Maxima H Minus Reverse Transcriptase

Engineered to impress

Thermo Scientific™ Maxima™ H Minus Reverse Transcriptase enables consistent success in cDNA synthesis in RT-PCR and RT-qPCR applications. Developed through molecular evolution, with the introduction of multiple favorable mutations into wild-type Moloney murine leukemia virus (M-MuLV) reverse transcriptase, the enzyme delivers impressive thermostability, processivity, and cDNA synthesis rates.

Why use Maxima H Minus Reverse Transcriptase?

Maxima H Minus Reverse
Transcriptase is an enzyme with
significantly increased thermostability,
enhanced processivity, and
diminished RNase H activity. Due
to these improved features, the
enzyme consistently produces high
yields of full-length cDNA even at
elevated temperatures. The enzyme
is available as a stand-alone enzyme
or formulated for convenient kits to
support different needs for first-strand
cDNA synthesis.

Features

- High yields of cDNA over a wide temperature range (up to 65°C) (Figure 1)
- Efficient RT-PCR even with extremely long amplification targets (up to 20 kb) (Figure 2)
- Enhanced linearity, sensitivity, and early C, in two-step RT-qPCR (Figure 3)
- Multiple formats, from stand-alone enzyme to first-strand cDNA synthesis kit, double-stranded cDNA synthesis kit, and one-tube reverse transcription (RT) master mix

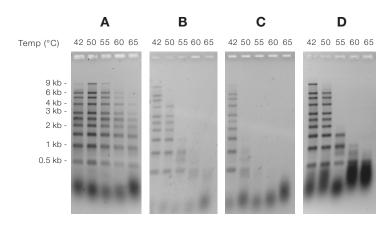


Figure 1. Maxima H Minus Reverse Transcriptase generates full-length cDNA at a wider range of reaction temperatures than reverse transcriptases from other suppliers. cDNA synthesis using 1 µg of Invitrogen™ Millennium™ RNA Markers (poly(A)-tailed) with an oligo(dT)₁₈ primer was performed with (A) Maxima H Minus Reverse Transcriptase; and reverse transcriptases from other suppliers: (B) TaKaRa™ PrimeScript™ Reverse Transcriptase, (C) Promega™ GoScript™ Reverse Transcriptase, at different temperatures per suppliers' instructions. The optimal temperatures are 50°C for Maxima H Minus Reverse Transcriptase and 42°C for reverse transcriptases from other suppliers. cDNA products were detected by alkaline gel electrophoresis.



Technical details

- Supports a wide range of starting total RNA amounts (1 pg–2.5 μg)
- Can synthesize RNA at elevated temperatures (42–65°C)
- RT reaction can be completed in only 15–30 minutes
- Up to 50x higher processivity compared with wild-type M-MuLV reverse transcriptase enzyme
- Demonstrates templateswitching activity
- Capable of incorporating modified nucleotides

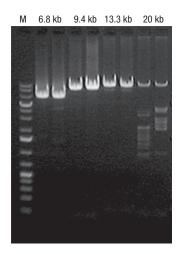


Figure 2. Two-step long RT-PCR from RNA transcripts up to 20 kb. Total RNA was extracted from human and mouse cells and used to reverse-transcribe 6.8 kb and 9.4 kb transcripts from the human RNA, and 13.3 kb and 20 kb transcripts from the mouse RNA, using 1 µg of total RNA per reaction and Maxima H Minus Reverse Transcriptase. The synthesized cDNAs were used as templates for PCR, and the PCR products were visualized by gel electrophoresis. M: Thermo Scientific™ GeneRuler™ 1 kb Plus DNA Ladder.

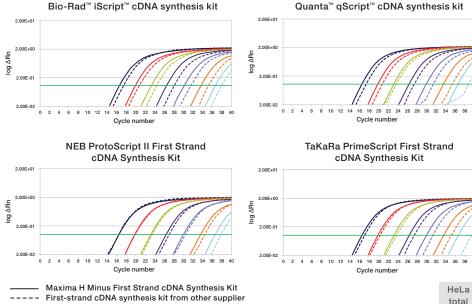


Figure 3. The Thermo Scientific™ Maxima™ H Minus First Strand cDNA Synthesis Kit demonstrates consistently better RT efficiency than that obtained with other suppliers' kits. RT-qPCR of the human β₂ macroglobulin gene was performed on 10-fold serial dilution of 1 μg–1 pg of total RNA from HeLa cells. First-strand cDNA was generated using the Maxima H Minus First Strand cDNA Synthesis Kit and four other commercial first-strand cDNA synthesis kits. cDNA was amplified using Applied Biosystems™ TaqMan® Universal Master Mix II with uracil N-glycosylase (UNG), on the Applied Biosystems™ ViiA™ 7 Real-Time PCR System.

total RNA
1 µg
100 ng
10 ng
1 ng
100 pg
1 ng

Ordering information

Quantity	Cat. No.
2,000 U	EP0751
10,000 U	EP0752
4 x 10,000 U	EP0753
20 rxns/100 rxns	K1651/K1652
20 rxns/100 rxns	K1681/K1682
10 rxns	K2561
50 rxns/200 rxns	M1661/M1662
50 rxns/200 rxns	M1681/M1682
	2,000 U 10,000 U 4 x 10,000 U 20 rxns/100 rxns 20 rxns/100 rxns 10 rxns 50 rxns/200 rxns

Find out more at thermofisher.com/maxima

