omega BIO-TEK
innovations in nucleic acid isolation

Product Manual

E.Z.N.A.® HP Total RNA Kit

R6812-00	5 preps
R6812-01	50 preps
R6812-02	200 preps

Manual Date: August 2019
Revision Number: v4.0

For Research Use Only

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E.Z.N.A.® HP Total RNA Kit

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Introduction

E.Z.N.A.® HP Total RNA Kit provides a rapid and easy method for RNA isolation from a small amount of cultured eukaryotic cells or tissues. This kit allows single or simultaneous processing of multiple samples in less than 40 minutes. Normally, 1 x 10^7 eukaryotic cells or 25-30 mg tissue can be used in a single experiment. There is no need for phenol/chloroform extractions, and time-consuming steps such as CsCl gradient ultracentrifugation, and precipitation with isopropanol or LiCl, are eliminated. RNA purified using the E.Z.N.A.® HP Total RNA method is ready for applications such as RT-PCR, RT-qPCR, Northern blotting, poly A+ RNA (mRNA) purification, nuclease protection, and *in vitro* translation.

The E.Z.N.A.® HP Total RNA Kit uses the reversible binding properties of HiBind® matrix, a silica-based material. This is combined with the speed of mini-column spin technology. A specifically formulated high salt buffer system allows 100 µg RNA fragments greater than 200 bases to bind to the matrix. Cells or tissues are lysed under denaturing conditions that inactivates RNase. After the homogenization process by either bead milling or rotor-stator homogenizer, samples are transferred to a HiBind® gDNA Removal Column to remove genomic DNA, and the filtrate is transferred to a HiBind® RNA Mini Column. After a few quick washing steps in which cellular debris and other contaminants are effectively washed away, high-quality RNA is eluted in Nuclease-free Water.

New in this Edition:

August 2019:

- DEPC Water has been replaced with Nuclease-free Water. DEPC Water is no longer provided in this kit.
- PR032 (DEPC Water, 100 mL) has been discontinued and is no longer available to purchase.

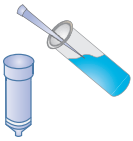
March 2017:

- VAC-08, Omega's vacuum manifold had been discontinued and is no longer available for purchase. Compatible vacuum manifolds are listed on Page 6.
- The amount of 2 mL Collection Tubes has been reduced to reflect the actual number of tubes required in the protocols.

Illustrated Protocols

Centrifugation Protocol

Add GTC Lysis Buffer
and Lyse



Transfer Lysate to
gDNA Removal Column

Add Ethanol
Apply Sample to Column

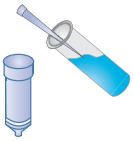
Wash 3X

Dry

Elute

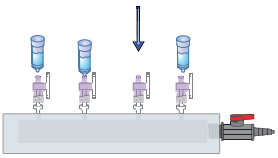
Vacuum Protocol

Add GTC Lysis Buffer
and Lyse



Transfer Lysate to
gDNA Removal Column

Add Ethanol
Apply Sample to Column



Wash 3X

Dry

Elute

Kit Contents

Product	R6812-00	R6812-01	R6812-02
Preparations	5	50	200
HiBind® RNA Mini Columns	5	50	200
gDNA Removal Columns	5	50	200
2 mL Collection Tubes	10	100	400
GTC Lysis Buffer	5 mL	40 mL	150 mL
RNA Wash Buffer I	5 mL	50 mL	200 mL
RNA Wash Buffer II	2 mL	12 mL	50 mL
Nuclease-free Water	2 mL	10 mL	40 mL
User Manual	✓	✓	✓

Storage and Stability

All E.Z.N.A.® HP Total RNA Kit components are guaranteed for at least 24 months from the date of purchase when stored at room temperature. During shipment, crystals or precipitation may form in the GTC Lysis Buffer. Dissolve by warming buffer to 37°C.

Important Notes

Please take a few minutes to read this booklet in its entirety to become familiar with the procedures. Prepare all materials required before starting to minimize RNA degradation.

- Whenever working with RNA, always wear gloves to minimize RNase contamination. Use only clean RNase-free disposable plastic pipette tips when using the supplied reagents.

- Equilibrate samples and reagents to room temperature before beginning this protocol. All steps should be carried out at room temperature unless otherwise noted. Work quickly, but carefully.

- Prepare all materials required before starting the procedure to minimize RNA degradation.

- Carefully apply the sample or solution to the center of the HiBind® RNA Mini Columns. Avoid touching the membrane with pipet tips.

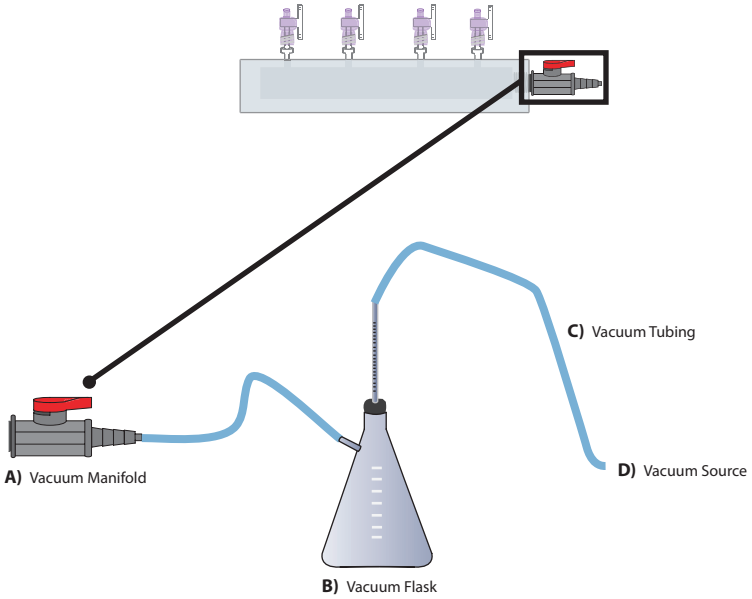
Guidelines for Vacuum Manifold

The following is required for use with the Vacuum Protocol:

A) Vacuum Manifold
Compatible Vacuum Manifolds: Qiagen QIAvac24, Sigma Aldrich VM20, Promega Vacman®, or manifold with standard Luer connector
B) Vacuum Flask
C) Vacuum Tubing
D) Vacuum Source (review tables below for pressure settings)

Conversion from millibars:	Multiply by:
millimeters of mercury (mmHg)	0.75
kilopascals (kPa)	0.1
inches of mercury (inHg)	0.0295
Torrs (Torr)	0.75
atmospheres (atm)	0.000987
pounds per square inch (psi)	0.0145

Illustrated Vacuum Setup:



Quantification and Storage of RNA

To determine the concentration and purity of RNA, measure absorbance at 260 nm and 280 nm with a spectrophotometer. One OD unit measured at 260 nm corresponds to 40 µg/mL RNA. Nuclease-free Water is slightly acidic and can dramatically lower absorbance values. We suggest that you dilute the sample in a buffered solution (TE) for spectrophotometric analysis. The A_{260}/A_{280} ratio of pure nucleic acids is 2.0, while an A_{260}/A_{280} ratio of 0.6 denotes pure protein. A ratio of 1.8-2.0 corresponds to 90%-100% pure nucleic acid. Phenol has a maximum absorbance at 270 nm and can interfere with spectrophotometric analysis of DNA or RNA. Store RNA samples at -70°C in water. Under these conditions, RNA is stable for more than a year.

Integrity of RNA

It is highly recommended that RNA quality be determined prior to beginning all downstream applications. The quality of RNA can be best assessed by denaturing agarose gel electrophoresis with ethidium bromide staining. The ribosomal RNA bands should appear as sharp, clear bands on the gel. The 28S band should appear to be double that of the 18S RNA band (23S and 16S if using bacteria). If the ribosomal RNA bands in any given lane are not sharp and appear to be smeared towards the smaller sized RNA, it is very likely that the RNA undergone degradation during the isolation, handling, or storage procedure. Although RNA molecules less than 200 bases in length do not efficiently bind to the HiBind® matrix, a third RNA band, the tRNA band, may be visible when a large number of cells are used.

