

**MATERIAL DATA SHEET****His<sub>6</sub>-NEDP1/SENPs8, human recombinant****Cat. # E-800**

The covalent modification of proteins by NEDD8 is reversible and is mediated by NEDP1. NEDP1 cleaves residues from the C-terminus of NEDD8 precursor proteins to generate the mature and active form which contains the conserved C-terminal di-glycine. This enzyme also reverse the process of NEDDylation by removing poly-NEDD8 conjugates from substrate proteins. NEDP1 has been shown to deconjugate NEDD8 from CUL2 *in vitro* and from CUL4A *in vivo*, and does not show activity against ubiquitin or SUMO proteins. Accession # NP\_660205.

**Product Information**

<b>Quantity:</b>	50 µg
<b>Stock:</b>	X mg/ml (X mM) in 50 mM Hepes pH 8.0, 100 mM NaCl, 10% glycerol, 5 mM DTT. Concentration will vary with specific Lot #.
<b>MW:</b>	26.5 kDa
<b>Purity:</b>	> 95% by SDS-PAGE

**Use & Storage**

<b>Use:</b>	Typical enzyme concentration to support hydrolysis of substrates <i>in vitro</i> is 50-500 nM depending on conditions. Pre-incubation (15 min) with 10 mM DTT is recommended to achieve maximum activity.
<b>Storage:</b>	Store at -80°C. Avoid multiple freeze/thaw cycles.

**Literature**

<b>References:</b>	Gan-Erdene T., <i>et al.</i> (2003) <u>J.Biol.Chem.</u> <b>278</b> :28882-28900 Gong L. and Yeh. E.T.H. (1999) <u>J.Biol.Chem.</u> <b>274</b> :212036-12042 Hemelaar J., <i>et al.</i> (2004) <u>Mol. Cell. Biol.</u> <b>24</b> : 84-95 Hochstrasser M. (2002) <u>Science</u> , <b>298</b> : 549-552 Mendoza H.M., <i>et al.</i> (2003) <u>J.Biol.Chem.</u> <b>278</b> :25637-25643 Reverter D., <i>et al.</i> (2005) <b>345</b> :141-151 Shen L.N., <i>et al.</i> (2005) <u>EMBO. J.</u> <b>24</b> :1341-1351 Wada H., <i>et al.</i> (1999) <u>Biophys. Biochem. Res. Comm.</u> <b>275</b> :100-105 Wu K., <i>et al.</i> (2003) <u>J.Biol.Chem.</u> <b>278</b> :288882-288891
--------------------	--

***For Laboratory Research Use Only, Not For Use in Humans***