

Small Molecules

(±)-Nutlin-3

p53 pathway activator; Inhibits MDM2

Catalog # 73752
73754

1 mg
10 mg



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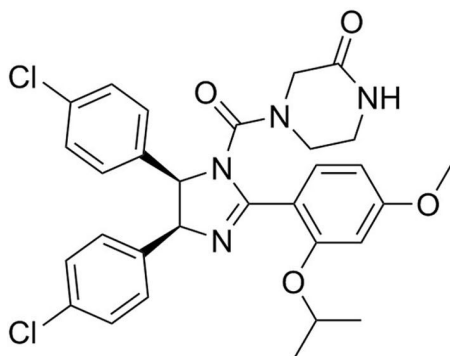
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Product Description

(±)-Nutlin-3 is a small-imidazoline-based mouse double minute 2 (MDM2) protein antagonist which disrupts MDM2-p53 interaction ($IC_{50} = 0.09 \mu M$; Vassilev et al.). MDM2 binds the p53 tumor suppressor protein with high affinity and negatively modulates its transcriptional activity and stability. By disrupting this interaction, (±)-Nutlin-3 therefore promotes the expression of p53-regulated genes and exhibits potent antiproliferative activity in cells with functional p53, but not in cells with mutated p53 (Vassilev et al., Gu et al.).

Molecular Name:	(±)-Nutlin-3
Alternative Names:	Not applicable
CAS Number:	548472-68-0
Chemical Formula:	$C_{30}H_{30}Cl_2N_4O_4$
Molecular Weight:	581.5 g/mol
Purity:	> 98%
Chemical Name:	4-[4,5-bis(4-chlorophenyl)-2-(4-methoxy-2-propan-2-yloxyphenyl)-4,5-dihydroimidazole-1-carbonyl]piperazin-2-one

Structure:



Properties

Physical Appearance:	A crystalline solid
Storage:	Product stable at $-20^{\circ}C$ as supplied. Protect product from prolonged exposure to light. For long-term storage store with a desiccant. For product expiry date, please contact techsupport@stemcell.com .
Solubility:	· DMSO ≤ 40 mM · Absolute ethanol ≤ 40 mM For example, to prepare a 10 mM stock solution in DMSO, resuspend 1 mg in 172 μL 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 $-20^{\circ}C$. 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

CANCER RESEARCH

- Inhibits the proliferation of exponentially growing human skin fibroblasts ($IC_{50} = 2.2 \mu M$) and mouse embryonic fibroblasts ($IC_{50} = 1.3 \mu M$), and suppresses the growth of established tumor xenografts in mice (Vassilev et al.).
- Leads to G1 cell-cycle arrest in HCT116 colon carcinoma cell line expressing wild-type p53 and p21 (Benson et al.).
- Induces p53-mediated apoptosis in solid tumor and pediatric acute lymphoblastic leukemia cell lines (Gu et al.; Vaseva et al.)

References

- Benson EK et al. (2014) p53-dependent gene repression through p21 is mediated by recruitment of E2F4 repression complexes. *Oncogene* 33(30): 3959–69.
- Gu L et al. (2008) MDM2 antagonist nutlin-3 is a potent inducer of apoptosis in pediatric acute lymphoblastic leukemia cells with wild-type p53 and overexpression of MDM2. *Leukemia* 22(4): 730–9.
- Vaseva A V et al. (2011) Blockade of Hsp90 by 17AAG antagonizes MDMX and synergizes with Nutlin to induce p53-mediated apoptosis in solid tumors. *Cell Death Dis* 2: e156.
- Vassilev LT et al. (2004) In vivo activation of the p53 pathway by small-molecule antagonists of MDM2. *Science* 303(5659): 844–8.

Related Small Molecules

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