Bst DNA Polymerase Product Handling Guide		Storage and stability: Bst DNA Polymerase is shipped on dry or blue ice. On arrival store at -20 °C for optimum stability. Repeated freeze/thaw cycles should be avoided. Solutions should be mixed/equilibrated after each thawing to avoid phasing.	
		Expiry: When stored under the recommended conditions and handled correctly, full activity of the kit is retained until the expiry date on the outer box label.	
Shipping: Catalog number:	On dry/blue ice MDX012	Safety precautions: Read and understand the SDS (Safety Data Sheets) before handling the reagents. Hardcopies of the SDS will be provided with the first shipment, thereafter they will be available upon request.	
Batch No.: Concentration:	See vial 8 units/µL Store at −20 °C	Quality control: Bioline operates under ISO 13485 Quality Management System. Bst DNA Polymerase and its components are extensively tested for activity, processivity, efficiency, heat activation, sensitivity, absence of nuclease contamination and absence of nucleic acid contamination.	
merid		Notes: For research or further manufactured use only.	

Description

Bst DNA Polymerase is a DNA polymerase (exonuclease minus), with strand-displacement properties. Bst DNA Polymerase is used for Isothermal DNA amplification and LAMP (Loop-mediated Isothermal Amplification).

Kit components

Table 1

Component	
Bst DNA Polymerase (8 U/μL)	
Bst Reaction Buffer, 10x	
Enzyme Dilution Buffer, 1x	

Users Guidelines

Thawing during transportation does not affect the product performance. Prior to use or storing at -20 °C, the thawed reagents must be thoroughly mixed by 10 inversions.

LAMP Optimisation

It is recommended that $MgSO_4$ (not supplied) is supplemented to a total concentration of 8 mM as indicated in the example LAMP protocol, but can be optimised according to individual assay requirements.

dNTPs can be optimised between 1.6 mM (0.4 mM each) and 6 mM (1.5 mM each) final concentration.

Bst DNA polymerase can be diluted using the Enzyme Dilution Buffer provided to between 0.12 and 0.32 U/ μ L final concentration, depending on individual assay requirements.

Working Concentration of Bst

Bst DNA Polymerase is ready-to-use at a working concentration of 8 U/ μ L.

Typical LAMP reaction conditions:

Incubate at 60 °C for 60 minutes.

The following protocol is for a standard 25 μ L LAMP reaction to be used as a starting point for optimization.

Reagent	Volume	Final Concentration
Bst Reaction Buffer, 10x	2.5 µL	1x (contains 2 mM MgSO ₄)
dNTP Mix (100 mM -25 mM each)	0.4 - 1.5 µL	1.6 - 6 mM
MgSO4 (100 mM)	1.5 µL	6 mM (8 mM total)
FIP/BIP Primers (25X)	1 µL	1.6 µM
F3/B3 Primers (25X)	1 µL	0.2 µM
Loop F/B Primers (25X)	1 µL	0.4 µM
Bst DNA Polymerase (3-8 U/µL)	1 µL	0.12 - 0.32 U/µL
Sample DNA	variable	> 10 copies
Water (ddH ₂ O)	to 25 µL	

Related Products	Cat. No.
High Conc. Glycerol-Free Bst, 100 U/µL	MDX018
Bst Reaction Buffer, 10x	MDX076
Enzyme Dilution Buffer, 10x	MDX080
Enzyme Dilution Buffer, 1x	MDX078
dNTP Mix, 100mM	MDX051

Technical Support

For any technical enquiries, please contact our Technical Support team via email at: mbi.tech@meridianlifescience.com

Bioline USA Inc. USA

Tel: +1 901 382 8716 Fax: +1 901 382 0027