

# **TelN Protelomerase**

### T767021

<b>Product Component</b>	250 U	1000 U
TelN Protelomerase	25µL	100µL

10×TelN Reaction Buffer 250μL 1mL

**Storage/Transportation Condition** Store at -20°C for up to 12 months. Avoid repeated freeze/thaw cycles. Transport on dry ice.

Form Liquid

Source E.coli strain that carries TelN from phage N15

Storage Buffer 10 mM Tris-HCl, 100 mM NaCl, 0.1 mM EDTA, 1 mM DTT, 50% Glycerol, pH 7.4

**10X TelN Reaction Buffer** 200 mM Tris-HCl, 100 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>,100 mM KCl, 20 mM MgSO<sub>4</sub>, 1% Triton X-100, pH 8.8

Concentration 10U/µL

**Unit Definition** One unit is defined as the amount of enzyme required to cleave 0.5  $\,\mu g$  of Bsal linearized pMiniT-TelN control plasmid (313 fmol TelN recognition site) in a total reaction volume of 50  $\,\mu L$  at 30°C for 30 minutes.

#### **Product Description**

TelN Protelomerase is cloned from bacteriophage N15. TelN cuts dsDNA at the recognition site TelRL (56 bp), which consists of a palindromic sequence-TelO in the middle and palindromic sequences R3 and L3 at both ends of 14 bp form (Figure 1). TelN has cutting-ligating enzyme activity and covalently ligates at the cleavage site forming hairpin termini (Figure 2).



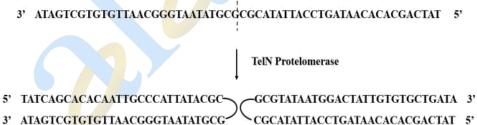


Figure 1. TelRL site



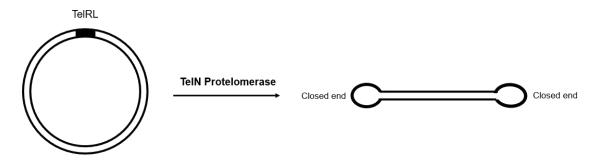


Figure 2. Circular Plasmid Linearization

### **Applications**

In vitro enzymatic synthesis of DNA constructs.

# **Recommended Protocol for Digestion**

1. Make the reaction mixture according to the table below:

Reagent Quantity dsDNA X  $\mu g$  10X TelN Reaction Buffer 2  $\mu L$  TelN Protelomerase (10U/ $\mu L$ ) 1  $\mu L$  Nuclease-free H<sub>2</sub>O Up to 20  $\mu L$ 

2.Gently mix the reaction by pipetting up and down and spin for a few seconds.

- 3.Incubate at 30°C for 30 minutes.
- 4. Heat inactivation at 75°C for 5 minutes.

#### **Notes**

1. For research use only.

