

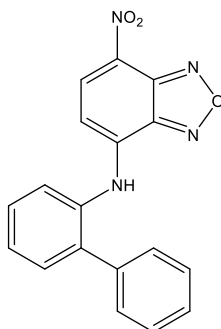
Catalog # 10-5627

10074-G5

CAS# 413611-93-5

7-Nitro-N-(2-phenylphenyl)-2,1,3-benzoxadiazol-4-amine

Lot # S106255



10074-G5 inhibits c-Myc by disrupting c-Myc/Max heterodimer formation and inhibiting its transcriptional activity.¹ 10074-G5 binds to the c-Myc bHLHZip domain with $K_d=2.8 \mu\text{M}$.¹ It was cytotoxic in C-Myc-overexpressing cell lines ($\text{IC}_{50}\text{s} = 13$ to $15 \mu\text{M}$) but was not active *in vivo* due to rapid metabolism.² 10074-G5 is a useful tool to probe for involvement of c-Myc involvement in cellular processes.^{3,4} Binds to and sequesters the intrinsically disordered amyloid- β (A β) in its monomeric soluble state and rescues a *C. elegans* model of A β -associated toxicity.⁵

- 1) Follis *et al.* (2008), *Structural rationale for the coupled binding and unfolding of the c-Myc oncoprotein by small molecules*; Chem..Biol. **15** 1149
- 2) Clausen *et al.* (2010), *In vitro cytotoxicity and in vivo efficacy, pharmacokinetics, and metabolism of 10074-G5, a novel small-molecule inhibitor of c-Myc/Max dimerization*; J. Pharmacol. Exp. Ther. **335** 715
- 3) Velpula *et al.* (2012), *Transcriptional repression of Mad-Max complex by human umbilical cord blood stem cells downregulates extracellular signal-regulated kinase in glioblastoma*; Stem Cells Dev. **21** 1779
- 4) Wu *et al.* (2018), *Oxidative stress enhances tumorigenicity and stem-like features via the activation of the Wnt/ β -catenin/MYC/Sox2 axis in ALK-positive anaplastic large-cell lymphoma*; BMC Cancer **18** 361
- 5) Heller *et al.* (2020), *Small-molecule sequestration of amyloid- β as a drug discovery strategy for Alzheimer's disease*; Sci. Adv. **6** eabb5924

PHYSICAL DATA

Molecular Weight:	332.32
Molecular Formula:	C ₁₈ H ₁₂ N ₄ O ₃
Purity:	>98% by TLC
	NMR: (Conforms)
Solubility:	DMSO (40 mg/ml)
Physical Description:	Orange solid
Storage and Stability:	Store as supplied at -20°C for up to 2 years from the date of purchase. Solutions in DMSO may be stored at -20°C for up to 3 months.

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