TTNPB

Small Molecules

Retinoid pathway activator; Activates retinoic acid receptor (RAR)

Catalog # 72892 10 mg



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Product Description

TTNPB is an analog of retinoic acid that potently and selectively activates retinoic acid receptors (RAR; $EC_{50} = 21$, 4, and 2.4 nM for RAR α , RAR β , and RAR γ , respectively; Beard et al.; Wong et al.). It does not act on retinoid X receptors and weakly agonizes farnesoid X receptor ($EC_{50} > 1 \mu$ M; Maloney et al.; Wong et al.). TTNPB is used to study RAR action in diverse processes, including epidermal cell proliferation, embryogenesis, and stem cell differentiation (Araoka et al.; Hou et al.; Minucci et al.; Thacher et al.).

Molecular Name: TTNPB

Alternative Names: AGN 191183; Arotinoid Acid; Ro 13-7410

CAS Number: 71441-28-6 Chemical Formula: $C_{24}H_{28}O_2$ Molecular Weight: 348.5 g/mol Purity: \geq 98%

Chemical Name: 4-[(1E)-2-(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)-1-propen-1-yl]-benzoic acid

Structure:

Properties

Physical Appearance: A crystalline solid

Storage: Product stable at -20°C as supplied. Protect from prolonged exposure to light. For product expiry date, please

contact techsupport@stemcell.com.

Solubility: · Absolute ethanol ≤ 280 µM

 \cdot DMSO ≤ 5.7 mM

For example, to prepare a 1 mM stock solution in DMSO, resuspend 1 mg in 2.86 mL of fresh DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported, however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

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Published Applications

REPROGRAMMING

- · Enables chemical reprogramming (without genetic factors) of mouse embryonic fibroblasts to induced pluripotent stem (iPS) cells, in combination with CHIR99021, Tranylcypromine, Valproic Acid, 3-Deazaneplanocin A, and RepSox (Hou et al.). DIFFERENTIATION
- · In combination with CHIR99021 or Activin A, induces intermediate mesoderm formation from human or mouse pluripotent stem cells, respectively (Araoka et al.; Oeda et al.).
- · Promotes neuronal differentiation in cultured chick caudal neural plate explants (Diez del Corral et al.). CANCER RESEARCH
- · Induces the in vitro growth and differentiation to granulocytes of myeloid progenitors isolated from myelodysplastic syndrome (MDS) patients (Fabian et al.).

References

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Fabian I et al. (1987) In-vitro growth and differentiation of marrow cells from myelodysplastic patients in the presence of a retinoidal benzoic acid derivative. Leuk Res 11(7): 635–40.

Hou P et al. (2013) Pluripotent stem cells induced from mouse somatic cells by small-molecule compounds. Science 341(6146): 651–4. Maloney PR et al. (2000) Identification of a chemical tool for the orphan nuclear receptor FXR. J Med Chem 43(16): 2971–4.

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Thacher SM et al. (1997) Receptor specificity of retinoid-induced epidermal hyperplasia: effect of RXR-selective agonists and correlation with topical irritation. J Pharmacol Exp Ther 282(2): 528–34.

Wong MF et al. (1997) Synthesis and receptor binding affinity of conformationally restricted retinoic acid analogues. Bioorg Med Chem Lett 7(17): 2313–2318.

Related Small Molecules

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