## Blue MegaMix-Double

Double concentrated PCR mix
Containing Taq polymerase (recombinant) in $2 x$ reaction buffer ( $5 \mathbf{~ m M ~ M g C l} 2$ ) with $400 \boldsymbol{\mu}$ dNTPs, blue agarose loading dye \& stabiliser

## For instant and accurate PCRs

## Protocol

Thaw the Blue MegaMix-Double
Add the desired volume ( $25 \mu \mathrm{l}$ or less) into the PCR tube*
Add DNA, primers and water of total equal volume to Blue MegaMix-Double
Overlay with mineral oil if necessary
Place in a Thermal Cycler

Cycling profile (guide only)
Initial denaturation step: $95^{\circ} \mathrm{C}$ for 3 mins
Then cycle 25-30 times:
Step 1: $95^{\circ} \mathrm{C}$ for 30 secs
Step 2: Optimal annealing temp. of primers for 30 to 60 secs
Step 3: $72^{\circ} \mathrm{C}$ for 45 to 60 secs


Cool to room temperature

After PCR, samples can be loaded direct onto the agarose gel
NO GEL LOADING BUFFER REQUIRED
*Refreeze the rest of the Blue MegaMix-Double. It can be frozen/thawed many times without loss of activity

## Related Products

100 bp DNA ladder = An agarose size standard supplied 'ready to load'
in room temperature stable loading buffer microCLEAN = The easy way to clean-up your PCR fragments

Just Water = Molecular grade water in handy sizes

