

# **Certificate of Analysis**

TOPO<sup>™</sup> TA Cloning<sup>™</sup> Kit for Sequencing, without competent cells

Product No.	4500	30, 11533127
	Lot No.	2852468
Date of Manufacture		05-Jan-2024

## **Restriction Enzyme Analysis**

The parental supercoiled plasmid is qualified by restriction digest to confirm its identity prior to linearization and adaptation with topoisomerase I. Restriction digests must demonstrate the correct banding pattern when electrophoresed on an agarose gel. The table below lists the restriction enzymes and the expected fragments.

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Restriction Enzyme	Expected Fragments	(bp)
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Pme I	3957 (linearize)
Xba I	3957 (linearize)
<i>Bsp</i> H I	1008, 2949
Fsp I	899, 1132, 1926
EcoR I and Afl III	16, 408, 716, 2789

**Note:** The *Xba* I site is removed during the linearization and adaptation process and is not present in the sequence of the linearized, adapted vector.

### **Results: Meets specification**

### **TOPO Cloning**

Each lot is qualified using control reagents included in the TOPO TA Cloning® Kit for Sequencing. Under conditions described in the on-line manual, a 750 bp control PCR product is amplified and TOPO® Cloned into pCR®4-TOPO® and subsequently transformed into the One Shot® competent *E. coli*. The following results must be obtained:

 <u>></u> 95% cloning efficiency when vector + PCR insert colonies are analyzed by EcoR I digestion and agarose gel electrophoresis.

2)  $\leq$  5% of foreground will be produced in the vector-only reaction.

### **Results: Meets specification**

### **Sequencing Primers**

Sequencing primers are lot-qualified by DNA sequencing experiments using the dideoxy chain termination

technique. Each primer must yield  $\geq$  250 bp of quality sequence from a supercoiled plasmid template using standard sequencing conditions.

### **Results: Meets specification**

For Research Use Only. Not for use in diagnostic procedures.

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