

TBB

Catalog No:	tcsc1079
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Available Sizes

Size: 10mg

Size: 50mg

Specifications

CAS No: 17374-26-4

Formula:

 $C_6HBr_4N_3$

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

Target:

Casein Kinase;Casein Kinase

Purity / Grade:

>98%

Solubility: DMSO : ≥ 430 mg/mL (989.17 mM)

Alternative Names:

NSC 231634; Casein Kinase II Inhibitor I

Observed Molecular Weight:

434.71

Product Description

TBB is a cell-permeable and ATP-competitive **CK2** inhibitor with an IC_{50} of 0.15 μ M for rat liver CK2.

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IC50 & Target: IC50: 0.15 μM (CK2), 14 μM (CDK2)^[5]

In Vitro: Investigation of the inhibitory power of TBB with a panel of 33 protein kinases shows highest potency for CK2 (casein kinase 2) (human CK2: IC_{50} =1.6 µM at 100 µM ATP). TBB also inhibits three other kinases with less potency: CDK2 (IC_{50} =15.6 µM), phosphorylase kinase (IC_{50} =8.7 µM) and glycogen synthase kinase 3β (GSK3β) (IC_{50} =11.2 µM). All other kinases tested have IC50 values 50-fold greater than that for CK2^[1]. The viability of the androgen insensitive PC-3 cells may be diminished by TBB (60 µM TBB) acting either alone or combined with anticancer agents CPT or TRAIL when a proper time schedule of the administration is applied. The time schedule-dependent activity of TBB does not come from its effect on apoptosis in PC-3 cells^[2]. TBB is an ATP/GTP competitive inhibitor of protein kinase casein kinase-2 (CK2), has been examined against a panel of 33 protein kinases, either Ser/Thr- or Tyr-specific. In the presence of 10 µM TBB (and 100 µM ATP) only CK2 is drastically inhibited (>85%) whereas three kinases (phosphorylase kinase, glycogen synthase kinase 3L and cyclin-dependent kinase 2/cyclin A) underwent moderate inhibition, with IC₅₀ values one-two orders of magnitude higher than CK2 (IC_{50} =0.9 µM). TBB also inhibits endogenous CK2 in cultured Jurkat cells^[3].

In Vivo: The extent of retinal neovascularization in a mouse OIR model is reduced by approximately 60% after treatment with TBB (6 days at 60 mg/kg per day)^[4].



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