An Introduction to Polymerase Chain Reaction (PCR)

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Introduction

- The technique was invented by Dr. Kary Mullis, 1986
- for which he received the Nobel Prize in Chemistry in 1993.

PCR Achieves Fame and Fortune

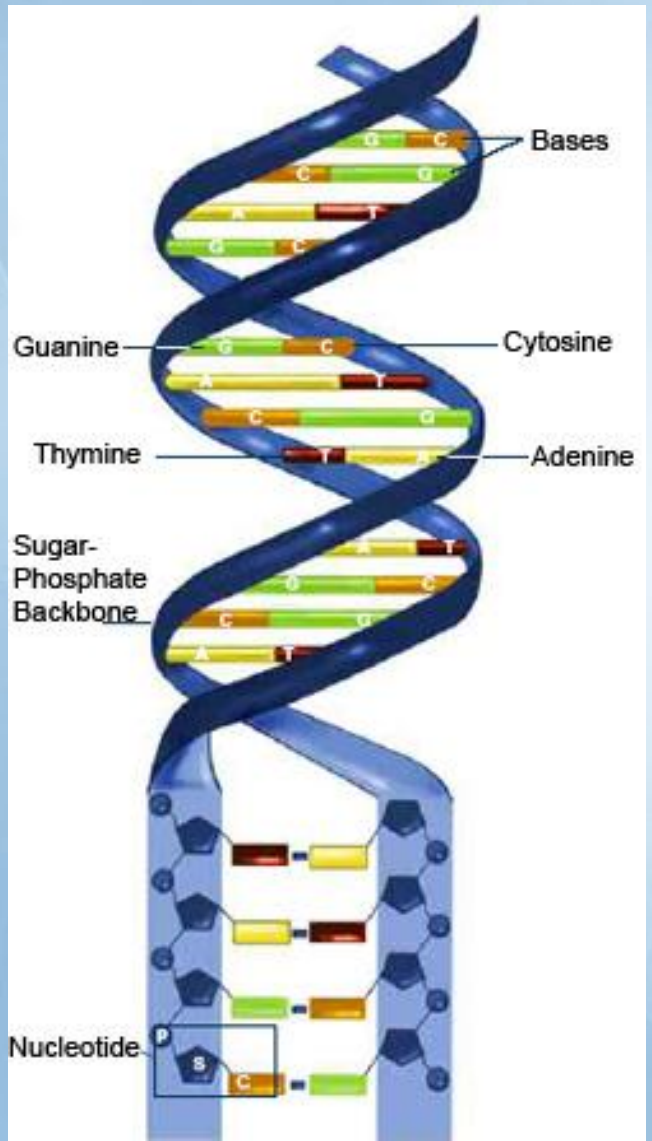
--becomes standard in molecular biology tool box--

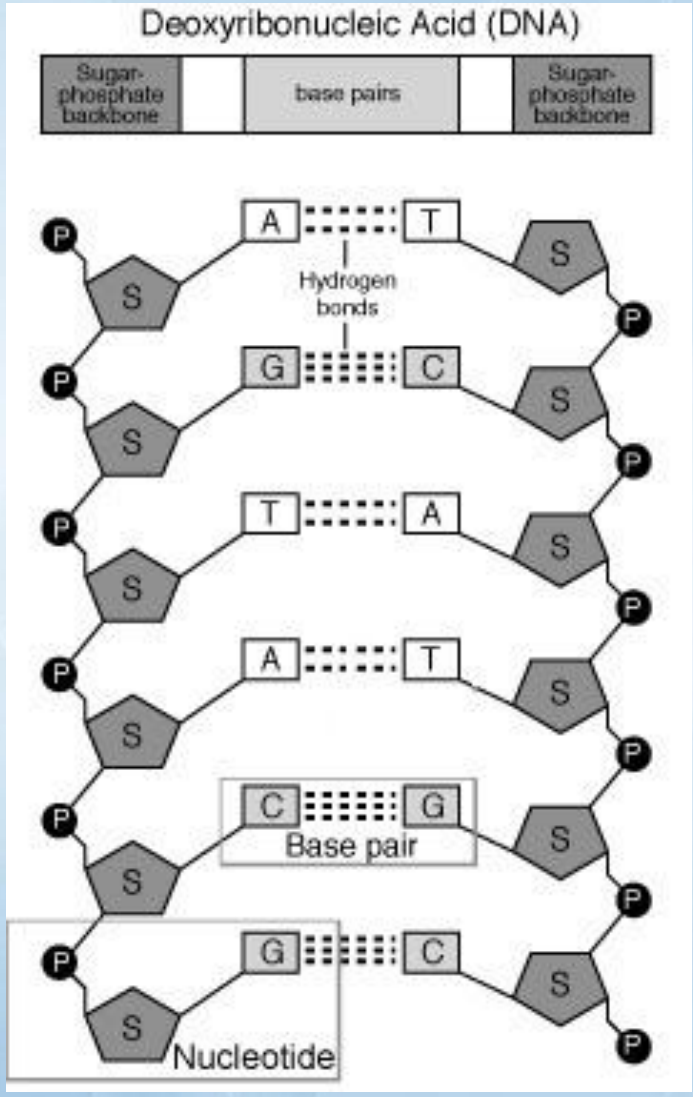
The Molecule of the Year

Ruth Levy Guyer and Daniel E. Koshland, Jr.

