

HelixAmp™ Ready-2x-Go [*Premium-Taq*]

Kit Contents

HelixAmp™ Ready-2x-Go [<i>Premium-Taq</i>]		
Cat. No.	PM007L	PMD007L
Packing size	1ml x 5ea	1ml x 5ea
Ready-2x-Go [<i>Premium-Taq</i>]	without dye	with dye
N-Solution™	1ml	1ml
Instructions for Use	1ea	1ea

HelixAmp™ Ready-2x-Go [*Premium-Taq*] are the mixtures of HelixAmp™ *Premium-Taq* Polymerase, PCR buffer, dNTPs and stabilizing agents. For the optimization of difficult PCR, N-Solution™ is separately provided.

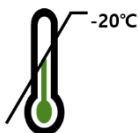
Description

HelixAmp™ Ready-2x-Go [*Premium-Taq*] are optimized mixtures of HelixAmp™ *Premium-Taq* Polymerase with reaction buffer and dNTPs as 2-fold concentration. This pre-mixed formulation is designed to save time and reduce the error and contamination opportunities. HelixAmp™ Ready-2x-Go [*Premium-Taq*] mixture contains NanoHelix's *Premium-Taq* Polymerase, which is an anti-*Taq* antibody complex form and ideal for automatic hot-start PCR. HelixAmp™ Ready-2x-Go [*Premium-Taq*] provides the most suitable condition for efficient and reproducible PCR.

Store

Store the products containing dye below -20°C and keep away from light during storage.

Storage



Store below -20°C

Shelf life



12 months

Quality control

By NanoHelix's ISO 13485-certified Quality Management System, each lot of **HelixAmp™ Ready-2x-Go [*Premium-Taq*]** was tested against predetermined specifications to ensure consistent product quality.

Protocol

1. Recommended amount of template DNA.

Human genomic DNA : 10 ~ 100ng

Bacterial genomic DNA : 5 ~ 50ng

Purified plasmid or phage DNA : 1 ~ 5ng

2. Mix following components in a PCR tube.

Components	Volumes (μ l)
Template	X μ l
Forward Primer (10 μ M)	2 μ l
Reverse Primer (10 μ M)	2 μ l
N-Solution™ [optional]	0 ~ 5 μ l
Ready-2x-Go [<i>Premium-Taq</i>]	25 μ l
RNase-free Water	to 50 μ l

※ **N-Solution™** is an additive altering the binding behavior of primer and template and can help the amplification that do not work well under standard PCR condition. Especially, **N-Solution™** can be used for the amplification of problematic template, such as high G+C content and repeat sequence regions. The optimal concentrations of **N-Solution™** are vary upon the primer-template sets and should be set by adding into the PCR reaction mixture from 2 to 10% volume. Most of the PCR reactions are not required the **N-Solution™** and we recommend to use the **N-Solution™** only in case of the PCR amplification is not works well or too much non-specific products are observed.

3. PCR condition.

Temperature & time	Cycles
95°C, 2 min	x 1
95°C, 20 sec	} x 25 ~ 40
Annealing Temp., 40 sec	
72°C, 1 min/kb (Expected size of product)	
72°C, 5 min	x 1

Annealing Temp. = $T_m - (4 \sim 6^\circ\text{C})$

T_m (Melting Temp.) = $[4^\circ\text{C} \times (\text{G} + \text{C})] + [2^\circ\text{C} \times (\text{A} + \text{T})]$

Precautions

Store the product containing dye in a place protected from light, as prolonged exposure to light may degrade its performance.

Products

Cat. No.	Products	Size
PM007L	HelixAmp™ Ready-2x-Go [<i>Premium-Taq</i>], N-Solution™	1ml x 5ea
PMD007L	HelixAmp™ Ready-2x-Go [<i>Premium-Taq</i>] (with dye), N-Solution™	1ml x 5ea