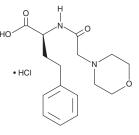
PRODUCT INFORMATION



Carfilzomib Impurity 13 (hydrochloride)

Item No. 22385

CAS Registry No.: Formal Name:	2319881-95-1 αS-[[2-(4-morpholinyl)acetyl]amino]- benzenebutanoic acid, monohydrochloride	
MF:	$C_{16}H_{22}N_2O_4 \bullet HCI$	
FW:	342.8	• HCI
Purity:	≥95%	
Supplied as:	A solid	
Storage:	-20°C	
Stability:	≥4 years	
Special Conditions: Store in hygroscopic conditions		
Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.		



Laboratory Procedures

Carfilzomib impurity 13 (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the carfilzomib impurity 13 (hydrochloride) in the solvent of choice, which should be purged with an inert gas. Carfilzomib impurity 13 (hydrochloride) is soluble in the organic solvent DMSO.

Description

Carfilzomib impurity 13 (hydrochloride) is a potential impurity found in commercial carfilzomib preparations. Carfilzomib (Item No. 17554) is a second-generation, irreversible, peptide epoxyketone class proteasome inhibitor that targets the chymotrypsin-like ß5 subunit of the constitutive 20S proteasome $(IC_{50} = 5.2 \text{ nM})$ and the β 5i subunit of the immunoproteasome 20Si (LMP7; IC₅₀ = 14 nM) with minimal cross reactivity to other proteases.^{1,2} It can induce cell cycle arrest and apoptosis in human cancer cell lines including multiple myeloma, lymphoma, and various solid tumors ($IC_{50}s = 2.4-20$ nM).³

References

- 1. Dou, Q.P., and Zonder, J.A. Overview of proteasome inhibitor-based anti-cancer therapies: Perspective on bortezomib and second generation proteasome inhibitors versus future generation inhibitors of ubiquitin-proteasome system. Curr. Cancer Drug Targets 14(6), 517-536 (2014).
- 2. Zhou, H.J., Aujay, M.A., Bennett, M.K., et al. Design and synthesis of an orally bioavailable and selective peptide epoxyketone proteasome inhibitor (PR-047). J. Med. Chem. 52(9), 3028-3038 (2009).
- Demo, S.D., Kirk, C.J., Aujay, M.A., et al. Antitumor activity of PR-171, a novel irreversible inhibitor of the 3. proteasome. Cancer Res. 67(13), 6383-6391 (2007).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFFTY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

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