

Purpose: To add gene into a plasmid

Ligation

1. Add 6 μL of dH_2O to a clean 1.5 mL Eppendorf tube, 1 μL of T4 DNA Ligase Buffer, 1 μL of plasmid DNA and 1 μL of DNA part and mix.
2. Add 1 μL of T4 DNA Ligase.
3. Pipet mix the tube and incubate at room temperature for 10 minutes.