

aTAG 2139 Biochemical | aTAG2139 biochemical

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CATALOG #
AAA227029

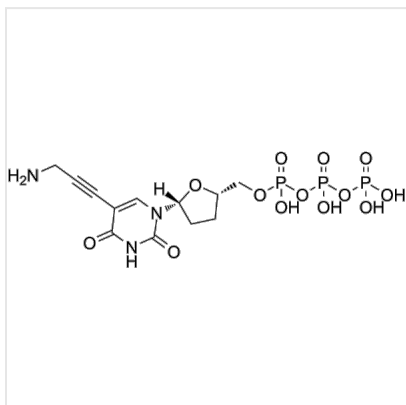
PRODUCT SYNONYMS

aTAG 2139; aTAG2139 biochemical

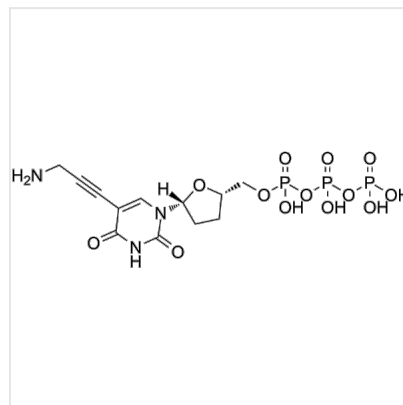
PRODUCT SPECIFICATIONS

Purity/Purification	98%
Solubility	DMSO: Soluble <1mg/ml refers to the product slightly soluble or insoluble
Preparation and Storage	Powder: -20 degree C for 3 years In solvent: -80 degree C for 2 years

IMAGES



Structure
5-Propargylamino-ddUTP, CAS 114748-59-3



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ADDITIONAL INFORMATION

Related Product Information for aTAG2139 biochemical	Degrader of MTH1 fusion proteins for use within the aTAG system. Comprises a ligand selective for MTH1, a linker and the cereblon-binding ligand Thalidomide. Induces highly potent and selective degradation of fusion proteins after a 4 h incubation (DC50 = 0.27 nM; Dmax = 92.1%). Cell-permeable. Suitable for in vitro and in vivo applications. Mouse DMPK properties are provided in the supplementary file (see below). (MTH1 can be expressed as a fusion with a target protein of interest using genome engineering techniques via CRISPR-mediated locus-specific knock-in - see protocol for more information.)
References	1. Duthie RS, et, al. Novel cyanine dye-labeled dideoxynucleoside triphosphates for DNA sequencing. Bioconjug Chem. Jul-Aug 2002;13(4):699-706.

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Small volumes of product may become entrapped in the seal of the vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the cap.