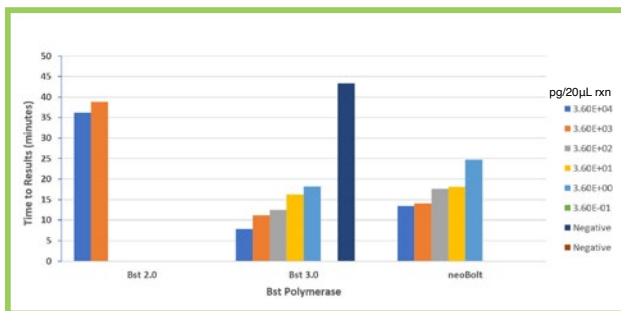


PR1MA™ Bst Polymerase

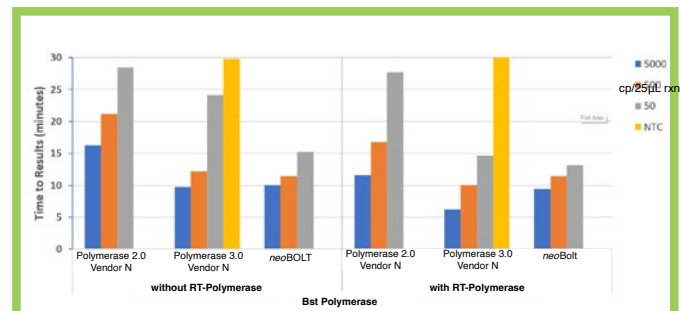
Recombinant, thermostable DNA polymerase for isothermal amplification

PR1MA™ Bst polymerase is a true single-enzyme reagent for developing RT-LAMP assays and is available either in glycerol or in a glycerol-free buffer as a custom order.

DNA LAMP



RT-LAMP



Bst Polymerase 2.0 and 3.0 are commercially available

Properties

- Only Bst Polymerase on the market with robust RT and DNA polymerase activity
- Thermostable, working temperature range 64 - 72°C
- Tolerant to inhibitors
- Use PR1MA™ Bst polymerase to develop LAMP assays with high sensitivity and specificity

MIDSCI also offers a variety of encapsulated RNA controls for common targets, and we would love to work with you to develop a unique control to suit your needs. Contact us at tech@midsci.com for more information.

*These products are intended for research use only, not for diagnostic use. The safety and efficacy of these products in diagnostic or other clinical uses has not been established.

PR1MA™ Bst Polymerase

Contents

PR1MA™ Bst polymerase is provided at a concentration of 8 U/μL with 10X Isothermal buffer.

Background

Bst polymerase is a recombinant, truncated (lacks 5' to 3' exonuclease activity), thermostable *Bacillus stearothermophilus* DNA polymerase with high reverse transcriptase and strand-displacement activities, ideal for isothermal amplification of RNA and DNA targets. PR1MA™ Bst polymerase is engineered to perform at temperatures up to 73°C and tolerate inhibitors, has increased sensitivity and speed relative to other Bst polymerases, and can incorporate dUTP.

Application Notes

Bst polymerase (exonuclease minus), with strong strand-displacement and RT activities can be used for amplification of DNA and RNA in loop-mediated isothermal amplification (LAMP).

Shipping and Storage

- Bst polymerase is supplied in a buffer of 50% glycerol, 50 mM Tris-HCl, 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 0.05% Tween-20, 0.05% NP-40 substitute, pH=7.5. Can be supplied in a glycerol-free buffer as a custom order.

Important note: use of the supplied buffer will yield optimal results.

PR1MA™ Bst polymerase is shipped on dry or blue ice. On arrival store at -20°C for optimum stability. Repeated freeze/thaw cycles should be avoided.

Quality Control

- Bst polymerase unit activity: A known polymerase is used to create a standard curve with a real-time primer extension assay against which the activity of this enzyme is measured.
- Purity: >95% as determined by SDS-PAGE analysis
- Bst polymerase is free of detectable RNase and DNase (exo- and endonuclease).
- <0.05 ng contaminating host DNA per 8 U

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Setting Up LAMP Reaction

- Prior to setting up LAMP reaction, thaw all reaction components.
- Before use, mix all components by vortexing (5 sec) followed by centrifugation (5 sec).
- Setting up reaction on ice (4°C) is highly recommended.

Reaction set up:

Component	Stock	Volume	Final Concentration
¹ 10X Isothermal Buffer	10x	2.5 µL	1x
² MgSO ₄	100 mM	1 µL	4 mM
dNTP Mix	25 mM	1.25 µL	1.25 mM
³ Dye	Variable	Variable	Variable
⁴ Primer Mix	20x	1.25 µL	1x
neoBolt™ Bst polymerase	8 U/µL	1 µL	0.32 U/µL
⁵ RNase Inhibitor (optional)			
⁶ RT polymerase (optional)			
Template	Variable	Variable	Variable
Nuclease-Free Water		to 25 µL	
Total			25

1. 10X Isothermal Buffer contains 20 mM MgSO
2. We recommend adding 4 mM MgSO₄ (on top of the 2 mM MgSO₄ contributed by the 10X Isothermal buffer) to start and optimize your assay from there
3. Intercalating dye (such as SYTO-82, SYTO-9, EvaGreen) are recommended for real time monitoring of amplification in LAMP reactions
4. A LAMP primer mix can be prepared with all 4 or 6 (with Loop) primers. A 20X primer mix should contain: 31.2 uM FIP, 31.2 uM BIP, 4 uM F3, 4 uM B3, 15.6 uM LoopF, 15.6 uM LoopB in TE or water
5. PR1MA™ RNase Inhibitor recommended when using RNA target
6. For RNA targets only. PR1MA™ RT polymerase has strong RT activity, however, for faster time to results and increased sensitivity, the use of RT polymerase is recommended

Recommended Companion Products

- RNA Controls
- RNase Inhibitor
- MS2 Phage
- Cod Uracil-DNA glycosylase