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Dot mutation

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Works for me

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SAFETY WARNINGS

Please wear gloves for the experiment.

- 1 The entire plasmid was amplified reversely by PCR using primers with the fragment sequence that you want to replace.
- 2 The temple plasmids in the PCR process were digested by DpnI enzyme.
- 3 The digested product was transferred into DH5 α . Overnight culture them at 37°C.
- 4 To determine whether the vector was constructed successfully, colony PCR and enzyme digestion were done.
- 5 Select the positive results for sequencing and the final results were obtained.

