Purpose: To purify and concentrate DNA

1. Add 1:10 ratio of Sodium Acetate: Gel extraction volume
2. Add chilled ethanol in a ratio of 2 times the volume of the gel extraction
3. Centrifuge at 13,000 rpm for 30 minutes
4. Remove supernatant, being careful not to disturb the clear pellet
5. Resuspend in 200 µL of 70% chilled ethanol
6. Centrifuge for 15 minutes at 13,000 rpm
7. Remove supernatant
8. Air dry under hood overnight
9. Resuspend in 100 µL of EB
10. Measur the concentration