## onTaq DNA Polymerase HOT START



experiments with quality

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onTaq DNA Polymerase is a chemically modified recombinant enzyme. onTaq DNA Polymerase provides very tight inhibition of the polymerase activity at moderate temperatures allowing room temperature reaction setup. The polymerase activity is restored during 5-minute initial denaturation step.

Use of the onTaq DNA Polymerase allows for the increase of PCR specificity, sensitivity and yield in comparison to the conventional PCR assembly method.

Both increased specificity and reduced mispriming improve multiplex PCR.

**Fig. 1** PCR amplification using EURx onTaq DNA Polymerase. 2 kb amplicon of the human β-globin gene was amplified using EURx onTaq DNA Polymerase,  $10 \times Pol$  Buffer B and 0.2 mM dNTPs in  $50 \, \mu l$  reaction volume.

**Lane 1**: molecular size marker Perfect Plus 1 kb DNA Ladder (E3113).

**Lane 2**: PCR amplification reactions using 1.25 U Taq DNA Polymerase. Reactions were incubated 30 min at 25 °C before PCR.

**Lanes 3, 4**: PCR amplification reactions using 1.25 U onTaq DNA Polymerase. Reactions were incubated 30 min at 25 °C before PCR.





**Fig.2** PCR amplification using EURx onTaq DNA Polymerase. 2 kb amplicon of the human β-globin gene was amplified using EURx onTaq DNA Polymerase in  $50 \, \text{ml}$  reaction volume.

Lane 1: molecular size marker Perfect Plus 1 kb DNA Ladder (E3113).

**Lane 2**: PCR amplification reaction using 1.25 U Taq DNA Polymerase. Reaction was incubated 30 min at 25 °C before PCR.

**Lanes 3, 4:** PCR amplification reactions using 1.25 U onTaq DNA Polymerase stored 3 months at  $-20^{\circ}$ C. Reactions were incubated 30 min at  $25^{\circ}$ C before PCR.

**Lanes 5, 6:** PCR amplification reactions using 1.25 U onTaq DNA Polymerase stored 3 months at  $4^{\circ}$ C. Reactions were incubated 30 min at  $25^{\circ}$ C before PCR.