# **ADS Sequencing Reaction Cleaning Beads User Manual**

#### **Product catalog numbers**

Catalog Number	Unit Size
080008	8 ml
080050	50 ml
080250	250 ml
081000	4x 250 ml

#### **Product description:**

ADS Sequencing Reaction Cleaning Beads have been specifically optimized to remove salts and unincorporated dye terminators from sequencing reaction mixes. It offers a simple, efficient way for sequencing reaction purification with a fast workflow.

**Storage temperature**: 4°C

### Sequencing reaction protocol

- 1. Prepare wash solution: 85% ethanol (use of freshly prepared solution is recommended to avoid ethanol evaporation).
- 2. Add 10 ul magnetic beads to 10 ul sequencing reaction.
- 3. Add 45 ul wash solution.
- 4. Mix well and let sit for 5 minutes.
- 5. Put the plate on a magnetic plate holder for 2 minutes.

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- 6. Without removing magnetic plate, reverse the plate and dump extra liquid into the sink.
- 7. Put two layers of paper towel on the plate holder. Put the plate with magnetic plate holder upside down on the holder.
- 8. Centrifuge at 250 rpm (10-15 xg) for 2 minutes.
- 9. Take plate out and discard paper towels.
- 10. Add 100 ul wash solution.
- 11. Repeat steps 6 to 9 one time (two times if there is still dye terminator in the final elution solution).
- 12. Add 45-65 ul deionized H<sub>2</sub>O to elute sequencing reaction products.
- 13. Mix and incubate for 5 minutes at room temperature. Put the plate on the magnetic plate holder and load them on the genetic analyzer for capillary electrophoresis.

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