

FAST HotStart DNA polymerase with Buffer with included dNTPs

FAST HotStart DNA polymerase is a robust enzyme with hotstart function amplifying with extreme speed, yield, and consistency

Component	Cat#	M3287.0500	M3287.2500	Colour code of cap
FAST HotStart DNA polymerase (5 units/μL)		500 units	5x 500 units	red
5X buffer for FAST Hotstart DNA polymerase (buffer contains dNTPs and MgCl ₂)		4x 1 mL	20x 1mL	colourless

Product description

Genaxxon's FAST HotStart DNA polymerase is a robust enzyme with a hotstart function, ideally suited for applications like genotyping and screening, amplifying with extreme speed of up to 2 seconds per 1kb, as well as high yield and consistency. PCR products generated with this enzyme are A-tailed and can thus be cloned into TA cloning vectors. The extreme speed of Genaxxon's FAST HotStart DNA polymerase allows the use of an extension rate of 4-8 kb/min. The buffer composition has been optimized via high-throughput screening and makes the enzyme particularly resistant to PCR inhibitors. Therefore, Genaxxon's FAST HotStart DNA polymerase is suitable for direct PCR from unpurified samples, including blood, urine, and bacterial colonies, making this the ideal choice for consistent results in fast complex PCR amplifications.

Product Specifications

Concentration:	5 units/μL
Extension rate:	Up to 30 kb/min. at 72 °C
Amplicon size	Up to 5kb
Hotstart	Yes
5'-3' exonuclease activity:	Yes
Extra addition of A:	Yes
3'-5' exonuclease activity:	No
Nuclease contamination:	No
Protease contamination:	No
RNase contamination:	No

FAST Hotstart DNA Polymerase is supplied at a concentration of 5U/μL. The also supplied reaction buffer (5x) includes not only 15mM MgCl₂, but also 5mM dNTPs, enhancers, and stabilizers, optimized to increase PCR success rates.

Source

Recombinant, purified from *E.coli*

Applications

Direct PCR
 Fast amplification of complex targets (including GC-rich and AT-rich templates)
 Colony PCR
 Multiplex PCR

Storage

FAST DNA HotStart polymerase, including buffers and reagents, should be stored immediately upon receipt at -20 °C. When stored under these conditions and handled correctly, these products can be kept at least until the expiration date (see tube label) without showing any reduction in performance. No loss of performance is detected after 20 freeze/thaw cycles.

Quality Control

Functionally tested in PCR. Absence of endonucleases, exonucleases, and ribonucleases was confirmed by appropriate assays.

Product Use Limitations

FAST DNA HotStart polymerase is developed, designed, and sold for research purposes only. It is not to be used for human, diagnostic or drug purposes or to be administered to humans unless expressly cleared for that purpose by the Food and Drug Administration in the USA or the appropriate regulatory authorities in the country of use. All due care and attention should be exercised in the handling of many of the materials described in this manual.

Safety information

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate material safety data sheets (MSDSs). These are available online as pdf-file or on request (info@genaxxon.com).

More information can be found in the REGULATION (EC) No. 1272/2008 OF THE EUROPEAN PARLIAMENT AND THE COUNCIL or contact Genaxxon bioscience (info@genaxxon.com)

PCR Protocol Part

Protocol using FAST HotStart DNA Polymerase

This protocol serves as a guideline for PCR amplification. Optimal cycling conditions such as incubation times, temperatures, and amount of template DNA depend on the DNA target (GC-content, size, quantity, purity, etc.), primers, buffer composition, MgCl₂-concentration and enzyme concentration and must be determined individually. We recommend starting with the basic protocol described below and subsequently optimize the annealing temperature, incubation times and cycling numbers.

Please note: The FAST HotStart Reaction Buffer already includes dNTPs and has been optimized with respect to the MgCl₂-concentration and other components to guarantee highest success rates. Therefore, we do not recommend adding additional MgCl₂ or other enhancers.

Important notes before getting started

Set up all reaction mixtures in an area separate from that used for DNA preparation or PCR product analysis.

Use disposable tips containing hydrophobic filters to minimize cross-contamination.

Procedure

- 1. Prepare a master mix according to Table 1.**
To minimize the risk of contamination or reagent loss and to improve pipetting accuracy, we recommend preparing a master mix for multiple samples containing all the components needed for PCR except the template DNA. Prepare a volume of master mix 10% greater than that required for the total number of PCR assays to be performed. A negative control (without template DNA) should always be included in every experiment.
- 2. Mix your master mix with all components except the template DNA** starting with the component with the greatest volume (usually water) and ending with the FAST HotStart DNA polymerase. Keep the master mix on ice.

