

Fiaure 1

HA-Ahx-Ahx-Ub-VAE (VAE= vinyl azido ester, human sequence, synthetic)	Figure 1
UbiQ code : UbiQ-216	
Batch # : B01045014-001	
Amount : 50 ug, lyophilized powder	
Purity : ≥95% by RP-HPLC	
Mol. Weight : 9.98 kDa	
Storage : upon arrival, powder at –20°C; solution at –80°C. Avoid mult	iple freeze/thaw cycles.

## Productsheet

Background. HA-Ahx-Ahx-Ub-VAE (UbiQ-216, Figure 1) is an activity-based probe for deubiquitinating enzymes (DUBs). It is labelled on the N-terminus with the HA peptide sequence (YPYDVPDYA) derived from the influenza hemagglutinin protein and allows for the sensitive identification or purification of DUBs by anti-HA antibodies and/or anti-HA-agarose. The HA tag is separated from the N-terminus by two 6-aminohexanoic acid (Ahx) linkers for efficient recognition of the tag. The azide group in the vinyl azido ester warhead allows for further (post-labelling) modification by using click chemistry.

## sequence YPYDVPDYA-Ahx-Ahx-

MQIFVKTLTGKTITLEVEPSDTIENVKAKIQDKEGIPPDQQRLIFAGKQLEDGRTLSDYNIQKESTLHLVLRLRG-VAE

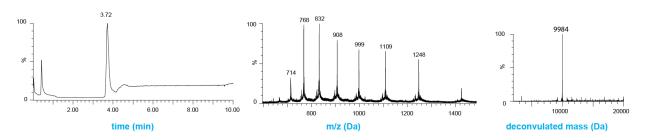


Figure 2 - LC-MS analysis. XBridge BEH300 C18 5µm 4.6x100 mm column; flow rate = 0.8 mL/min, runtime = 10 min, column T= 40°C. Mobile phase A = 1% CH<sub>3</sub>CN and 0.1% formic acid in water; B= 1% water and 0.1% formic acid in CH<sub>3</sub>CN. Gradient: 30-60% B over 6.5 min.

## Important: sample preparation

- dissolve the powder in as little DMSO as possible (e.g., 20 mg/mL)
- add this DMSO stock slowly to milliQ (please note the order of addition); mix by vortex
- next buffer as desired. For example:
  - 50 ug probe in 2.5 uL DMSO (20 mg/mL, 2 mM)
  - example 1: add to 47 uL water followed by addition of 0.5 uL 5M NaOAc pH 4.5 to prepare a 1 mg/mL stock in 50 mM 0 NaOAc pH 4.5 (100 uM); this stock is useful when working with low concentrations of probe
  - example 2: add to 45 uL water followed by addition of 2.5 uL 1M HEPES or Tris to prepare a 1 mg/mL stock in 50 mM 0 HEPES/Tris (100 uM); this stock is useful when working with high concentrations of probe

Literature. (1) (a) de Jong et al. ChemBioChem 2012, 13, 2251. (b) Borodovsky et al. EMBO J 2001, 20, 5187. (c) Borodovsky et al. Chem Biol 2002, 9, 1149. (2) El Oualid et al. Angew Chem Int Ed 2010, 49, 10149. (3) Hewings et al. Nat Commun 2018, 9, article number 1162.

Science Park 408 1098 XH Amsterdam The Netherlands

t +31 20 303 1970 e info@ubiqbio.com i www.ubiqbio.com

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