

Cat No. SM213-0250

Size: 250 Reactions (2 X 1.25 ml)

Cat No. SM213-0010

Size: 10 Reactions (1X 100 µl)

Storage: Store at RT up to 3 months

Store at 4°C up to 6 months

Store at -20°C up to 1 year

Shipping Temperature: 4°C

Note: amaR OnePCR is light sensitive and should be stored and protected from light.

## Description

The amaR OnePCR is a ready-to-use PCR reaction mixture. Simply add primers, template, and water, the reagent will execute primer extensions and other molecular biology applications. The amaR OnePCR is a pre-mixed solution containing *Taq* DNA polymerase, PCR buffer, dNTPs, gel loading dyes, and fluorescence dye. The amaR OnePCR which contains the *Taq* DNA polymerase, is purified from the *E. coli.*, and expressing the *Thermus aquaticus* DNA polymerase gene. This enzyme has a 5' → 3' DNA polymerase and the 5' → 3' exonuclease activity but lacks the 3' → 5' exonuclease activity. The amaR OnePCR, which contains the fluorescence dye, is directly detected on BLook LED transilluminator or UV epi-illuminator after the DNA electrophoresis. The amaR OnePCR contents red tracking dyes, provide a safe, non-toxic and non-mutagenic alternative to ethidium bromide for instantaneous band visualization, that are environmentally friendly containing no hazardous chemicals. The tracking dyes that run at 10 bp on a 1% agarose gel. The amaR OnePCR mixture is supplied at the 2X concentration to allow approximately 50% of the final reaction volume to be used for the addition of primer and template solutions. The reagents are provided with the sufficient amplification reactions of 20 µl each.

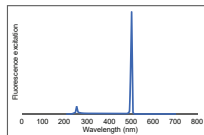


Fig. 1a. Fluorescence excitation spectra of the fluorescence dye

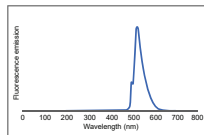


Fig. 1b. Fluorescence emission spectra of the fluorescence dye

## Features

- No post-staining processing of DNA required.
- No need to prepare PCR Reagents.
- Direct loading onto your agarose gel for analysis.
- Sensitivity – High degree of sensitivity as the ethium bromide.
- Speed – No destaining requirement.
- Compatibility – Use the blue light or UV to detect the signal.

## Tracking dye

- Amaranth

## Protocol

Standard PCR with amaR OnePCR :

1. For each 20 µl reaction, assemble the following components in a 0.2 ml PCR tube on ice just prior to use:

	Volume (µl)	Final Concentration
amaR OnePCR	10	1X
Forward primer, 5~10 µM	Variable	0.1-0.2 µM
Reverse primer, 5~10 µM	Variable	0.1-0.2 µM
DNA template	Variable	4 pg~500 ng
Add ddH <sub>2</sub> O to	20	

2. Mix gently. If necessary, centrifuge briefly. Cap tubes and place in the thermal cycler.

3. Process in the thermal cycler for 25~35 cycles as follows:

Initial Denaturation	2~5 minutes at 94°C	} 30 cycles
Denaturation	20~40 seconds at 94°C	
Annealing	1 minute at the proper annealing temperature	
Extension	2 minutes at 72°C	
Final Extension	5 minutes at 72°C	

Note: Optimal conditions for amplification will vary depending on the primers and thermal cycler used. It may be necessary to optimize the system for individual primers, template, and thermal cycler.

4. After the PCR reaction, DNA electrophoresis will detect the PCR product.

5. Use the UV or blue-light transilluminator or UV epi-illuminator to photograph the gel.

Note: When the DNA concentration is less than 4 pg, it may cause the migratory shift when performing the electrophoresis. To remedy this observation, we recommend to conduct the following steps (please refer to the experimental procedures), or use the PCR Clean-Up & Gel Extraction Kit (see SN006-0100) to remove the fluorescence dye prior to post-staining with the Novel Green (SL002-0500) or Novel Green *plus* (SL003-0500) again for restoring the DNA molecular weight in the original position.

### Removal of Fluorescence Dye

1. Immerse the PCR product containing the fluorescence dye into the 100 mM NaCl and add 2.5 volumes of absolute or 95% ethanol.
2. Incubate on ice for 20 minutes.
3. Centrifuge the mixture at 4°C for at least 10 minutes.
4. Remove the suspension of ethanol and wash the pellet with 1ml of 70% ethanol.
5. Dry the residual ethanol and resuspend the double-stranded DNA in the TE.

### Related Ordering Information

Cat. No.	Description	Package
BK001	BLoK LED transilluminator	1 each
SL001-1000	Novel Juice	1 ml
SL003-0500	Novel Green Plus	500 µl (20,000X in DMSO)
SM215-0250	amaR OnePCR HiFi	250 Reactions
SM216-0250	amaR OnePCR HotStar	250 Reactions
SD101-0100	OneMARK 100	600 µl
SD110-0100	OneMARK B	600 µl

#### Caution:

1. During operation, always wear a lab coat, disposable gloves, and protective equipment.
2. Research Use Only. Not intended for any animal or human therapeutic or diagnostic uses.