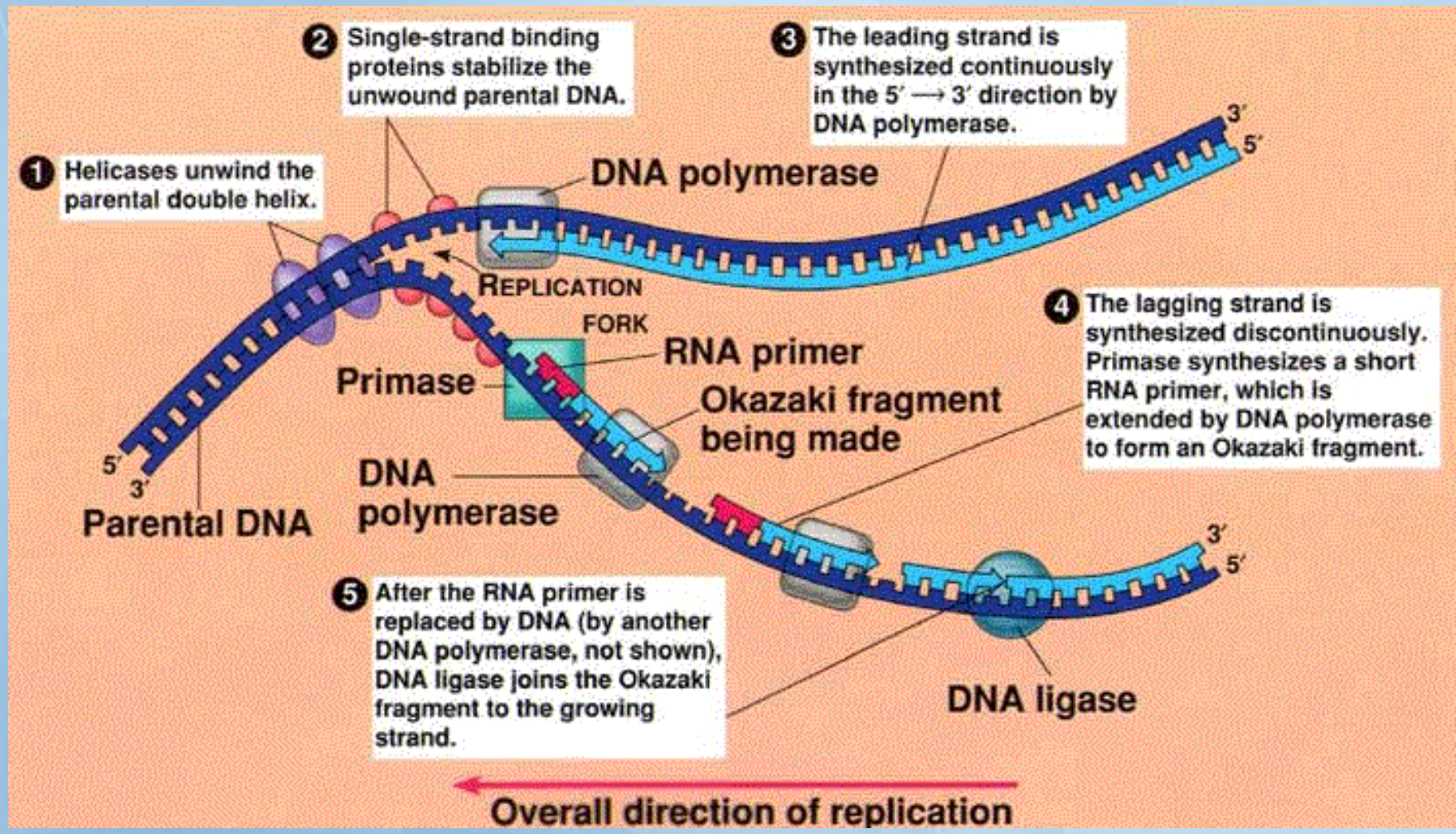
Science has selected the polymerase chain reaction as the major scientific development of 1989 and has chosen for its first "Molecule of the Year" the DNA polymerase molecule that drives the reaction. The list from which the polymerase chain reaction (PCR) was chosen included an impressive array of accomplishments in many areas of science and technology; additional kudos are therefore conferred below to 17 of the other big "stories" that made 1989 an exciting year for scientists and for followers and beneficiaries of science. Although the PCR procedure was introduced several years ago, use of the technique truly burgeoned in 1989; in much the same way, the full potentials of many of the interesting "runner-up" scientific achievements of this year are likely to be realized sometime in the years to come.

DNA Structure





DNA Replication



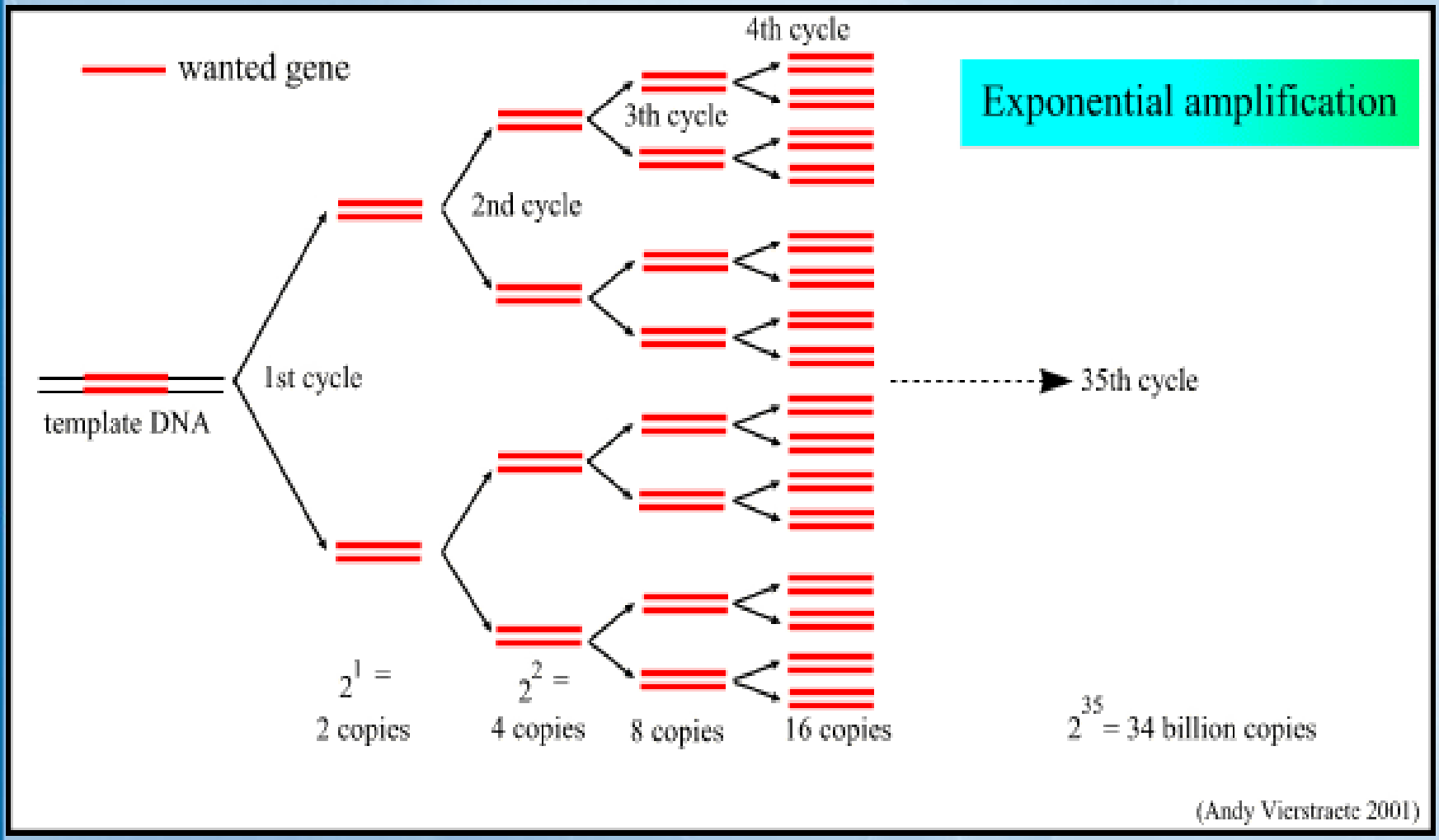
Polymerase Chain Reaction (PCR)

- PCR is a technique which is used to amplify the number of copies of a **specific region of DNA**, (usually fewer than 3000 base pairs) in order to produce enough DNA to be adequately tested.

- **Millions** of copies of a segment of DNA can be made within a few hours

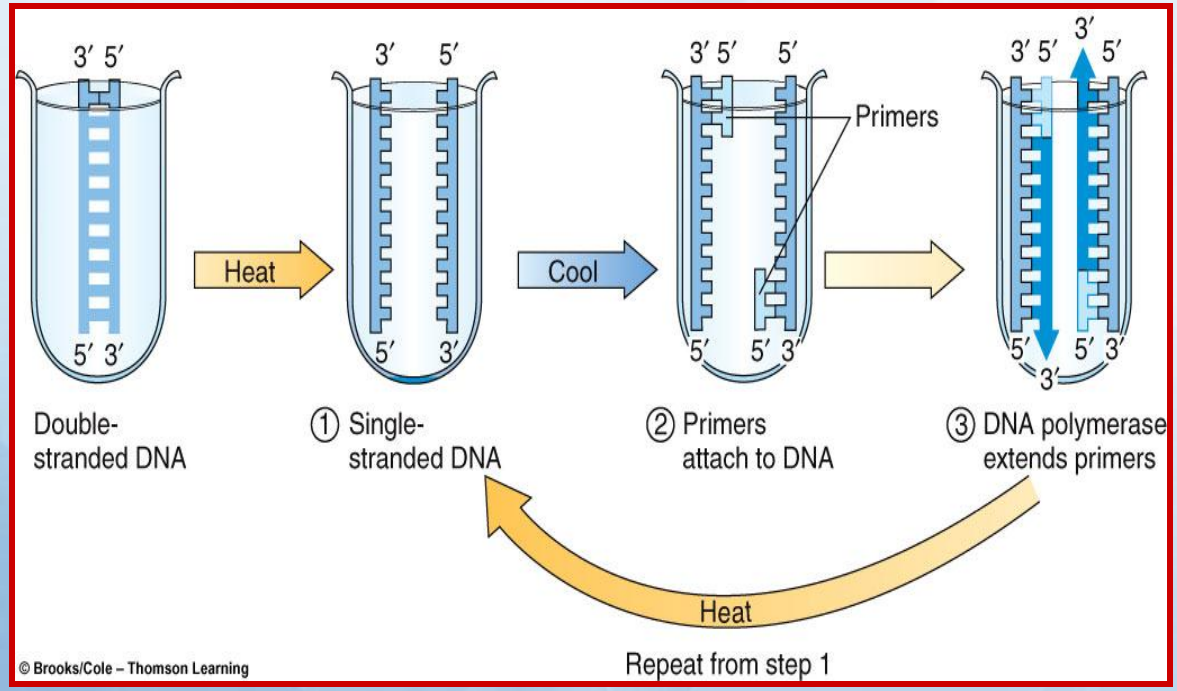
- As a result, it now becomes possible to analyze and characterize the DNA.

DNA amplification by PCR (overview)

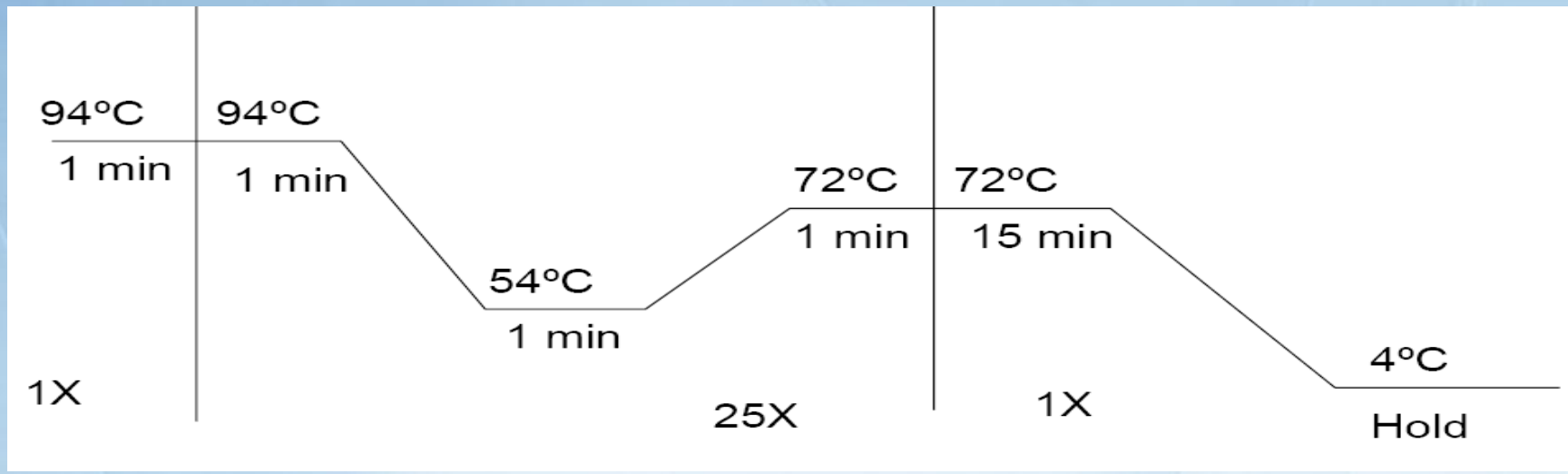


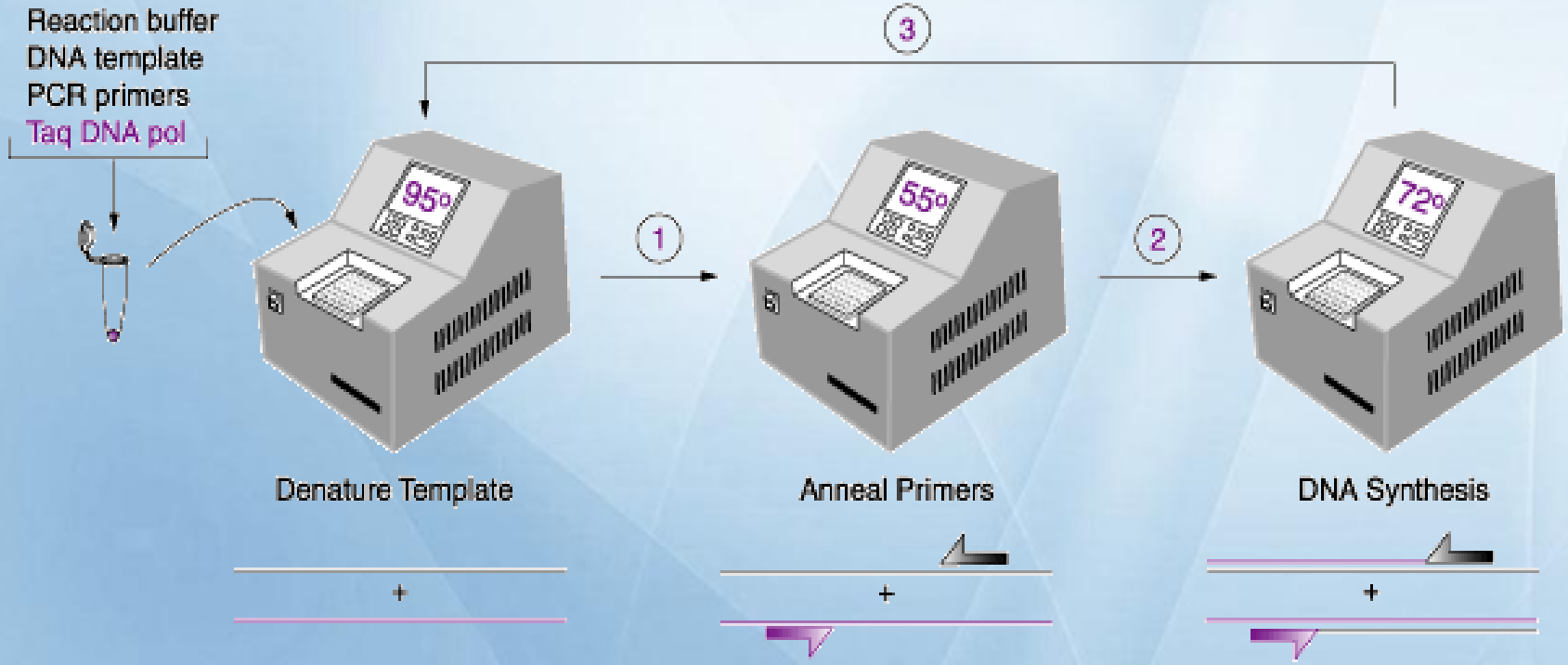
PCR Cycle

- **Each cycle (Round) of PCR contains 3 steps:**
  - **1- Denaturation**
  - **2- Primer annealing**
  - **3- Primer extension**
- **The cycle usually repeated for 25 – 40 times.**

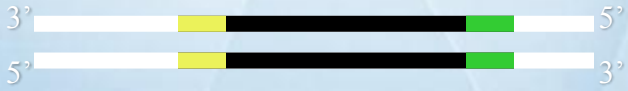


Programming the Thermocycler

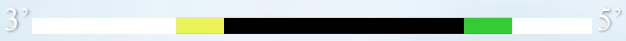


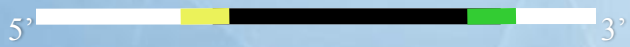


PCR

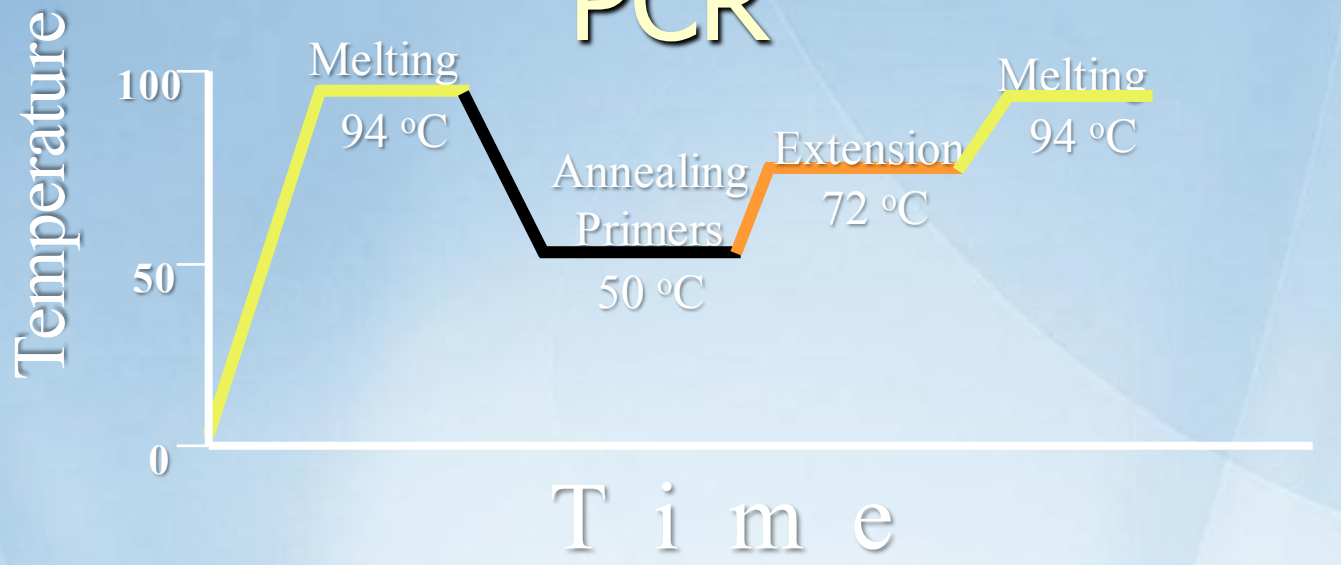


PCR





PCR


Temperature
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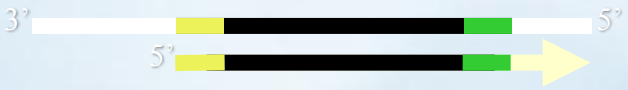
Melting
94 °C
Annealing
Primers
50 °C
Extension
72 °C
Melting
94 °C
T i m e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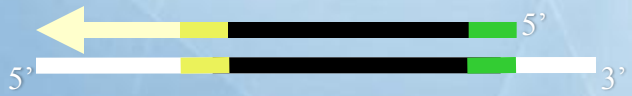
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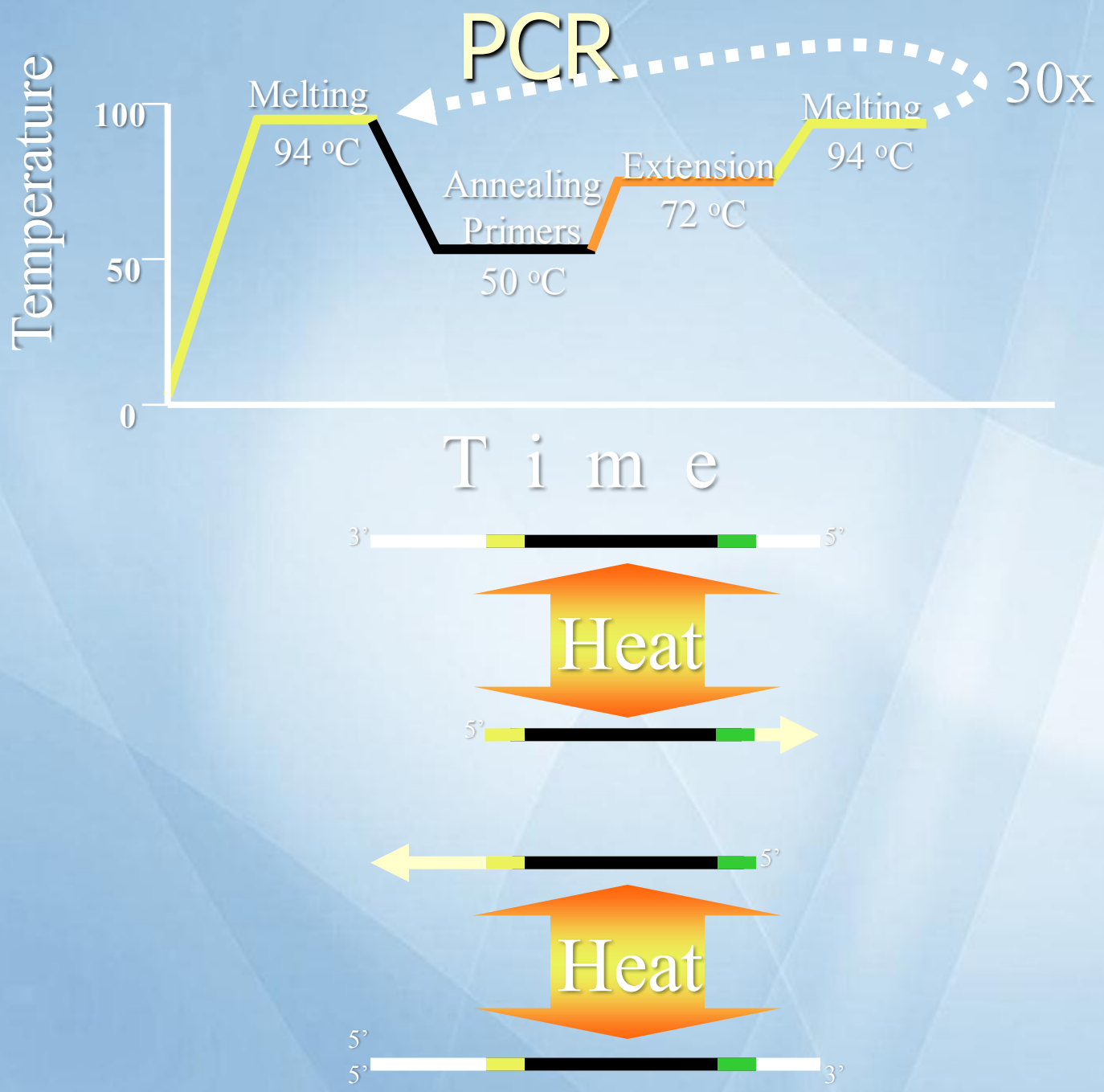
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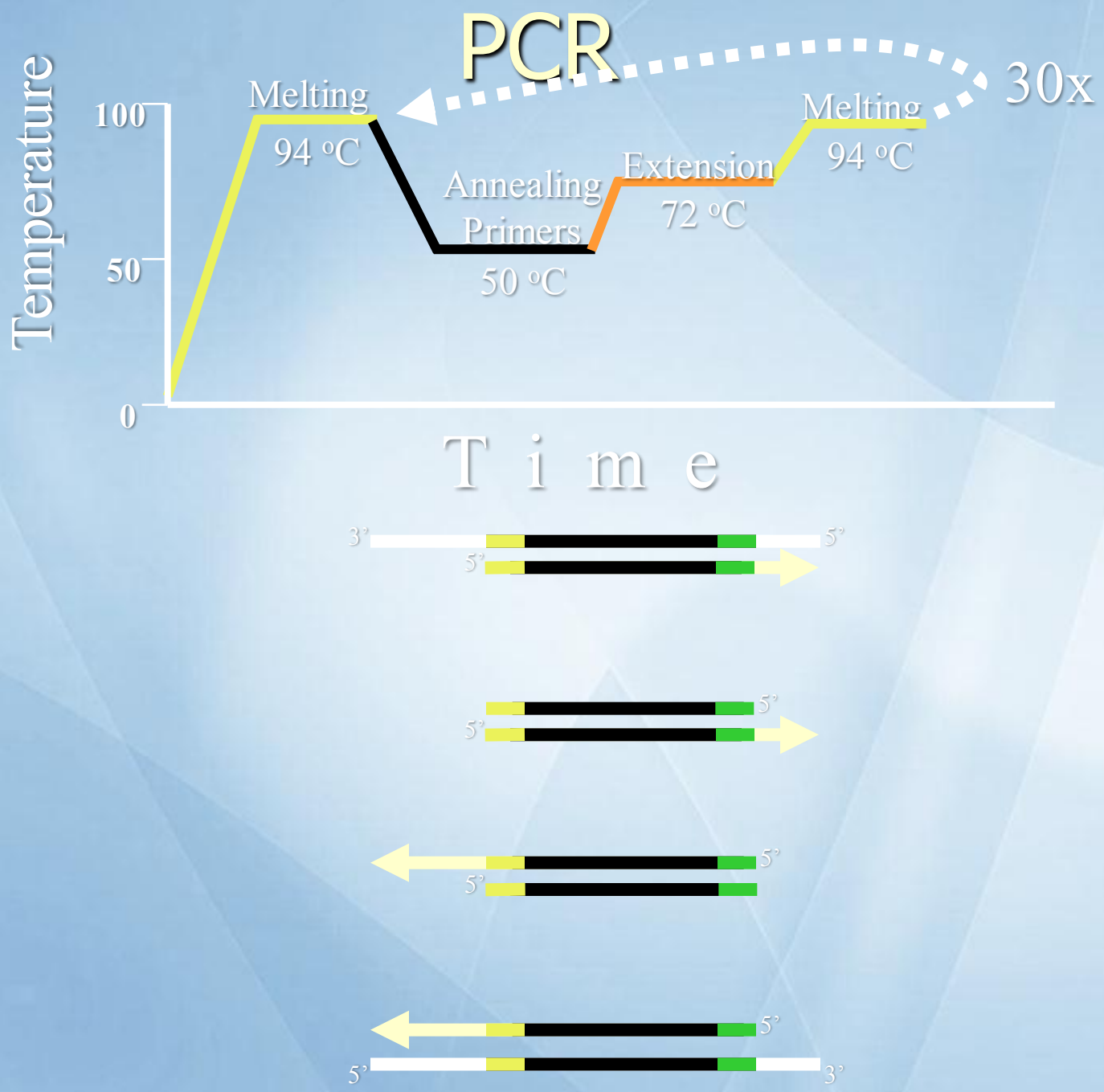
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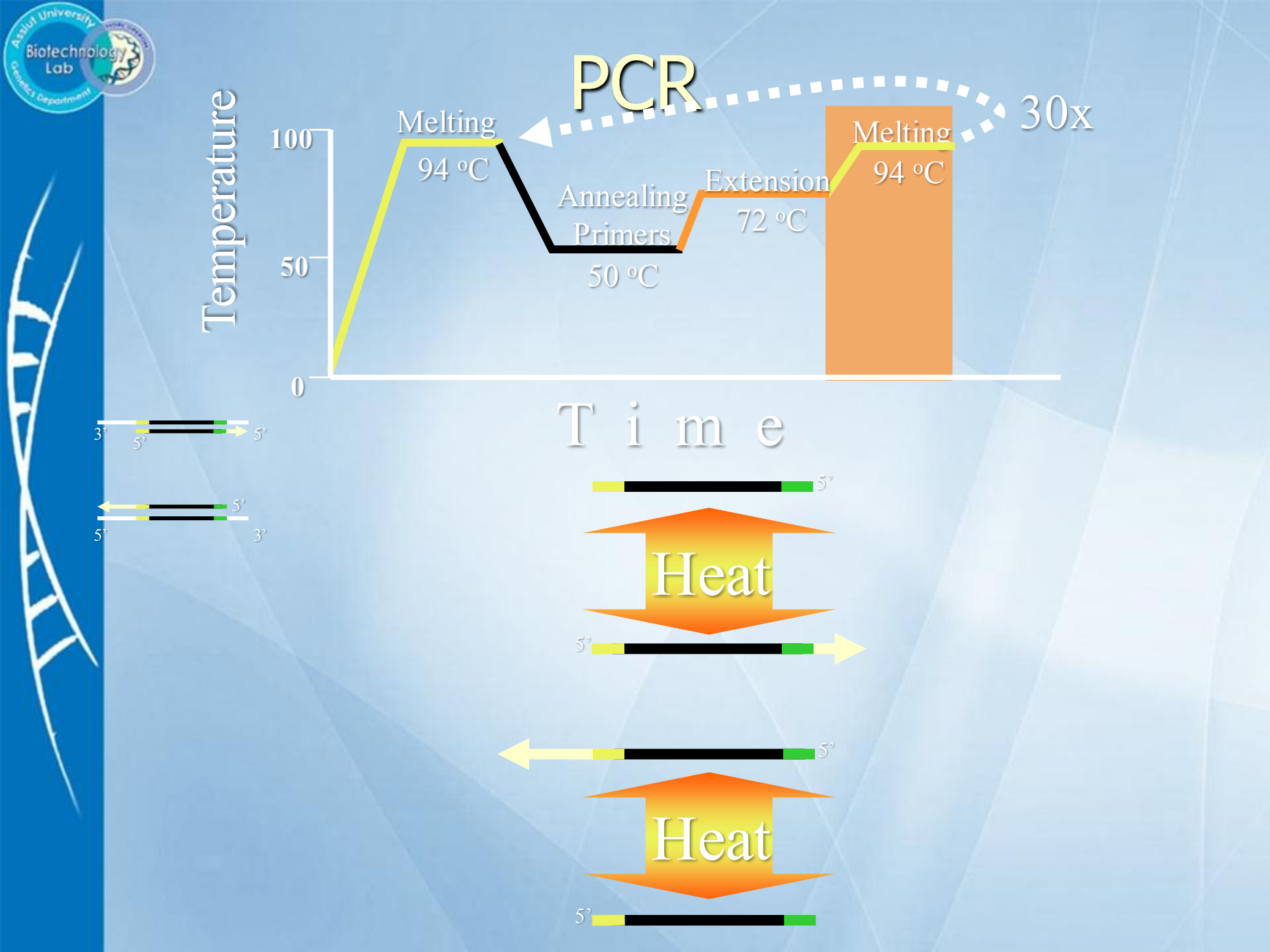
