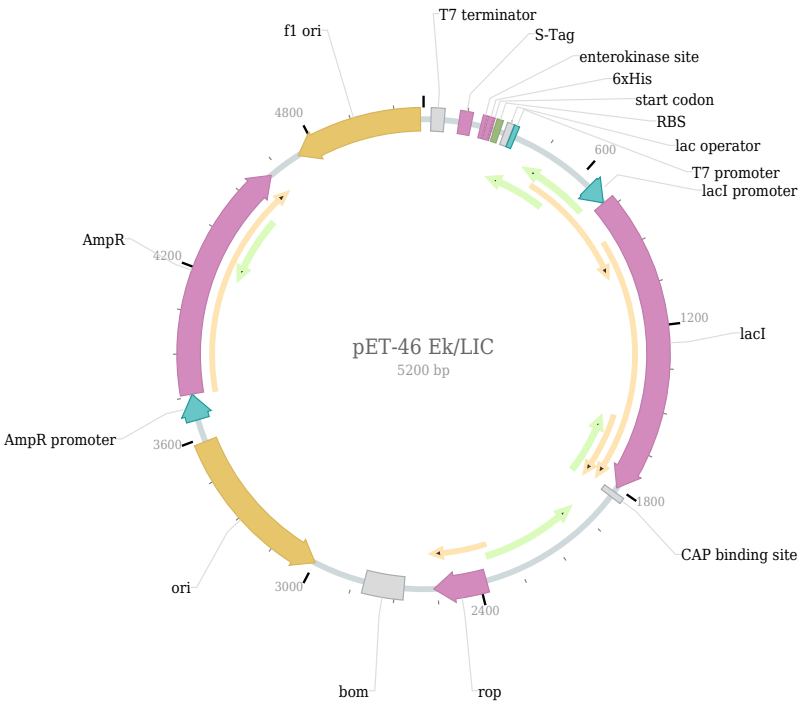
pET-46 Ek/LIC

Vector ID	V010959
Antibiotic Resistance	Ampicillin
Length	5200 bp
Vector Type	pET & Duet Vectors (Novagen)

Product Information

Replication Origin	ori
Source/Author	Novagen (EMD Millipore)
Copy Number	High copy number
Cloning Method	Enzyme digestion and ligation
3' Primer	T7t: 5'-TGCTAGTTATTGCTCAGCGG-3'
Fusion Tag	N-His, N-EK, C-S
Growth Strain(s)	Top10
Comment	The pET-46 Ek/LIC vector is prepared for rapid, directional cloning of PCR-amplified DNA for highlevel expression of polypeptides. Using specifically designed primers for amplification and the pET-46 Ek/LIC Cloning Kit (Cat. No. 71335-3), inserts can be efficiently cloned without the need for restriction digestion or ligation. Fusion proteins contain an N-terminal cleavable His•Tag sequence. Fusion to an optional C-terminal S•TagTM sequence can also be created for detection, purification and quantification of fusion proteins. Unique sites are shown on the circle map. Note that the sequence is numbered by the pBR322 convention, so the T7 expression region is reversed on the circle map. The f1 origin is oriented so that infection with helper phage will produce virions containing single stranded DNA that corresponds to the coding strand. Therefore, single stranded sequencing should be performed using the T7 Terminator Primer (Cat. No. 69337-3).

Vector Map



Plasmid Resuspension Protocol

1. Centrifuge at 5,000×g for 5 min.
2. Carefully open the tube and add sterile water to dissolve the DNA: add 20 µl for 5 µg plasmid, and 100 µl for 100 µg plasmid.
3. Close the tube and incubate for 10 minutes at room temperature.
4. Briefly vortex the tube and then do a quick spin to concentrate the liquid at the bottom. Speed is less than 5000×g.
5. Store the plasmid at -20 ℃.