

GeneRead™ rRNA Depletion Nano Kit, Part 2

The RNeasy® MinElute® Cleanup Kit is supplied with the GeneRead rRNA Depletion Nano Kit. Store RNeasy MinElute spin columns immediately upon receipt at 2–8°C. The remaining components of the RNeasy MinElute Cleanup Kit should be stored at room temperature (15–25°C) under dry conditions. All components of the RNeasy MinElute Cleanup Kit are stable for at least 9 months under these conditions. Refer to part 1 of the protocol for information on storage of the components of the GeneRead rRNA Depletion Nano Kit.

Further information

- GeneRead rRNA Depletion Handbook: www.qiagen.com/handbooks
- Safety Data Sheets: www.qiagen.com/safety
- Technical assistance: toll-free 00800-22-44-6000, or www.qiagen.com/contact

Notes before starting

- This protocol is for the cleanup of depleted RNA with the RNeasy MinElute Cleanup Kit, which is supplied with the GeneRead rRNA Depletion Nano Kit.
 - Add 4 volumes of ethanol (96–100%) to Buffer RPE for a working solution.
 - Buffer RLT may form a precipitate during storage. If necessary, redissolve by warming, and then place at room temperature (15–25°C).
1. Add 350 µl Buffer RLT to the RNA sample and mix well.
 2. Add 250 µl of 96–100% ethanol to the diluted RNA and mix well by pipetting. Do not centrifuge. Proceed immediately to step 3.
 3. Transfer the sample to an RNeasy MinElute spin column placed in a 2 ml collection tube (supplied). Close the lid gently, and centrifuge for 15 s at $\geq 8000 \times g$ ($\geq 10,000$ rpm). Discard the flow-through.

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4. Place the RNeasy MinElute spin column in a new 2 ml collection tube (supplied). Add 500 μ l Buffer RPE to the spin column. Close the lid gently, and centrifuge for 15 s at $\geq 8000 \times g$ ($\geq 10,000$ rpm) to wash the spin column membrane. Discard the flow-through.
5. Add 500 μ l of 80% ethanol to the RNeasy MinElute spin column. Close the lid gently, and centrifuge for 2 min at $\geq 8000 \times g$ ($\geq 10,000$ rpm) to wash the spin column membrane. Discard the flow-through and collection tube.
6. Place the RNeasy MinElute spin column in a new 2 ml collection tube (supplied). Open the lid of the spin column, and centrifuge at full speed for 5 min to dry the membrane. Discard the flow-through and collection tube.
7. Place the RNeasy MinElute spin column in a new 1.5 ml collection tube (supplied). Add 14 μ l RNase-free water directly to the center of the spin column membrane. Close the lid gently, and centrifuge for 1 min at full speed to elute the RNA.

For up-to-date licensing information and product-specific disclaimers, see the respective QIAGEN kit handbook or user manual.

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