

## *TransStart*<sup>®</sup> *FastPfu* Fly DNA Polymerase

Please read the datasheet carefully prior to use

Cat. No. AP231

Concentration: 2.5 units/μl

Storage: at -20°C for two years

### Description

*TransStart*<sup>®</sup> *FastPfu* Fly DNA Polymerase is a hot start, high-fidelity and high processivity DNA Polymerase used for fast PCR. Compared with *TransStart*<sup>®</sup> *FastPfu* DNA Polymerase, *TransStart*<sup>®</sup> *FastPfu* Fly DNA Polymerase has higher extension rate (6kb/min vs 4kb/min), higher fidelity, and higher amplification efficiency.

- Offers 108-fold fidelity as compared to *EasyTaq*<sup>®</sup> DNA Polymerase.
- PCR products can be directly cloned into *pEASY*<sup>®</sup>-Blunt vectors.
- Amplification of genomic DNA fragment up to 15 kb.
- Amplification of plasmid DNA fragment up to 20 kb.

### Features

- Hot start, high specificity.
- High amplification efficiency.
- Fast and high fidelity.
- High sensitivity.
- High production.

### Applications

- Amplifies complex and high GC/ AT templates.
- High fidelity and fast PCR, blunt end cloning, site-directed mutagenesis.
- Amplifies long fragment.

### Kit Contents

Component	AP231-01/11	AP231-02/12	AP231-03/13
<i>TransStart</i> <sup>®</sup> <i>FastPfu</i> Fly DNA Polymerase	250 U×1	500 U×1	500 U×6
5× <i>TransStart</i> <sup>®</sup> <i>FastPfu</i> Fly Buffer	1.2 ml ×1	1.2 ml ×2	1.2 ml ×12
50 mM MgSO <sub>4</sub>	200 μl×1	400 μl×1	1 ml×1
6×DNA Loading Buffer	500 μl×1	1 ml×1	1 ml×2
2.5 mM dNTPs	- /500 μl×1	- / 1 ml×1	- / 1 ml×6
Complimentary Component	200 μl×1	400 μl×1	1 ml×1
PCR Stimulant			



#### Reaction Components(50 $\mu$ l reaction volumes)

Component	Volume	Final Concentration
Template	Variable	As required
Forward Primer (10 $\mu$ M)	1 $\mu$ l	0.2 $\mu$ M
Reverse Primer (10 $\mu$ M)	1 $\mu$ l	0.2 $\mu$ M
5 $\times$ TransStart <sup>®</sup> FastPfu Fly Buffer	10 $\mu$ l	1 $\times$
2.5 mM dNTPs	4 $\mu$ l	0.2 mM
TransStart <sup>®</sup> FastPfu Fly DNA Polymerase	1 $\mu$ l	2.5 units
Nuclease-free Water	Variable	-
Total volume	50 $\mu$ l	-

#### Optimized parameters (50 $\mu$ l reaction volumes)

Template	Input
Genomic DNA	10-500 ng
Plasmid DNA	1-30 ng
cDNA	1-2 $\mu$ l cDNA from RT reaction (50-500 ng RNA for RT reaction)

#### PCR

Number of Cycles	Temperature	Time
1 cycle	95°C	2 min
30-35 cycles	95°C	20 sec
	T <sub>m</sub> -5°C	20 sec
	72°C	6 kb/min
1 cycle	72°C	5 min

#### PCR Stimulant

PCR Stimulant is used to optimize the amplification of complex templates or high GC/ AT templates. The amplification of the Pfu series of enzymes is enhanced significantly. The concentration of the storage solution is 5 $\times$ , and the concentration of the working solution can be adjusted between 0.5 $\times$ -2.5 $\times$

#### Notes

- For GC-rich templates, the recommended denaturation temperature is 98°C
- To ensure high fidelity, we recommend using high quality dNTPs. dNTPs containing dUTP cannot be used.
- It is recommended to add TransStart<sup>®</sup> FastPfu Fly DNA Polymerase to the reaction system in the last step.
- If 5 $\times$ TransStart<sup>®</sup> FastPfu Fly Buffer has a small amount of precipitation after thawing, please heat it in a 37°C water bath and mix it for use.

FOR RESEARCH USE ONLY

