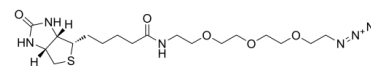


## Data Sheet

<b>Product Name:</b>	Biotin-PEG3-azide
<b>Cat. No.:</b>	CS-0105250
<b>CAS No.:</b>	875770-34-6
<b>Molecular Formula:</b>	C <sub>18</sub> H <sub>32</sub> N <sub>6</sub> O <sub>5</sub> S
<b>Molecular Weight:</b>	444.55
<b>Target:</b>	PROTAC Linkers
<b>Pathway:</b>	PROTAC
<b>Solubility:</b>	10 mM in DMSO



### BIOLOGICAL ACTIVITY:

Biotin-PEG3-azide is a PEG-based **PROTAC linker** can be used in the synthesis of PROTACs. Biotin-PEG3-azide is a click chemistry reagent, it contains an Azide group and can undergo copper-catalyzed azide-alkyne cycloaddition reaction (CuAAC) with molecules containing Alkyne groups. Strain-promoted alkyne-azide cycloaddition (SPAAC) can also occur with molecules containing DBCO or BCN groups. *In Vitro*: Biotin-C2-PEG3-azide is a biotin with the azide tag, it can be conjugated to antiviral inhibitors, for example, RYL-634, which shows excellent broad-spectrum inhibition activity against various pathogenic viruses, including hepatitis C virus, dengue virus, Zika virus, chikungunya virus, enterovirus<sup>[1]</sup>.

PROTACs contain two different ligands connected by a linker; one is a ligand for an E3 ubiquitin ligase and the other is for the target protein.

PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins.

### References:

[1]. Yang Y, et al. Discovery, Optimization, and Target Identification of Novel Potent Broad-Spectrum Antiviral Inhibitors. *J Med Chem.* 2019 Apr 25;62(8):4056-4073.

### CAIndexNames:

1H-Thieno[3,4-d]imidazole-4-pentanamide, N-[2-[2-[2-(2-azidoethoxy)ethoxy]ethoxy]ethyl]hexahydro-2-oxo-, (3aS,4S,6aR)-

### SMILES:

O=C(NCCOCCOCCOCCN=[N+]=[N-])CCCC[C@@H]1SC[C@]([C@]1([H])N2)([H])NC2=O

**Caution: Product has not been fully validated for medical applications. For research use only.**

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