

MATERIAL DATA SHEET**SUMO E1 (SAE1/UBA2), *S. cerevisiae***
Cat. # E-311

Conjugation of the ubiquitin-like modifier SUMO (Sentrin) requires the activities of the heterodimeric E1 (Aos1/Uba2) and the Ubch9 E2 enzyme. The dimeric activating enzyme utilizes ATP to adenylate the C-terminal glycine residue of SUMO-1 (also SUMO-2 and SUMO-3), forming a high-energy thiolester bond with the cysteine residue of Uba2 and the release of AMP and PPi. The second step is the trans-esterification reaction whereby SUMO-1 is transferred to Cys⁹³ of Ubch9.

Product Information

Quantity:	25 µg
Stock:	X mg/ml (X µM) in 50 mM HEPES pH 8.0, 150 mM NaCl. Actual concentration will vary with specific Lot #.
Purity:	> 90% by SDS-PAGE
MW:	114 kDa

Use & Storage

Use:	Typical enzyme concentration to support conjugation <i>in vitro</i> is 50-200 nM depending on conditions.
Storage:	Store at -80°C. Avoid multiple freeze/thaw cycles.

Literature

References:	Dohmen R.J., <i>et al.</i> (1995) <i>J. Biol. Chem.</i> 270 :18099-18109 Gong L., <i>et al.</i> (1999) <i>FEBS Lett.</i> 448 :185-189 Johnson E.S., <i>et al.</i> (1997) <i>EMBO J.</i> 16 :5509-5519 Okuma T., <i>et al.</i> (1999) <i>Biochem Biophys Res Commun.</i> 254 :693698 Tatham M.H., <i>et al.</i> (2001) <i>J. Biol. Chem.</i> 276 :35368-35374
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