

Notes on Use:

CERTIFICATE OF ANALYSIS

Ligation reaction protocol (for 20 µl)

2 x Quick Ligation Buffer -

2. Incubate at room temperature (+25 °C) for 5 min.

1.Quick T4 DNA Ligase (recombinant, 2000 u/µl) - 20µl

(132 mM Tris-HCl (pH 7.6 at 25 °C); 20 mM MgCl₂, 2 mM

3. Cool the mixture on ice, store at -20°C.

The Quick Ligation Kit includes:

2. 2 x Quick Ligation Buffer - 200µl

DTT; 2 mM ATP; 15% PEG 6000). Mix thoroughly before use.

Add 1 µl Quick T4 DNA Ligase

calculated quantity

10 µl

H₂O -

Add DNA

Before the first use the 2X SE-DNA Quick Ligation Buffer should be divided into small aliquots and store at -20 °C. Avoid defrosting this Buffer more than 2-3 times. The Buffer can be stored at +4 °C during 7 days. The efficiency of ligation starts to decrease after 2 hours of incubation and is reduced by up to 75% if the reaction is allowed to go overnight at 25 °C. Before using the products of a Quick Ligation reaction for electrotransformation, it is necessary to reduce the PEG concentration by precipitation mith Ethanol, and resuspending in H₂O.