

Small Molecules

AM580

Retinoid pathway activator; RAR alpha activator

Catalog # 72962
72964

1 mg
10 mg



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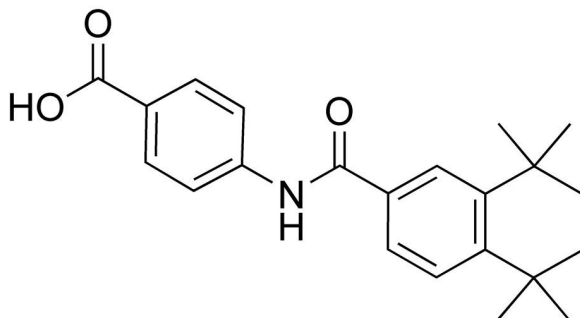
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Product Description

AM580 is a retinoic acid receptor (RAR) agonist that is selective for RAR α (EC_{50} = 0.36 nM) compared to RAR β (EC_{50} = 24.6 nM) and RAR γ (EC_{50} = 27.9 nM; Bernard et al.) It is a derivative of retinoic acid (RA), however it demonstrates greater specific binding to RAR α compared to RA, which exhibits little selectivity across RAR α , β , or γ (Gianní et al.; Bernard et al.; Kim et al; Rochette-Egly & Germain).

Molecular Name: AM580
Alternative Names: CD336, NSC 608001, Ro 40-6055
CAS Number: 102121-60-8
Chemical Formula: $C_{22}H_{25}NO_3$
Molecular Weight: 351.4 g/mol
Purity: $\geq 98\%$
Chemical Name: 4-[[[(5,6,7,8-tetrahydro-5,5,8,8-tetramethyl-2-naphthalenyl)carbonyl]amino]-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:

- DMSO ≤ 55 mM
- Absolute ethanol ≤ 25 mM

For example, to prepare a 10 mM stock solution in DMSO, resuspend 1 mg in 285 μ L of 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.

Published Applications

REPROGRAMMING

- Promotes reprogramming of somatic cells to induced pluripotent stem cells (Wang et al.).

DIFFERENTIATION

- Induces differentiation of human induced pluripotent stem cells into intermediate mesoderm, in combination with the GSK3 β inhibitor CHIR99021 (Araoka et al.).

CANCER

- Inhibits tumor cell proliferation and survival signaling pathways, and induces apoptosis, leading to inhibition of mouse mammary tumor virus (MMTV)-neu- and MMTV-wnt1-induced mammary gland hyperplasia (Lu et al.).
- Inhibits tumor growth in MMTV-Myc mice (Bosch et al.).
- Inhibits endometrial cancer cell proliferation (Cheng et al.).
- Induces differentiation in acute promyelocytic leukemia cells (Gianní et al.).

References

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