Super-Run Taq DNA Polymerase

Date 2009.14 Version 1_03

Cat No : PT-T535-250U **Size :** 250 units

Concentration : 5 units/μl Volume : 50 μl

Purity degree:

Nicking activity, endonuclease and exonuclease activity were not detected after the incubation of 0.6 μg of supercoiled pBR322 DNA, 0.6 μg of λ DNA or 0.6 μg of λ -Hind III digest with 10 units of this enzyme for 1 hour at 74°C.

PCR products:

As most PCR products amplified with Super-Run Taq that is high fidelity thermostable DNA polymerases, have one A to add at 3'-termini, the obtained PCR product can be directly used for cloning into T-vector. Also it is possible to clone the product in blunt-end vectors after blunting and phosphorylation of the end.

Components: Store at -20°C.

- 1 tubes of 5 μl Super-Run Taq DNA polymerase
- 1 tube of 100 ul 10X Super-Run Taq Buffer (20 mM Mg²⁺ plus)

General Protocol: (total 50 ul)

Set up as follows:	Experimental Sample

Super-Run Taq ™ (5 units/μl)	0.25 ul
10X Super-Run Taq Buffer (20 mM Mg ²⁺ plus)	5 ul
dNTP Mixture (2.5mM)	4 μl (1ul for 10 mM dNTP)
Primer 1 (0.2 ~ 1.0 uM)	Xμl
Primer 1 (0.2 ~ 1.0 uM)	Yμl
Template < 500 ng	Zμl
Sterilized distilled water	Up to 50 ul
Total	50 μl

- For Experimental Sample, go to your own Cycle Conditions.
- When amplifying 1 kb DNA fragment: 30 cycles of 98 °C denaturation for 10 sec, 55 °C annealing for 30 sec and 72°C extension for 1 mins, Or, 30 cycles of 98 °C denaturation for 10 sec, 68°C annealing/extension for 1 min. Denatureation condition varies depending on an used thermal cycler and tube. It is recommended for 10-30 sec at 94°C, or 1-10 sec at 98°C

For Research Using Only. Please do not hesitate to contact us if you have any questions.

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