

CleanSpin

Gel DNA Recovery Kit

1. Add 3 volumes of ADB Buffer to each volume of gel.
2. Incubate at 55 °C for 5-10 minutes (do not incubate above 60 °C).
3. Add the melted agarose solution into a CleanSpin Column and place it into a Collection Tube.
4. Centrifuge for 30 seconds. Empty the collection tube when necessary.
5. Add 200 μ l of Wash Buffer to the column and centrifuge for 30 seconds. Repeat the wash step.
6. Place the CleanSpin Column into a new 1.5 ml tube. Add 6-10 μ l of water directly to the column matrix and spin to elute the DNA.



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