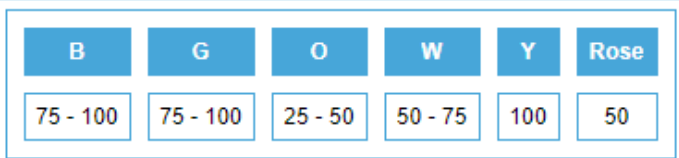


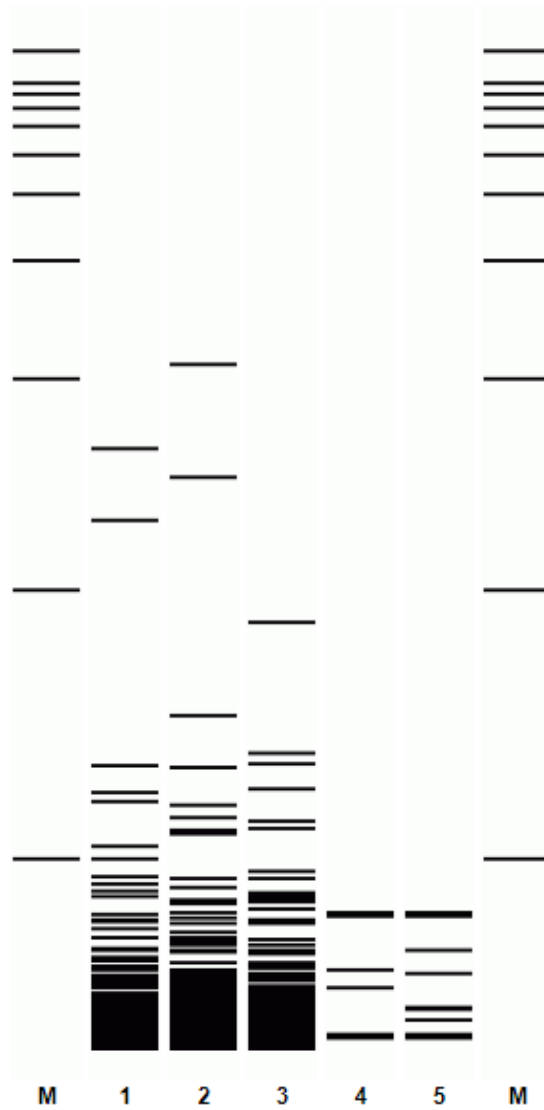
# Mbo II

Cat# RE0471/RE0472– 200 u.a./1000 u.a.

Storage -20°C for 3 years powder | -80°C for 6 months in solvent

## INTFORMATION

<b>Product Name</b>	Mbo II
<b>Cat NO.</b>	RE0471/RE0472
<b>Size</b>	200 u.a./1000 u.a.
<b>Concentration</b>	5000 u.a./mL
<b>Recognition site</b>	GAAGA(N) <sub>8</sub> ↑ CTTCT(N) <sub>7</sub> ↓
<b>Source</b>	An E.coli strain, that carries the cloned gene Mbo II from Moraxella bovis
<b>Assayed on</b>	Lambda DNA (dam-)
<b>Unit definition</b>	One unit of the enzyme is the amount required to hydrolyze 1 µg of Lambda DNA (dam-) in 1 hour at 37°C in a total reaction volume of 50 µl.
<b>Optimal SE-buffer</b>	Y (33 mM Tris-acetate (pH 7.9 at 25°C); 10 mM magnesium acetate; 66 mM potassium acetate; 1 mM DTT.)
<b>Enzyme activity (%)</b>	
<b>Optimal temperature</b>	37°C
<b>Storage buffer</b>	10 mM Tris-HCl (pH 7.5); 50 mM NaCl; 0,1 mM EDTA; 1 mM DTT; 200 µg/ml BSA; 50% glycerol. Store at -20°C.
<b>Reagents Supplied with Enzyme</b>	10 X SE-buffer Y
<b>Ligation</b>	After 5-fold overdigestion with enzyme approximately 60% of the DNA fragments can be ligated and recut. In presence of 10%PEG ligation is better.
<b>Non-specific hydrolysis</b>	No nonspecific activity was detected after incubation of 1 µg of Lambda DNA with 5 u.a. of enzyme for 16 hours at 37°C.
<b>Methylation sensitivity</b>	Blocked by overlappin dam-methylation(G <sup>m</sup> ATC): <b>GAAGATC</b> .
<b>Inactivation</b>	20 minutes under 65°C.
<b>References</b>	Brown N.L., Hutchison C.A. III, Smith M. J. Mol. Biol. 140: 143-148 (1980).



M - ladder, 1 - Adeno-2 DNA, 2 - Lambda DNA, 3 - T7 DNA, 4 - pUC19, 5 - pBR322

**Theoretical diagrams of DNA digestion by this enzyme for the most known DNA substrates:**

To view the fragments length values please point mouse cursor over diagram

Fragment lengths

**PRODUCT USE LIMITATION**

These products are intended for research use only.