Table 1: PCR reaction components (25µL PCR reaction)

Components	Quantities	Final Concentration
FAST HotStart Reaction Buffer (5X)	5µL	1X
Forward primer (5pmol/µL)	2µL	0.4µM
Reverse primer (5pmol/µL)	2µL	0.4µM
Template DNA*	0.25 - 10µL	1 - 250ng**
FAST HotStart DNA polymerase	0.1µL***	0.5U***
PCR grade water	up to 25µL	

* In case of Direct PCR: use 1µL of mammalian blood or urine per 25µL reaction volume

In case of Colony PCR: use a sterile tip to resuspend a colony (or 5µL of an overnight culture) directly in the PCR reaction mix.

** In case of cDNA <50ng and in case of gDNA <250ng (total amount)

*** For difficult targets, e.g., GC-rich, higher enzyme concentration may be required (up to 2.5U).

For smaller/larger reaction volumes, scale it down/up proportionally.

- 3. Mix the master mix thoroughly** and dispense appropriate volumes into PCR tubes.
Mix gently, for example, by pipetting the master mix up and down. Do not vortex as this may harm the enzyme. It is recommended that PCR tubes are kept on ice before placing in the thermal cycler.
- 4. Add template DNA** (or PCR grade water in case of the control) to the individual tubes containing the master mix.
- 5. Program the thermal cycler** according to the manufacturer's instructions.
A typical PCR cycling program is outlined in Table 2. For maximum yield and specificity, temperatures and cycling times should be optimized for each new target or primer pair.
- 6. Place PCR tubes in the thermal cycler** and start program.

Table 2: PCR conditions (Thermal cycler)

Step	time	temperature	comments
Initial denaturation:	1 min ¹	95 °C	denaturation of template DNA including removal of all secondary DNA structures such as hairpins
3-step cycling²			
Denaturation:	15 sec	95 °C	
Annealing:	15 sec	Ta ³	Ta ³ should be 2 °C below the Tm ⁴ value of the primer with the lowest Tm
Extension:	2-15 sec ⁵	72 °C	2 sec for targets <1kb, 15sec/kb for targets >1kb
Number of Cycles	40		
Final extension	3 min	72 °C	

¹ In case of Colony PCR, increase initial denaturation time to 10 min.

² In case of Multiplex PCR, use 90 seconds in total per cycle independent of the sizes of the amplicons.

³ Ta is the annealing temperature

⁴ Tm is the melting temperature of a primer which is defined as the temperature at which 50% of the primer bind to the complementary sequence of the target DNA.

Please note: By increasing Ta above Tm, this percentage decreases, however, primers will still anneal up to a certain degree and initiate extension. Therefore, PCR would still work with a Ta of several degrees higher than the Tm but with a dramatically reduced efficiency. Hence, we recommend optimizing the Ta by performing a temperature gradient (e.g., starting at the lowest Tm or a few degrees below and increasing with 2 °C increments).

⁵ Largest tested amplicon size: 5kb. Please note that longer extension times may be required for targets larger than 5kb!

Recommendations

- **Annealing temperature (Ta)**
Optimization of the annealing temperature is important to guarantee an optimal amplification. We recommend performing a temperature gradient and using primers with a Tm >60 °C.
- **Primer concentration**
The final primer concentration should be in the range of 0.2µM to 0.6µM.

Optimization potential for incubation times and number of cycles

Denaturation and annealing steps may require less time depending on the thermocycler you're using (ramp rate), the reaction volume, the PCR tube (varies with the efficiency of heat transfer) and your template primer combination.

Therefore, it might be worthwhile to optimize times and cycle number depending on the success of your amplification. To greatly reduce the overall PCR time, you can try to:

- reduce times to as low as 10 seconds for both steps, annealing and denaturation.
- reduce the number of cycles to 25-30.

Please note: Fast cycling conditions should not be applied in multiplex PCR. The recommended 90 seconds per cycle may be further extended to increase the yield.

Trouble shooting

Non-specific amplification	Ta too low	Increase Ta in 2 °C increments. Choose primers with a Tm above 60 °C.
	MgCl ₂ concentration too high	Decrease MgCl ₂ if possible.
	Primer concentration too high	Final concentration should be between 0.2 and 0.6 µM for each primer.
No amplification	Ta too high	Decrease Ta in 2 °C increments.
	MgCl ₂ concentration too low	Increase MgCl ₂ concentration.
	Forgot to pipet template DNA	Make sure that template DNA and primers are included and repeat PCR.
	Primer concentration too low	Final concentration should be between 0.2 and 0.6 µM for each primer.