

## Product datasheet for **TP760839**

### **NSE2 (NSMCE2) (NM\_173685) Human Recombinant Protein**

#### **Product data:**

<b>Product Type:</b>	Recombinant Proteins
<b>Description:</b>	Purified recombinant protein of Human non-SMC element 2, MMS21 homolog ( <i>S. cerevisiae</i> ) (NSMCE2), full length, with N-terminal HIS tag, expressed in <i>E. coli</i> , 50ug
<b>Species:</b>	Human
<b>Expression Host:</b>	<i>E. coli</i>
<b>Expression cDNA Clone or AA Sequence:</b>	A DNA sequence encoding human full-length NSMCE2
<b>Tag:</b>	N-His
<b>Predicted MW:</b>	27.8 kDa
<b>Concentration:</b>	>0.05 µg/µL as determined by microplate BCA method
<b>Purity:</b>	> 80% as determined by SDS-PAGE and Coomassie blue staining
<b>Buffer:</b>	25 mM Tris-HCl, pH 8.0, 150 mM NaCl, 1% sarkosyl, 10% glycerol
<b>Note:</b>	For testing in cell culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process.
<b>Storage:</b>	Store at -80°C.
<b>Stability:</b>	Stable for 12 months from the date of receipt of the product under proper storage and handling conditions. Avoid repeated freeze-thaw cycles.
<b>RefSeq:</b>	<a href="#">NP_775956</a>
<b>Locus ID:</b>	286053
<b>UniProt ID:</b>	<a href="#">Q96MF7</a> , <a href="#">A0A024R9J6</a>
<b>RefSeq Size:</b>	1225
<b>Cytogenetics:</b>	8q24.13
<b>RefSeq ORF:</b>	741
<b>Synonyms:</b>	C8orf36; MMS21; NSE2; ZMIZ7



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**Summary:**

This gene encodes a member of a family of E3 small ubiquitin-related modifier (SUMO) ligases that mediates the attachment of a SUMO protein to proteins involved in nuclear transport, transcription, chromosome segregation and DNA repair. The encoded protein is part of the structural maintenance of chromosomes (SMC) 5/6 complex which plays a key role genome maintenance, facilitating chromosome segregation and suppressing mitotic recombination. A knockout of the orthologous mouse gene is lethal prior to embryonic day 10.5. Naturally occurring mutations in this gene, that abolish the SUMO ligase activity, are associated with primordial dwarfism and extreme insulin resistance. [provided by RefSeq, Mar 2017]

**Product images:**