

KAPA2G Robust DNA Polymerase

KAPA2G Robust DNA Polymerase is a highly robust and versatile second-generation (2G) enzyme derived through a process of molecular evolution. The novel amino acid mutations in the polymerase offer higher processivity and specific activity, which translates to robust performance across a wide range of GC- and AT-rich templates, higher yields and sensitivity, and improved tolerance to common PCR inhibitors.

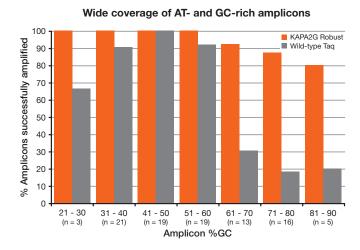
KAPA2G Robust DNA Polymerase is ideally suited for:

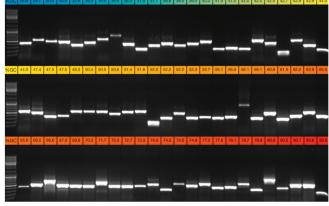
- · Consolidating PCR protocols and reagents
- Broad coverage of both AT- and GC-rich targets
- · Applications requiring higher yield per unit enzyme
- Colony PCR
- Tolerance to inhibitor carryover and crude sample PCR (e.g. FFPE)
- Routine PCR, using the HotStart or ReadyMix formulation

A second-generation DNA polymerase evolved for robustness and versatility.

Consolidate PCR protocols and increase success rates with a single enzyme.

The improved processivity and tolerance to common PCR inhibitors of the KAPA2G Robust DNA Polymerase offers consistent amplification, high yields and wide coverage of both easy and challenging amplicons. The unique features of the enzyme supports versatile and robust amplification of a broad range of AT- and GC-rich targets and allows for the simplification of PCR workflows, through the consolidation of reagents and protocols, while increasing success rates and turnaround time.

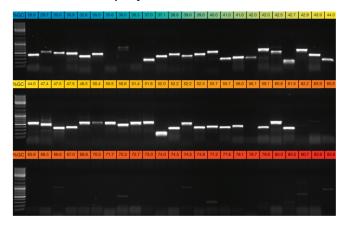




A total of 96 amplicons from human genomic DNA were amplified using a single PCR protocol (above): 3 min (95 °C) initial denaturation followed by 35 cycles of 15 sec (95 °C) denaturation, 15 sec (60 °C) annealing, and 15 sec (72 °C) extension for KAPA2G Robust HotStart ReadyMix or 60 sec (72 °C) extension for Taq polymerase. With KAPA2G Robust, high success rates were achieved across the full spectrum of GC contents, whereas wild-type Taq yielded poor results with AT-rich amplicons and amplicons with GC content >60%.

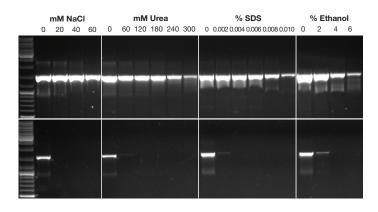
72 of the 96 amplicons (right), representing a range of primer lengths, sequence composition and melting temperatures, were electrophoresed in order of increasing GC content, with the lowest GC content (27%, blue) at the top left corner and the highest GC content (84%, red) at the bottom right corner of each composite gel image. All reactions contained 25 ng human genomic DNA.

Hot start Taq Polymerase achieved 66% success



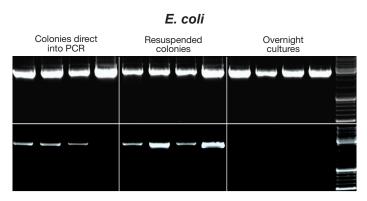
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Greatly improved tolerance to common PCR inhibitors.

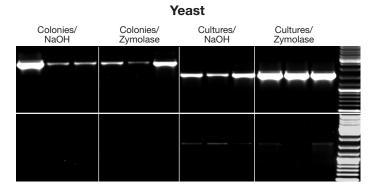


Amplification of a 1.5 kb fragment from 1 pg plasmid DNA in the presence of four common PCR inhibitors using KAPA2G Robust HotStart (top) or wild-type Taq (bottom). All reactions contained 0.5 units of enzyme per 25 μ l reaction. A standard 3-step cycling profile (35 cycles) with an annealing temperature of 65 °C and 1.5 min extension per cycle was used.

Unrivalled performance in Colony PCR for yeast and E. coli



Amplification of a cloned 2.7 kb insert from four commonly used $E.\ coli$ strains (DH5a, DH10B, JM109 or BL21) using KAPA2G Robust HotStart (top) or wild-type Taq (bottom). Colonies (grown on LB-agar + Amp plates) were either resuspended directly in individual PCR reactions (left) or first resuspended in PCR grade water and then added to PCR reaction mixes (middle). For overnight cultures (prepared in LB + Amp), 1 μ l was added directly to the PCR mix (right).



Amplification of a 2.5 kb (left) or 1.6 kb (right) fragment from the GSH1 gene from three commonly used *S. cerevisiae* strains (BY4742, FY23 and W303) using KAPA2G Robust HotStart (top) or wild-type Taq (bottom). Colonies (from YPD-agar plates) or YPD overnight cultures were first lysed in 50 μ l volumes with NaOH or Zymolase (as indicated).



ORDERING INFORMATION

Code	Kit contents
KK5023	100 units
KK5024	250 units
KK5004	100 units
KK5005	250 units
KK5522	100 units
KK5515	250 units
KK5517	500 units
KK5525	2500 units
KK5532	100 units
KK5516	250 units
KK5518	500 units
KK5701	100 rxns
KK5702	500 rxns
	KK5023 KK5024 KK5004 KK5005 KK5522 KK5515 KK5517 KK5525 KK5532 KK5516 KK5518