

# AccuGO™ Pfu proofreading DNA polymerase

Catalog number: C108300  
Store at -20°C

## Materials supplied:

1. AccuGO™ Pfu DNA polymerases 100U (2 units/μL) in storage buffer. (Cat.: C108301)
2. 5X AccuGO™ Pfu buffer 1.25mL (1 mL/vial) provides a final 2 mM Mg<sup>2+</sup> concentration in the PCR reaction. (Cat.: C108302)

## I. Description:

AccuGO™ Pfu DNA polymerase is a Pfu-based DNA polymerase specially provides both short extension time and high performance. It was designed to apply in the replication of DNA sequence required high fidelity and high efficiency. Like Pfu, AccuGO™ Pfu also exhibits 3'→5' exonuclease activity that provides the proofreading ability and results in higher fidelity during DNA synthesis. Moreover, AccuGO™ Pfu was engineered to have a better processivity than normal Pfu. Therefore, AccuGO™ Pfu can easily amplify up to 10 kb blunt-ended DNA fragment from variant template. Even if the template's GC content is up to 70%, AccuGO™ Pfu still has good performance. In short, AccuGO™ Pfu DNA polymerase is an ideal tool for molecular cloning containing long or complicated amplicons.

## II. Applications:

1. High fidelity PCR.
2. Routine PCR amplification of DNA fragments up to 10 kb.
3. PCR products for cloning and expression.

## III. Storage Condition

| -20°C     | 4°C      | 25°C    | 37°C    |
|-----------|----------|---------|---------|
| 1.5 years | 2 months | 21 days | 14 days |

## IV. Recommended PCR Condition:

| DNA         | Amount      |
|-------------|-------------|
| Genomic DNA | 10-250 ng   |
| Plasmid DNA | 1 pg -10 ng |
| cDNA        | 1 -100 ng   |

| Component                  | Amount per reaction |
|----------------------------|---------------------|
| DNA template               | Variable            |
| 10 μM forward Primer       | 1 μL                |
| 10 μM reverse Primer       | 1 μL                |
| 10 mM each dNTP            | 1 μL                |
| 5X AccuGO™ Pfu buffer      | 10 μL               |
| AccuGO™ Pfu DNA polymerase | 0.5 μL              |
| Nuclease-free water        | to 50 μL            |
| Total volume               | 50 μL               |

## V. Thermocycling Conditions for a Routine PCR:

The recommended parameters may be optimized for each new primer-template pair for optimal specificity and amplification.

| Cycles | Step                 | Temperature | Time       |
|--------|----------------------|-------------|------------|
| 1      | Initial denaturation | 95°C        | 1-5 mins   |
| 25-35  | Denaturation         | 95°C        | 10-30 s    |
|        | Annealing            | 45-72°C     | 10-30 s    |
|        | Extension            | 72°C        | 15-30 s/kb |
| 1      | Final Extension      | 72°C        | 5-10 mins  |
|        | Hold                 | 14°C        | --         |

## VI. Experimental Data

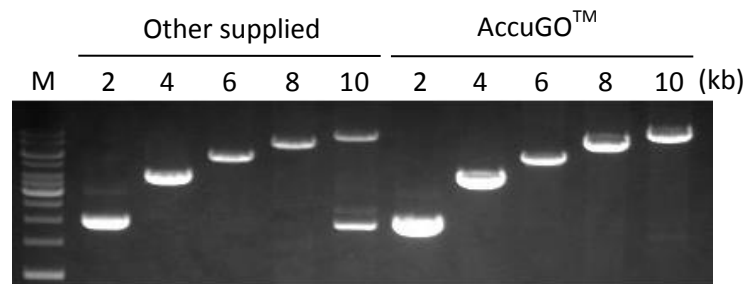


Figure 1. Comparison of PCR amplification performance between AccuGO™ and other suppliers' Pfu DNA polymerase.

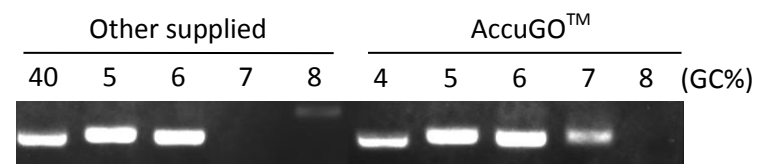


Figure 2. Comparison of the tolerance of GC content between AccuGO™ and other suppliers' Pfu DNA polymerase.

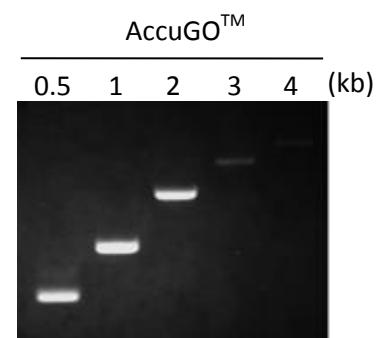


Figure 3. Comparison of the different PCR amplifications at extension time 30 (sec) of AccuGO™.