

## MATERIAL DATA SHEET

## Recombinant Human His6 USP19

Cat. # E-576

Ubiquitin specific processing protease 19 (USP19) is a 146 kDa member of the C19 peptidase family that contains two alpha crystallin domains and one MYND-type zinc finger. USP19 plays a role in the rescue of ER-associated protein degradation (ERAD) substrates such as cystic fibrosis transmembrane conductance regulator (CFTR)  $\Delta$ F508 and T-cell receptor-alpha (TCR $\alpha$ ) from proteasomal degradation. USP19 also interacts with the ubiquitin ligases cellular IAP 1 (cIAP1) and cIAP2. Knockdown of USP19 decreases levels of both cIAPs, whereas overexpression of USP19 results in a marked increase in cIAP levels. Although it effectively removes ubiquitin from cIAPs *in vitro*, USP19 may stabilize cIAP proteins *in vivo* mainly through deubiquitinase-independent mechanisms. Intriguingly, partial rescue of TCR $\alpha$  (but not CFTR  $\Delta$ F508) in the ERAD pathway is also observed using a catalytically dead USP19, suggesting that USP19 exerts a non-catalytic function for select substrates in multiple biological pathways. This recombinant protein contains amino acids a C-terminal 6-His tag.

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Product 1	Intorm	ation

**Quantity:** 50 μg

**MW:** 143 kDa

**Source:** Spodoptera frugiperda, Sf 21 (baculovirus)-derived

Met1 - Arg1290, Contains a C-terminal 6-His tag

Accession # O94966

Stock: X mg/ml (X μM) in 50 mM HEPES pH 7.5, 100 mM NaCl, 2 mM TCEP

**Purity:** >95%, by SDS-PAGE under reducing conditions and visualized by Colloidal

Coomassie® Blue stain.

**Use & Storage** 

**Use:** Recombinant Human His6-USP19 is a ubiquitin-specific deconjugating enzyme.

Reaction conditions will need to be optimized for each specific application. We

recommend an initial USP19 concentration of 10-100 nM.

Storage: Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

• 6 months from date of receipt, -70 °C as supplied.

• 3 months, -70 °C under sterile conditions after opening.





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## Literature

## **References:**

- 1. Hassink G.C., et al. (2009) EMBO Rep. 10: 755
- 2. Iphöfer, A. et al. (2012) ChemBioChem 13: 1416
- 3. Mei, Y. et al. (2011) J. Biol. Chem. 286: 35380

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