ADSTM PCR Cleaning Beads User Manual

Product catalog number

Catalog Number	Unit Size
170001	1 mL
170005	5 mL
170050	50 ml

Pleasanton, CA 94588

Product description

 $\mathsf{ADS}^\mathsf{TM}\ \mathsf{PCR}\ \mathsf{Cleaning}\ \mathsf{Beads}\ \mathsf{are}\ \mathsf{used}\ \mathsf{to}\ \mathsf{purify}\ \mathsf{PCR}\ \mathsf{products}\ \mathsf{for}\ \mathsf{different}\ \mathsf{applications},\ \mathsf{especially}\ \mathsf{for}$ Sanger Sequencing. The beads specifically bind to DNA and purify PCR products from other components in the PCR reaction such as dNTPS, primers, and DNA polymerase. The process takes ~30 minutes and is easy to scale up for high-throughput purification.

Product features and storage conditions

- Store at 4 °C for up to 6 months upon arrival
- Use the product at room temperature
- Use 1.5:1 beads volume : PCR product
- > 70% PCR products recovery
- High quality sequencing results from purified PCR product

Materials not provided with the product

- Magnetic rack
- 75% ethanol
- TE buffer (pH8.0)

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PCR product purification procedures

- 1. Vortex the ADS beads for 10 seconds at room temperature before use.
- 2. Add an appropriate volume of ADS beads to the PCR reaction according to the table below:

PCR Reaction Volume	ADS Beads Volume to Be Added (1.5X Vol.)
$10~\mu L$ (if the PCR reaction is <10 μl, make up to 10 μl with TE buffer, pH8.0)	15 μL
20 μL	30 μL
30 μL	45 μL
40 μL	60 μL
50 μL	75 μL

- 3. Vortex briefly to mix well and avoid any bubbles.
- 4. Let the mixture sit for 5 minutes for DNA binding.
- 5. Place the tube in a magnetic rack for 2 minutes.
- 6. Use a pipette to carefully remove the liquid without disturbing the beads.
- 7. With the tube in the magnet rack, add 200 μ L of 75% ethanol and incubate for 1 minute.
- 8. Repeat step 6 and 7.
- 9. Remove the tube from the magnetic rack and air dry the ADS beads for up to 5 minutes. Make sure no liquid is left in the tube.
- 10. Add 10 to 50 μ LTE buffer (pH 8.0) to the tube and vortex briefly. Let the tube sit for 5 minutes.
- 11. Put the tube back to the magnetic rack for 2 minutes.
- 12. Transfer the eluted DNA solution to a clean tube.
- 13. Check DNA quality and quantify DNA for downstream applications.

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