

Restriction Endonuclease**Fai I****E551**

SibEnzyme
SibEnzyme Ltd., Russia
Ph: +7 383 333 4991
Fax :+7 383 333 6853
info@sibenzyme.com
www.sibenzyme.com

**50 u****Lot: 20****1000 u/ml****Store at -20°C****Recognition Sequence:**

5'... YA↓TR ...3'

3'... RT↑AY ...5'

Source: *Flavobacterium aquatile B15*

Fail cleaves 4 expected recognition sites as well as several other sites with a weaker activity. In the case of long incubation with Fail DNA can be digested to small oligos.

Supplied in:

10 mM Tris-HCl (pH 7.5); 100 mM KCl;
0,1 mM EDTA; 200 µg/ml BSA;
7 mM 2-mercaptoethanol; 50% glycerol.

Reaction Conditions:

1×SEBuffer B.

Incubate at **50°C**.

Warranty period for the enzyme storage at-20°C is two years from the date of the last assay indicated on the enzyme vial.

1×SEBuffer B (pH 7.6 @ 25°C)10 mM Tris-HCl 10 mM MgCl₂ 1 mM DTT**Unit Definition:**

One unit is defined as the amount of enzyme required to cleave 1 pmol of the double-stranded oligonucleotide with the following structure

5'-CGAGTTCA^TAGCTGGGCCCAAC-3'**3'-GCTCAAGT^ATCGACCCGGGTG-5'**

in 1 hour at 50°C in a total reaction volume of 20 µl.

http://science.sibenzyme.com/article8_article_51_1.phtml

Quality Control Assays

Ligation: After 3-fold overdigestion with Fai I, ~50% of pUC19 DNA fragments can be ligated with T4 DNA Ligase and recut.

Oligonucleotide Assay:

No detectable degradation of a single- and double-stranded oligonucleotide was observed after incubation with 1 unit of enzyme for 3 hours.

Enzyme Properties**Activity in SEBuffers:**

SEBuffer B **100%**
SEBuffer G 50-75%
SEBuffer O 10-25%
SEBuffer W 25-50%
SEBuffer Y 25-50%
SEBuffer ROSE 100%

Heat Inactivation:**Yes** (80°C for 20 minutes)**Reagents Supplied with Enzyme:**

10×SEBuffer B

CERTIFICATE OF ANALYSIS