

HOT DISCO TAQ 5X MASTER MIX, LYOPHILISE

Cat. No.: 257701

500 reactions

Reaction size 25 µl



Foil bags CL0.500-0058 with:		
Content	Hot Disco Taq 5x Master Mix, Lyophilise 2 mM MgCl ₂	5x Buffer, Lyophilise
	1 foil bag x 4 tubes	2 tubes x 1.3 ml
ID No.	CL0.125-0056	CL1.300-0054
Cap colour	Blue	Black

Key Features

- Lyophilized Hot Disco Taq Master Mix
- Highly stable at room temperature
- Shipping at ambient temperature
- Storage at room temperature
- Reduced CO₂ impact
- For amplification up to 4 kb

Hot Disco Taq 5x Master Mix, Lyophilise is lyophilized for storage at room temperature and shipping at ambient temperature. Hot Disco Taq 5x Master Mix, Lyophilise is easily reconstituted with 5x Buffer, Lyophilise included in the kit.

The reconstituted Hot Disco Taq 5x Master Mix, Lyophilise is a ready-to-use 5x reaction mix with everything needed to perform PCR; Hot Disco Taq, optimized NH₄⁺ PCR buffer system, dNTPs and magnesium chloride present. Each reaction requires 5 µl of the 5x Master Mix. Simply add primers, template and water to a total reaction volume of 25 µl to successfully carry out PCR.

Hot Disco Taq 5x Master Mix, Lyophilise is a cost saving choice as it can be shipped at ambient temperature without the presence of dry ice. Furthermore, the reduced electricity consumption and greenhouse gas emission leave an improved CO₂ imprint.

Recommended Shipping, Storage and Stability

- Shipping at ambient temperature.
- Hot Disco Taq 5x Master Mix, Lyophilise and 5x Buffer, Lyophilise are stable at room temperature (≤ 25 °C) for 12 months from date of receipt.
- Expiry when stored at -20 °C is stated on the kit label.
- The reconstituted master mix is stable for 6 months at -20 °C or for 3 months at +4 °C.

Quality Control

Hot Disco Taq is tested for contaminating activities, with no traces of endonuclease activity, nicking activity or exonuclease activity.

Protocol

This protocol serves as a guideline to ensure optimal PCR results when using Hot Disco Taq 5x Master Mix, Lyophilise. Optimal reaction conditions such as incubation times, temperatures, and amount of template DNA may vary and must be determined individually.

Hot Disco Taq 5x Master Mix, Lyophilise must be reconstituted using the 5x Buffer, Lyophilise provided in the kit.

1. Reconstitute each vial of Hot Disco Taq 5x Master Mix, Lyophilise with 625 µl of 5x Buffer, Lyophilise. Vortex for 30 seconds then incubate at RT for 1 min and finally vortex again until fully dissolved.

Important: After reconstitution, the Hot Disco Taq 5x Master Mix, Lyophilise should be stored at -20 °C or +4 °C.

2. Thaw the reconstituted Hot Disco Taq 5x Master Mix, Lyophilise and primers. **It is important to thaw the solutions completely and mix thoroughly before use to avoid localized concentrations of salts.**
3. Prepare a reaction mix. Table 1 shows the reaction set up for a final volume of 25 µL. If desired, the reaction size may be scaled down. Use 5 µl of the Hot Disco Taq 5x Master Mix, Lyophilise in a final volume of 25 µl.

Table 1. Reaction components (reaction mix and template DNA)

Component	Vol./reaction*	Final concentration*
Hot Disco Taq 5x MM, Lyophilised	5 µl	1x
25 mM MgCl ₂	0 µl (0 – 2 µl)	2 mM (2 – 4 mM)
Primer A (10 µM)	0.5 µl (0.25 – 2.5 µl)	0.2 µM (0.1 – 1.0 µM)
Primer B (10 µM)	0.5 µl (0.25 – 2.5 µl)	0.2 µM (0.1 – 1.0 µM)
PCR-grade H ₂ O	X µl	-
Template DNA	X µl	Genomic DNA: 50 ng (10 – 500 ng) Plasmid DNA: 0.5 ng (0.1 – 1 ng) Bacterial DNA: 5 ng (1 – 10 ng)
TOTAL volume	25 µl	-

* Suggested starting conditions; theoretically used conditions in brackets

4. Mix the reaction mix thoroughly and dispense appropriate volumes into reaction tubes. Mix gently, e.g. by pipetting the reaction mix up and down a few times.
5. Add template DNA to the individual tubes containing the reaction mix.
6. Program the thermal cycler according to the manufacturer's instructions. See table 2 for an example. For maximum yield and specificity, temperatures and cycling times should be optimized for each new template target or primer pair.
7. Place the tubes in the thermal cycler and start the reaction.

Table 2. Three-step PCR program

Cycles	Duration of cycle	Temperature
1	15 minutes	95 °C
25 - 35	20 – 30 seconds ^a 20 – 40 seconds ^b 30 seconds ^c	95 °C 50 – 65 °C 72 °C
1	5 minutes ^d	72 °C

^a Denaturation step: This step is the first regular cycling event and consists of heating the reaction to 95 °C for 20 – 30 seconds. It causes melting of the DNA template by disrupting the hydrogen bonds between complementary bases, yielding single-stranded DNA molecules.

^b Annealing step: The reaction temperature is lowered to 50 – 65 °C for 20 – 40 seconds allowing annealing of the primers to the single-stranded DNA template. Typically, the annealing temperature is about 3 – 5 °C below the T_m (melting temperature) of the primers used.

^c Extension/elongation step: Hot Disco Taq polymerase has its optimum activity temperature at 72 °C. At this step the DNA polymerase synthesizes a new DNA strand complementary to the DNA template strand. The extension time depends on the length of the DNA fragment to be amplified. As a rule of thumb, at its optimum temperature the DNA polymerase will polymerize a thousand bases per minute.

^d Final elongation: This single step is occasionally performed at a temperature of 72 °C for 5 minutes after the last PCR cycle to ensure that any remaining single-stranded DNA is fully extended.

Notes:

- The final MgCl₂ concentration of this 5x Master Mix is 2 mM. In some applications, more than 2 mM MgCl₂ is required for best results. Use 25 mM to adjust the Mg²⁺ concentration according to table 3.

Table 3. Additional volume (µl) of MgCl₂ per 25 µl reaction:

Final MgCl ₂ conc. in reaction (mM)	2.0	2.5	3.0	3.5	4.0	4.5
Volume of 25 mM MgCl ₂	0	0.5	1.0	1.5	2.0	2.5

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Other product sizes, combinations and customized solutions are available. Please look at www.dutscher.com or ask for our complete product list for PCR Enzymes. For customized solutions please contact us.

Made in Europe

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