## **Blunt-end**

Protocol for blunting ends by 3' overhang removal and 3' recessed end fill in:

- 1. Dissolve the DNA in any 1X restriction enzyme reaction NEB Buffer or 1X T4 DNA Polymerase Reaction Buffer (supplemented with 100µM dNTPs).
- 2. Add 1 unit of T4 DNA Polymerase for each µg DNA.
- 3. Incubate for 15 minutes at 12°C.
- 4. Add EDTA to make the final concentration become 10mM.
- 5. Heat at 75°C for 20 minutes.