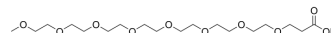


m-PEG7-CH₂CH₂COOH

| | | | |
|---------------------------|---|-------|----------|
| Cat. No.: | HY-130151 | | |
| CAS No.: | 1093647-41-6 | | |
| Molecular Formula: | C ₁₈ H ₃₆ O ₁₀ | | |
| Molecular Weight: | 412.47 | | |
| Target: | ADC Linker; PROTAC Linkers | | |
| Pathway: | Antibody-drug Conjugate/ADC Related; PROTAC | | |
| Storage: | Pure form | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |



BIOLOGICAL ACTIVITY

| | | |
|-------------------------------------|---|------|
| Description | m-PEG7-CH ₂ CH ₂ COOH is a non-cleavable ADC linker used in the synthesis of antibody-drug conjugates (ADCs). m-PEG7-CH ₂ CH ₂ COOH is also a PEG-based PROTAC linker that can be used in the synthesis of PROTACs. | |
| IC₅₀ & Target | Non-cleavable Linker | PEGs |
| In Vitro | ADCs are comprised of an antibody to which is attached an ADC cytotoxin through an ADC linker. 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. MCE has not independently confirmed the accuracy of these methods. They are for reference only. | |

REFERENCES

[1]. Tinya Abrams, et al. Antibody drug conjugates. WO2016203432A1.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA