Sample Preparation

Mix the samples of DNA with the 8X sample loading buffer (8:2)

A. The scientific Name of this material. EDTA Ethelen diamine tetraacetic acid

- B. Two application.
- 1- deactivate metal-dependent enzymes.
- 2- Used to suppress damage to DNA or proteins.

- A. Name of this equipment
- Mortal & pestle
 - B. Benefit of this step in nucleic aced extraction
- Physical grinding or vortexing to separated of the cells in a sample from each other.



Q4- What are the solution which add to lysate in step 2 (DNA binding)

Absolute ethanol (Elushin bufer)

Q5 Enumerate four materials used in RNA extraction

- 1- trizol
- 2- ethanol
- 3- isopropanol
- 4-chloroform
- 5-RNase free water

- Q6- Protocol of DNA extraction from human blood
- 1- samples preparation
- 2-cell lysis
- 3- DNA binding
- 4- wash step
- 5- DNA elution

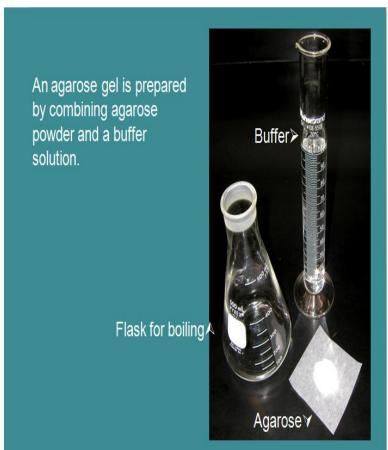
Prepare 1 liter of stock TBE buffer PH 8 (40ml.EDTA,0.5mol.)

108 g tris base55 g boric acid5.8 g EDTAComplete to 1 liter DW. and adjust pH 8.3 by HCl

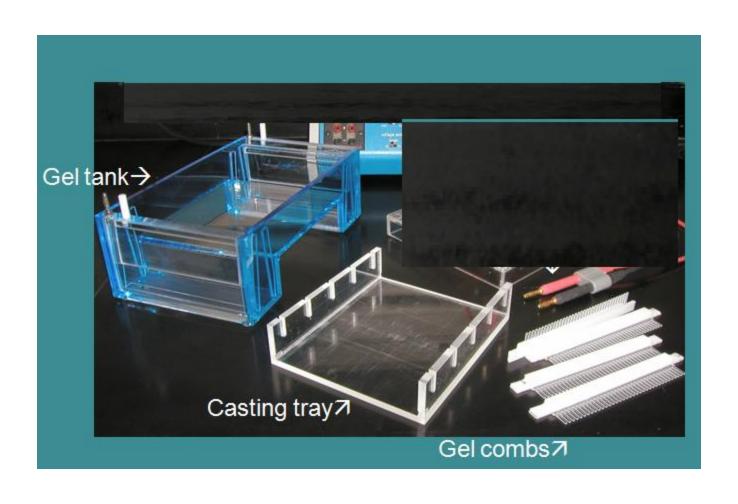
Prepared agarose gel electrophoresis TBE 1X

1- Add 0.7 gm agaros powder + 100 ml (1X TBE) in conical flask + Eth.Br 5 μ l and hooted with cover

2- cooling 50-60 °C and pouring in casting tray

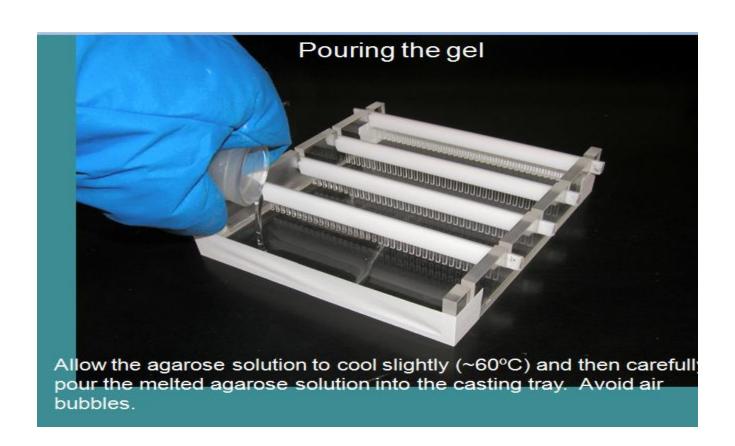


Q9 4-What are the pointed part

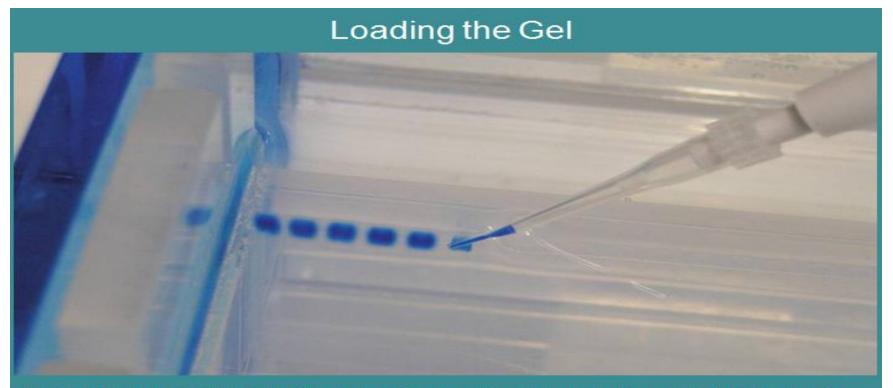


A- Name of this step

B- Explain of this step



Q11 A- Name of this step B- Explain of this step



Carefully place the pipette tip over a well and gently expel the sample. The sample should sink into the well. Be careful not to puncture the gel with the pipette tip.

Benefit of this material in gel electrophoresis

- 1- binds to DNA & fluoresce under UV light allowing the visualization of DNA on the gel and to increase the DNA density
- 2- Can be add to the gel &/or running buffer before the gel is run .

