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## 05 Agarose Gel Electrophoresis

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Working

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## GUIDELINES

## [M]0 Mass Percent

## MATERIALS

NAME ▾

CATALOG # ▾

VENDOR ▾

TAE (Tris-Acetate-EDTA) buffer, 1x

DNA samples

/

1% Agarose gel

/

10×green loading buffer

/

- 1 Use 1×TAE buffer to prepare 1% Agarose mix in a flask, then put it in the microwave and heat it as long as it takes to completely dissolve the Agarose.
- 2 Take out the conical flask, cool it in the wash basin to about 50°C. Add EB quickly, and then mix well. Pour the Agarose gel into gel tray and insert comb into slots. Let the gel solidify for 15-20min. Meanwhile, dilute the 10x green buffer to 1x and add to the DNA samples.  
 ⚡ 50 °C  
 ⌚ 00:15:00 ~ ⌚ 00:20:00
- 3 Place the gel onto the electrophoresis apparatus ensuring that it is totally submerged in 1xTAE buffer. Carefully load each sample into its designated lane and 2ul DNA marker into a separate lane.  
 🧴 2 µl
- 4 Run at 120V for 20-25 min. If the sample have not completely separated, the time may be extended appropriately.  
 ⌚ 00:15:00 ~ ⌚ 00:20:00
- 5 Check the gel using a gel imager or under UV light, then take a photo oNorma.



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