

cat.# 1017-10



10 000 U

## Mutanolysin recombinant



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## Mutanolysin recombinant

- \* Recombinant enzyme obtained in bacterial expression system
- \* High purity >95%

Mutanolysin (EC 3.2.1.17) (N-acetylmuramidase) is a muralytic enzyme that cleaves the  $\beta$ -N-acetylmuramyl-(1->4)-N-acetylglucosamine linkage of the bacterial cell wall polymer peptidoglycan-polysaccharide. Its carboxy terminal moieties are involved in the recognition and binding of unique cell wall structures abundant in many gram-positive bacteria.

Recombinant mutanolysin effectively lyses particularly problematic bacteria. Including but not limited to: *Streptococcus*, *Lactobacillus*, *Lactococcus*, *Listeria*.

Recombinant mutanolysin and lysozyme activity is synergistic. Using mutanolysin and lysozyme mixture leads to increased yield of bacteria lysis.

### Application:

1. Mild conditions formation of spheroplasts of gram-positive bacteria
2. Enzymatic cell lysis in DNA and RNA isolation process
3. Effective lysis of gram-positive bacteria in environmental studies and DNA-based microbial detection

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## Mutanolysin recombinant, lyophilisate

### Recommended protocol (for cell wall digestion):

1. Trasfer 0.2-1.0 ml of overnight bacterial culture to 1.5 ml Eppendorf tube and centrifuge (i.e. 2500 x g, 5 min).
2. Discard supernatant and suspend the bacterial pellet in 100  $\mu$ l of the digestion buffer (suggested buffer: 50 mM MES pH 6.0, 1 mM  $MgCl_2$ ). Different digestion buffers may also be tested.  
**Note:** The mutanolysin activity may strongly differ among various strains of Gram-positive bacteria tested.
3. Add 50 U of mutanolysin. Mix the contents and incubate for 20 min at 50 °C.

For best isolation results we suggest to use the mutanolysin with the following dedicated kits:  
Genomic Mini AX Bacteria+ Spin (cat. # 060-100MS) or  
Genomic Mini AX Bacteria+ (cat. # 060-60M)

**Concentration:** 10 U/ $\mu$ l  
**Enzyme form:** solution

**Store at -20 °C**

### Unit definition:

One unit will produce a  $\Delta A_{600}$  nm of 0.01 per minute at 50 mM MES pH 6.0, 1 mM  $MgCl_2$  at 37 °C in a 1 ml volume using suspension of *Streptococcus faecalis* cell wall as substrate.

For R&D use only

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