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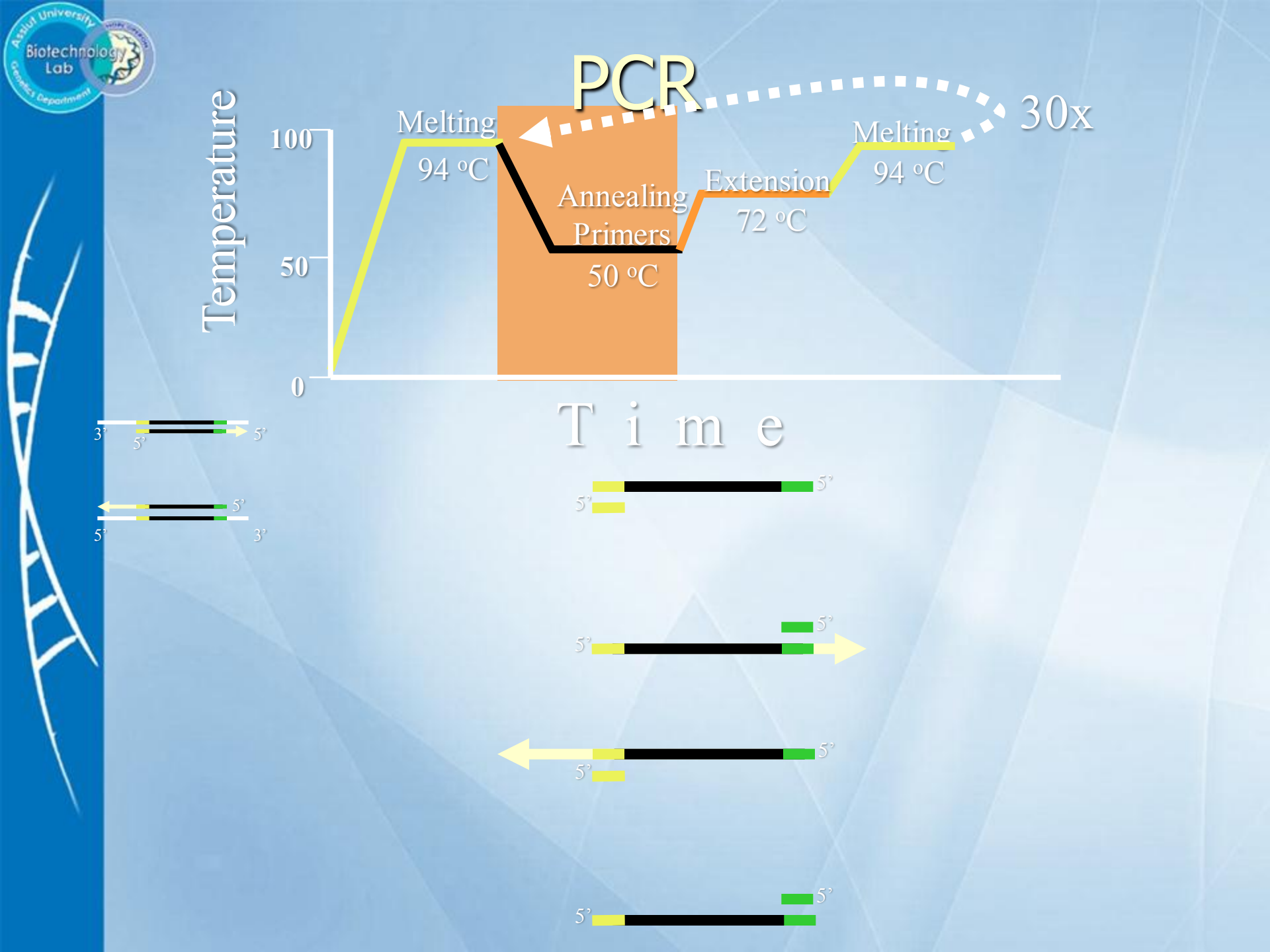
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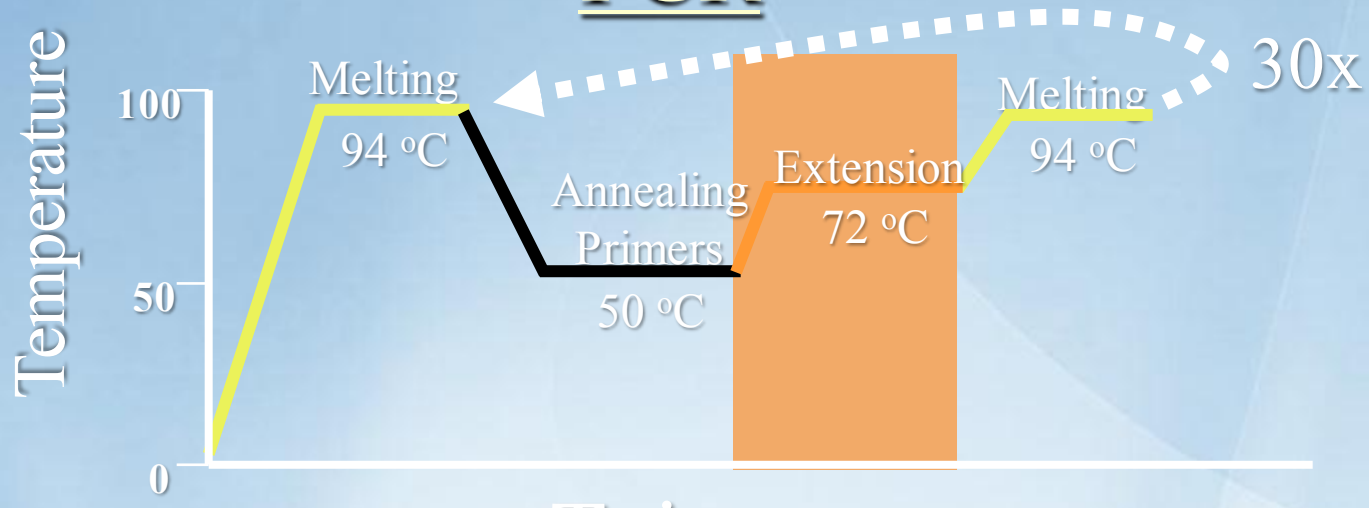
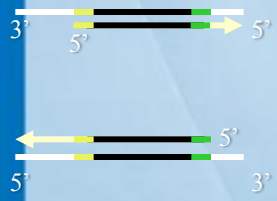
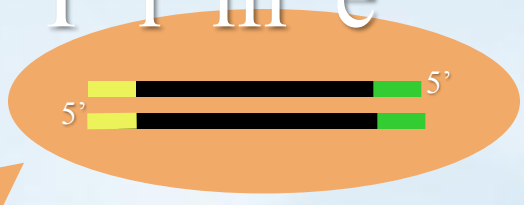
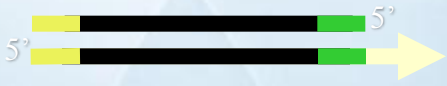
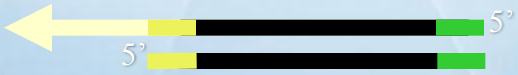


