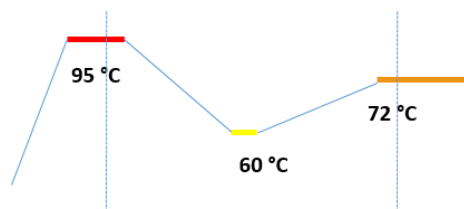




Fast PCR protocol:

All Ampliqon Taq DNA Polymerases and Taq master mixes

90 min



3-step Standard PCR protocol

PCR program for 3-step Standard PCR – 90 min total

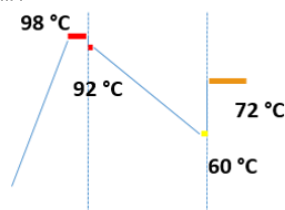
Cycler step	Temperature	Duration	Cycles
Initial heating	95 °C	3 min.	1
Denaturation	95 °C	30 sec.	30
Annealing*	60 °C	30 sec.	
Extension	72 °C	30 sec.	1
Final extension	72 °C	5 min.	

* the annealing temperature depends on the primer set

SAVE 1 HOUR

Save time - just by changing PCR cycler settings!

31 min

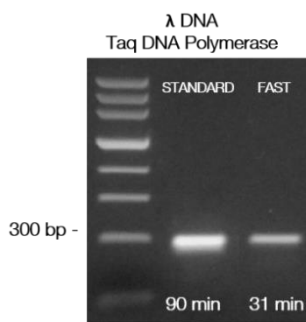


2-step Fast PCR protocol

PCR program for 2-step Fast PCR – 31 min total

Cycler step	Temperature	Duration	Cycles
Initial heating	98 °C	40 sec.	1
Denaturation	92 °C	2 sec.	30
Extension*	60 °C	2 sec.	
Final extension	72 °C	20 sec.	1

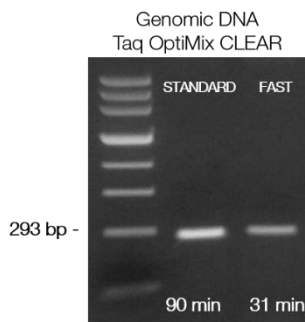
* the extension temperature depends on the primer set. For fast PCR choose highest possible T_m values



Amplification of λ DNA using Taq DNA Polymerase – Experimental setup

Reaction mix*	ID	Primer sequence (5'-3')	Length
Ammonium buffer			
1x	LAM300-F	ACGGATAGAAACTGCCGGTCAGGACA	300 bp
dNTP mix	LAM300-R	GTTATCGAAATCAGCCACAGGGC	
0,2 mM each			
MgCl ₂			
1,5 mM			
Primers			
0,2 μ M			
λ DNA			
1 ng			
Taq DNA polymerase			
0,5 – 1U			

* H₂O up to a total volume of 25 μ l



Amplification of gDNA using 2x Taq OptiMix CLEAR - Experimental setup

Reaction mix*	ID	Primer sequence (5'-3')	Length
Taq OptiMix			
1x	ENG9-F	AATGGCTGTGACTTGGGACCCCTG	293 bp
Primers	ENG9-R	GCACCAACCAGGCTGGTCTGATA	
0,2 μ M			
gDNA			
20 ng			

* H₂O up to a total volume of 25 μ l

Please require our Application note for Fast PCR: "Additional reduction of PCR run time – three approaches"