitro:* DMAT (1  $\mu$ M-2.5  $\mu$ M) DMAT is more efficient in killing antiestrogen resistant cells than parental antiestrogen sensitive MCF-7 cells. DMAT-induced cell death of antiestrogen resistant cells is mediated by caspases. DMAT inhibits CK2 activity but the inhibition is similar in the three cell lines, MCF-7, TAMR-1 and 182R-6<sup>[1]</sup>. DMAT has effects on H295R cell proliferation at concentrations of 10<sup>-4</sup> and 10<sup>-5</sup>mol/Las compared with the control. DMAT (100  $\mu$ M) significantly increases apoptosis of H295R cells. DMAT (1 nM-1  $\mu$ M) significantly decreases aldosterone release into supernatants of 72-h H295R cell cultures as compared with the control<sup>[2]</sup>. DMAT also inhibits PIM1 by a mechanism which is competitive with respect to ATP, and it is a powerful inhibitor of kinases other than CK2<sup>[3]</sup>.

*In Vivo:* DMAT application in vivo reduces tumor growth in a xenotransplant model by interference with tumor cell proliferation. Biochemical parameters and histology following DMAT administration revealed no alterations in liver tissue<sup>[4]</sup>.



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