

Enzyme digestion verification:

DNA	200ng~400ng	n μ l
E1 enzyme		0.2 μ l
10x Buffer		1 μ l
Distilled water (DDW)		To 10 μ l

1. Prepare a centrifuge tube.
2. Add 4.8 μ l of distilled water.
3. Prepare the plasmid, then take 100ng to 200ng of DNA (can take the middle).
4. Add 1 μ l of 10x buffer. This means a tenth of the system, so add 1 μ l to 10 μ l of the system.
5. Add 0.2 μ l of E1 enzyme. It is best to use the enzyme from the same buffer; otherwise, it cannot be used.
6. Mix well by pipetting.
7. Place in a 37°C metal bath, and let the enzyme digest for one to two hours.
8. Perform electrophoresis.