

Product: RNase H2 Enzyme, 500 Units, 20 U/μL (25 μL)

Product number: 11-03-02-03

Batch number: 0000511963

Expiration date: 2021-SEP-29

Test	Specification	Results
Physical purity assessment by electrophoresis separation analysis	Purity >90%	Pass
qPCR DNA contamination test for <i>E. coli</i> 16S rRNA subunit	<10 Copies/sample	Pass
Molecular weight determination by electrophoresis separation analysis	27.6 kDa ±15%	Pass
ESI mass spectral analysis of enzyme cleavage products	4335 ±0.05% Da 5132 ±0.05% Da	Pass
Enzyme activity concentration	20 ±25% U/μL	Pass

Storage: Store RNase H2 Enzyme in a sealed container at -20°C.

Verified by: Ekaterina Carlson Quality release date: 2020-APR-01

IDT verifies that the information contained herein is true and correct to the best of our knowledge. This document was produced electronically and is valid without signature.



certificate of analysis

General information: RNase H2 (rnhB) belongs to a family of enzymes that cleaves RNA bases in a DNA-RNA duplex. RNase H2 can cleave a duplex embedded with as little as a single RNA base leaving a 3′ OH and a 5′ phosphate. The enzyme does not cleave single-stranded RNA or DNA.

Enzyme source: RNase H2 gene from *Pyrococcus abyssi*, codon optimized and expressed in *E. coli*.

Storage buffer: 20 mM Tris-HCl, pH 8.4; 0.1 mM EDTA; 100 mM KCl; 50% glycerol; 0.1% Triton X-100

Concentration: 20 U/µL

Recommended reaction conditions:

- 20 mM Tris-HCl, pH 8.4; 60 mM KCl; 0.01% Triton X-100; 3 mM MgCl₂; 1% glycerol
- Optimal RNase H2 cleavage activity occurs at 70°C, but the enzyme works well in the range of 60–75°C. Activity drops off rapidly below 60°C. For use in conjunction with PCR (polymerase chain reaction), the recommended cycling parameters are (95°C, 15 sec; 60°C, 1 min) x 45. If a hotstart DNA polymerase is employed, then use the manufacturer's recommended activation temperature and time.

Unit definition: One enzymatic unit is the amount of enzyme needed to cleave 1 nmol of DNA-RNA-DNA heteroduplex substrate (S-rC 14-1-15) per minute at 70°C in Magnesium Cleavage Buffer (10 mM Tris-HCl, pH 8.0; 50 mM NaCl; 4 mM MgCl₂; 10 µg/mL BSA).

S-rC 14-1-15:

- 5' CTCGTGAGGTGATG<mark>rC</mark>AGGAGATGGGAGGCG 3'
- 3' GAGCACTCCACTAC GTCCTCTACCCTCCGC 5'

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