The full-length, active T7 Endo I is generated
in vitro by ligating a synthetic peptide, consisting of the truncated amino acid residues, to the
thioester-tagged tT7 Endo I (1).

Applications:
• Resolve four-way junction or branched DNA
• Detect or cleave heteroduplex and nicked DNA
• Randomly cleave linear DNA for shot-gun cloning

Supplied in: 200 mM NaCl, 20 mM Tris-HCl
(pH 7.5), 0.1 mM EDTA, 1 mM dithiothreitol,
0.15% Triton X-100 and 50% glycerol.

Reagents Supplied with Enzyme:
10X NEBuffer 2

Note: pUC(AT) is derived from pUC19 with a
modification of the polylinker between the EcoRI
site and the PstI site.

Reaction Conditions: 1X NEBuffer 2.
Incubate at 37°C.

The full-length, active T7 Endo I is generated
in vitro by ligating a synthetic peptide, consisting
of the truncated amino acid residues, to the
thioester-tagged tT7 Endo I (1).

Applications:
• Resolve four-way junction or branched DNA
• Detect or cleave heteroduplex and nicked DNA
• Randomly cleave linear DNA for shot-gun cloning

Supplied in: 200 mM NaCl, 20 mM Tris-HCl
(pH 7.5), 0.1 mM EDTA, 1 mM dithiothreitol,
0.15% Triton X-100 and 50% glycerol.

Reagents Supplied with Enzyme:
10X NEBuffer 2

Note: pUC(AT) is derived from pUC19 with a
modification of the polylinker between the EcoRI
site and the PstI site.

Reaction Conditions: 1X NEBuffer 2.
Incubate at 37°C.

The full-length, active T7 Endo I is generated
in vitro by ligating a synthetic peptide, consisting
of the truncated amino acid residues, to the
thioester-tagged tT7 Endo I (1).

Applications:
• Resolve four-way junction or branched DNA
• Detect or cleave heteroduplex and nicked DNA
• Randomly cleave linear DNA for shot-gun cloning

Supplied in: 200 mM NaCl, 20 mM Tris-HCl
(pH 7.5), 0.1 mM EDTA, 1 mM dithiothreitol,
0.15% Triton X-100 and 50% glycerol.

Reagents Supplied with Enzyme:
10X NEBuffer 2

Note: pUC(AT) is derived from pUC19 with a
modification of the polylinker between the EcoRI
site and the PstI site.

Reaction Conditions: 1X NEBuffer 2.
Incubate at 37°C.
T7 Endonuclease I is a structure-selective enzyme. It acts on a variety of DNA substrates with different specific activities. It is important to control the amount of enzyme and the reaction time used for cleavage of a particular substrate.

This enzyme is not recommended to be used at 55°C, as the activity is decreased.

References: