**PCRBIO Taq DNA Polymerase**

- Robust
- Reliable
- Convenient

PCRBIO Taq DNA Polymerase employs our advanced polymerase technology and buffer chemistry to enhance PCR speed, yield and specificity. The enzyme and buffer system allow for superior PCR performance on complex templates such as mammalian genomic DNA. Taq DNA polymerase is your starting point for every basic PCR application.

**Features**

- Increased PCR success rates with amplicons up to 6kb
- Ultra low background DNA
- Advanced buffer chemistry including Mg and dNTPs
- High yields under standard and fast PCR conditions
- Efficient and specific amplification from complex templates including GC and AT-rich sequences
- Also available as a 2x ready mix, with the option of a red dye for direct gel loading

**Applications**

- Routine PCR applications
- TA cloning
- High throughput PCR
- Methylated DNA amplification
- Crude sample PCR
- Standard and fast PCR

**Figure 1.** PCRBIO Taq DNA Polymerase outperforms most competitors at amplifying a 1kb fragment

PCR amplification of a 1kb fragment of the GAPDH gene using a 1 in 10 serial dilution of mouse genomic DNA (5000, 500, 50 and 5pg) with PCRBIO Taq DNA Polymerase (purple) and matching Taq polymerases from NEB (orange), Promega (yellow) and Thermo (red). Reactions were set up using master mix formats and following manufacturers’ recommendations. Cycling conditions were 95°C 2min, then 40 cycles of 95°C 15sec, 63°C 15sec, 72°C 30sec. 1/5 of the reaction volume was loaded in 1% agarose gel. L: PCRBIO Ladder II. PCRBIO Taq DNA Polymerase outperforms NEB’s OneTaq and Promega’s GoTaq G2, and is similar to Thermo’s DreamTaq.
PCRBIO Taq DNA Polymerase is a robust enzyme for all your everyday PCR applications, including genotyping, screening and library construction. PCRBIO Taq DNA Polymerase performs consistently well on a broad range of templates including both GC and AT-rich.

PCRBIO Taq DNA Polymerase has 5′-3′ exonuclease activities, but no 3′-5′ exonuclease (proofreading) activity. The enzyme has the same error rate as wild-type Taq DNA polymerase, approximately 1 error per $2.0 \times 10^5$ nucleotides incorporated. PCR products generated are A-tailed and may be cloned into TA cloning vectors.

PCRBIO Taq DNA Polymerase provides the research community with an affordable routine application polymerase that performs to the highest possible standard, with a versatility that allows you to amplify with the highest speed, yield, specificity and consistency on the market. The enzyme is produced using an enhanced 12 step purification strategy which includes physical, chemical and enzymatic removal of host DNA.

For added convenience PCRBIO Taq DNA Polymerase is also available as a 2x ready mix, with the option of a red dye suitable for direct loading and tracking during agarose gel electrophoresis.

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**Table:**

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<thead>
<tr>
<th>Catalogue Number</th>
<th>Product Name</th>
<th>Pack Size</th>
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</table>

PCR amplification of a 1kb fragment of the GAPDH gene using a 1 in 10 serial dilution of mouse genomic DNA (5000, 500, 50 and 5pg), with PCRBIO Taq Mix Red and matching Taq mixes from NEB (orange), Promega (yellow), Thermo (red) and Kapa Biosystems (blue). Reactions were set up using manufacturers’ recommendations. Cycling conditions were 95°C 2 min, then 40 cycles of 95°C 15 sec, 63°C 15 sec, 72°C 30 sec except for NEB: 94°C 2 min, then 40 cycles of 94°C 15 sec, 63°C 15 sec, 68°C 30 sec. L: PCRBIO Ladder II.