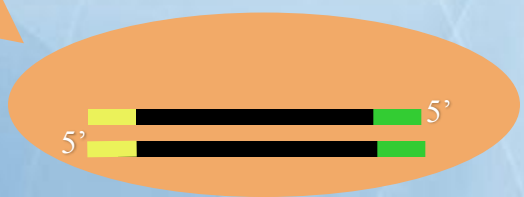




Fragments of defined length







Movie

What do we need for PCR?

PCR tube

THERMOCYCLER

متطلبات تقنيه تفاعل البلمره المتسلسل (PCR)

DNA Sample

Primers

نوعان :

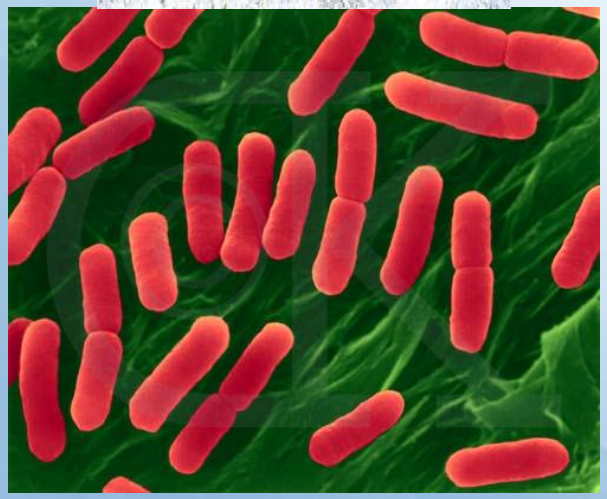
- أمامي (Forward).
- خلفي (Reverse).

وهي تتابع من القواعد النيتروجينيه في شريط واحد قصير(20-25 b) مكمل لبداية الجزء المراد تضخيمه في الـ DNA.

CCGAATGGGATGC

GGCTTACCCTACG

Taq polymerase



- مستخرج من سلالة بكتيريه تسمى *Thermus aquaticus* التي تعيش في المياه الحارة.
- لا يتأثر بدرجات الحرارة المرتفعه.
- درجه الحراة المثلى له ٧٢ ºم.

First reports using DNA polymerase from *Thermus aquaticus was at* (1988)

dNTPs

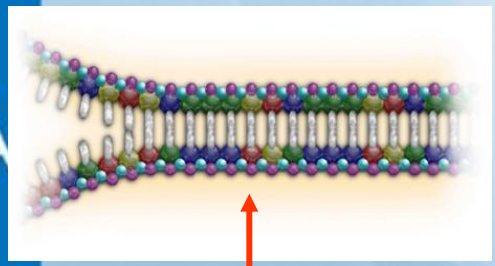
- **Adenine** أدنين
- **Thymine** ثايمين
- **Guanine** جوانين
- **Cytosine** سايتوسين

PCR Buffer 10x

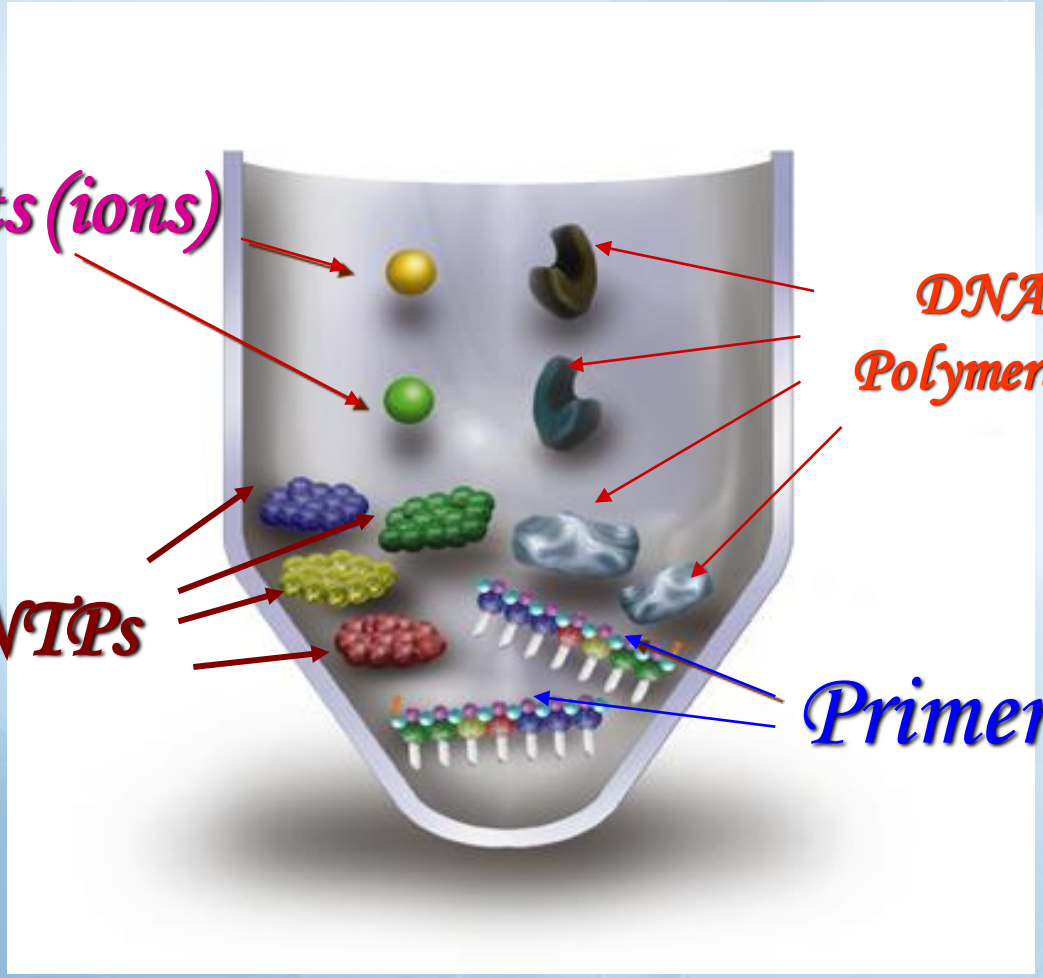
Distilled Water

PCR Procedure

All the required components are inserted into an Eppendorf tube



Template DNA

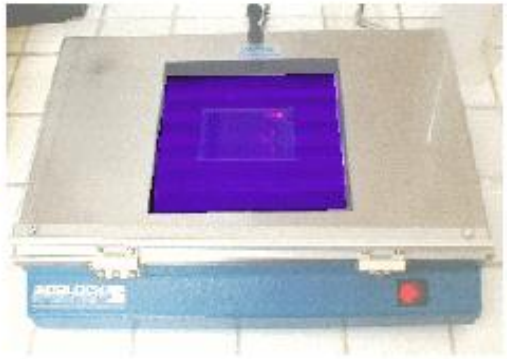


PROCEDURE .....

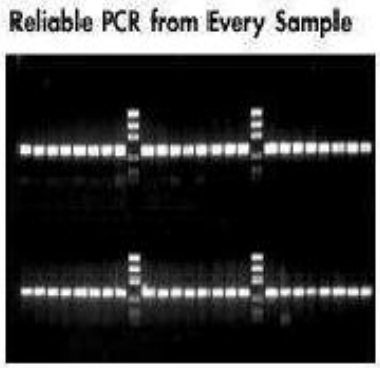
PCR

Agarose gel electrophoresis

3-4 hours



UV visualisation



The final product

PROCEDURE .....

- DNA (Template).
- Forward primer
- Reverse primer
- dNTP's
- *Taq* DNA Polymerase
- Buffer
- H_2O