AmpliPurifi™ ExTerminator
Nucleotide terminator removal kit for cycle sequencing reactions
Cat ## **Protocol**

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1. Add **5μl** of **mix blue** to cycle sequencing reaction mixture, performed in 20μl volume.
If cycle sequencing reaction is less than 20μl, add sterile water to reach the final volume of 20μl.

2. Add **100μl** of **bind-wash** solution and mix sample by pipetting

3. Load the whole sample onto the minicolumn

4. **Spin** at **12000 – 14000 RPM** for **1 minute**

5. Apply onto the minicolumn **400μl** of **bind-wash solution**

6. **Spin** at **12000 – 14000 RPM** for **2 minutes**

7. Transfer the minicolumn to a new 1.5ml tube (supplied) and apply precisely on the membrane (centre of dark blue circle) **25μl** of **water** or **TSR** (Template Suppression Reagent)

8. **Incubate** at **RT** for **2 minutes**

9. **Spin** at **12000 – 14000 RPM** for **1 minute**

10. A light blue colour of eluted sample confirms proper purification of sample which is ready for thermal denaturation. Storage (-20C) and denaturation can be performed directly in the tube with attached minicolumn.

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