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# 🌐 General Taq PCR Master Mix -- CHEM 384/584 V.3

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2X PCR Master Mixes are convenient to use as they include all the necessary PCR components except the template and primers. Most PCR Master Mixes also include agarose gel running dyes and a density reagent that allows direct loading of PCR products on an agarose gel for electrophoresis. The master mix format simplifies workflows and sample handling; simply add primers, template, and water and then begin PCR.

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**protocols.io**<https://dx.doi.org/10.17504/protocols.io.b3npqmdn>

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**Storage:** -20°C for long-term storage. 4°C for short-term storage (up to 3 months).  
(Note) If used frequently, store at 4°C ; the activity of the Master Mix may decrease with repeated freezing and thawing. Gently mix well before use and centrifuge briefly.

**Application:**

- DNA amplification by PCR
- Colony PCR

**PCR Products:** Since most PCR products amplified Taq PCR Master Mix have an A overhang added at 3'-termini, the obtained PCR product can be used directly for cloning into a TA cloning vector. Additionally, it is possible to clone the product in a blunt-end vector after blunting and phosphorylation.

**Dye Migration during Electrophoresis:** Check individual manufacturers information on the migration of the included running dyes.

Setup Reaction

- 1 To a **25 µL** aliquot of a 2X Taq PCR Master Mix (e.g. TaqDog, or Sapphire Amp), add template (10-20 µl cleared lysate for colony PCR or **20-50 ng** of purified DNA for typical PCR), forward and reverse primers to a final concentration of **200 nanomolar (nM)** . Adjust final volume to **50 µL** with nuclease free water or autoclaved water.

A	B
2X Master Mix	25 ul (pre-aliquoted and stored in the freezer)
Template	10-20 ul of bacterial lysate or 20-50 ng purified DNA
Forward Primer	1 ul of 10 uM primer dilution
Reverse Primer	1 ul of 10 uM primer dilution
ddH2O	to a final volume of 50 ul

Run Reaction

- 2 followed by 30 cycles of 98°C, 5 sec; 55°C, 5 sec; and 72°C, 40 sec.

A	B	C
Initial denature	94C	1 minute
Denature	94C	30 seconds
Anneal	55C	30 seconds
Extension	72C	1 min/kb
Repeat steps 2-4		25-40x
Final extension	72	minutes
Cool	4C	Until cancelled

A typical thermocycling program for a PCR for amplicons less than 1 kb is 1 minute. For longer amplicons, adjust the program to 1 minute/kb seconds for the extension and final extension times.