

# AGN 193109

Catalog No: tcsc0035405



## Available Sizes

Size: 5mg

Size: 10mg

Size: 25mg



## Specifications

**CAS No:**

171746-21-7

**Formula:**

$C_{28}H_{24}O_2$

**Pathway:**

Metabolic Enzyme/Protease

**Target:**

RAR/RXR

**Purity / Grade:**

>98%

**Solubility:**

DMSO : 2 mg/mL (5.10 mM; Need warming)

**Observed Molecular Weight:**

392.49

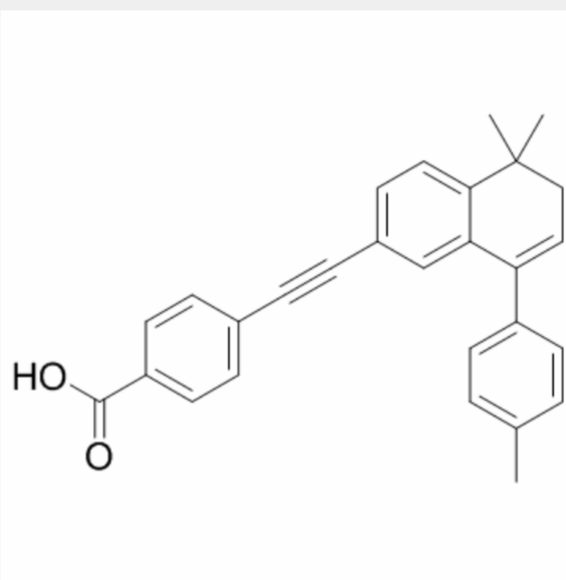
## Product Description

AGN 193109 is a retinoid analog, and acts as a specific and highly effective antagonist of **retinoic acid receptors (RARs)**, with  $K_d$ s of 2 nM, 2 nM, and 3 nM for **RAR $\alpha$** , **RAR $\beta$** , and **RAR $\gamma$** , respectively.

IC50 & Target: Kd: 2 nM (RAR $\alpha$ ), 2 nM (RAR $\beta$ ), 3 nM (RAR $\gamma$ )<sup>[1]</sup>

**In Vitro:** AGN 193109 is a highly effective antagonist of retinoic acid receptors, with K<sub>d</sub>s of 2 nM, 2 nM, and 3 nM for RAR $\alpha$ , RAR $\beta$ , and RAR $\gamma$ , respectively. AGN 193109 is completely RAR specific, because it does not bind to or transactivate through any of the RXRs <sup>[1]</sup>. AGN 193109 (100 nM) inhibits the TTNPB (a retinoic acid receptor agonist)-dependent morphological change in ECE16-1 cells. AGN193109 half-reverses retinoid-dependent growth suppression at 10 nM, and completely shows this effect at 100 nM in ECE16-1 cells. AGN193109 (100 nM) also eliminates TTNPB-induced decrease in levels of K5, K6, K14, K16, and K17 and increase in levels of K7, K8, and K19<sup>[2]</sup>.

**In Vivo:** AGN 193109 (1.15  $\mu$ mol/kg) does not cause overt toxicity and has no effect on spleen weight on the mice, but it suppresses TTNPB-induced increase in spleen weight of the mice. AGN 193109 also significantly reduces the cutaneous toxicity induced by ATRA. AGN 193109 (0.30 or 1.20  $\mu$ mol/kg) by topical treatment significantly reduces both weight loss and cutaneous toxicity caused by oral TTNPB cotreatment<sup>[3]</sup>.



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!