Expected Yields:

For animal cell yields, see Page 10.

For animal tissue yields, see Page 16.

Preparing Reagents

1. Dilute RNA Wash Buffer II with 100% ethanol as follows and store at room temperature.

Kit	100% Ethanol to be Added
R6812-00	8 mL
R6812-01	48 mL
R6812-02	200 mL

2. Add 20 µL 2-mercaptoethanol (β-mercaptoethanol) per 1 mL GTC Lysis Buffer. This mixture can be stored for 4 weeks at room temperature.

Disruption Techniques for Tissue Samples

Efficient sample disruption and homogenization is essential for successful total RNA isolation. Cell wall and plasma membrane disruption is necessary for the release of RNA from the sample and homogenization is necessary to reduce the viscosity of the lysates. Homogenization shears genomic DNA and other high-molecular-weight cell components creating a homogeneous lysate. Incomplete homogenization can cause the HiBind® RNA Mini Column to clog resulting in low or no yield.

Rotor-Stator Homogenizer

Using a rotor-stator homogenizer for sample disruption can simultaneously disrupt and homogenize most samples. The process usually takes less than a minute depending on sample type. Many rotor-stator homogenizers operate with differently sized probes or generators that allow sample processing in 50 mL tubes.

Bead Milling

By using bead milling, cells and tissue can be disrupted by rapid agitation in the presence of glass beads and a lysis buffer. The optimal size of glass beads to use for RNA isolation are 0.5 mm for yeast/unicellular cells and 3-6 mm for animal tissue samples.

E.Z.N.A.® HP Total RNA Kit - Animal Cell Protocol

All centrifugation steps used are performed at room temperature.

Materials and Equipment to be Supplied by User:

- Microcentrifuge capable of at least 14,000 x *g*
- Vortexer
- RNase-free pipette tips
- RNase-free 1.5 mL microcentrifuge tubes
- 100% ethanol
- 70% ethanol in sterile DEPC-treated water
- 2-mercaptoethanol (β-mercaptoethanol)
- Disruption equipment
  - Glass beads
  - Rotor-stator homogenizer

Before Starting:

- Prepare GTC Lysis Buffer and RNA Wash Buffer II according to the "Preparing Reagents" section on Page 8.

1. Determine the proper amount of starting material.

   Note: It is critical to use the correct amount of cultured cells in order to obtain optimal yield and purity with the HiBind® RNA Mini Column. The maximum amount of cells that can be processed with the HP Total RNA Protocol is dependent on the cell line and its RNA content. The maximum binding capacity of the HiBind® RNA Mini Column is 100 µg. The maximum number of cells that GTC Lysis Buffer can efficiently lyse is 1 x 10^7. Use the following table as a guideline to select the correct amount of starting material. If no information regarding your starting material is available, begin with 1 x 10^6 cells. Based on RNA yield and quality obtained from 1 x 10^6 cells, the starting amount can be adjusted for the next purification.

Source	Number of Cells	RNA Yield (µg)
IC21	1 x 10^6	12
HeLa	1 x 10^6	15
293HEK	1 x 10^6	10
HIN3T3	1 x 10^6	15

2. Harvest cells using one of the following methods. Do not use more than 1 x 10^7 cells.

   - For cells grown in suspension:

     1. Determine the number of cells.
     2. Centrifuge at 500 x *g* for 5 minutes.
     3. Aspirate and discard the supernatant.
     4. Proceed to Step 3 on Page 12.

   - For cells grown in a monolayer:

     Note: These cells can either be lysed directly in the cell culture dish or trypsinized and collected as a cell pellet prior to lysis. Cells grown in cell culture flasks should always be trypsinized.

     - For direct cell lysis:

       1. Determine the number of cells.
       2. Aspirate and discard the cell culture medium.
       3. Immediately proceed to Step 3 on Page 12.

          Note: Incomplete removal of the cell-culture medium will inhibit lysis and dilute the lysate. This will affect the RNA binding conditions of the HiBind® RNA Mini Column and may reduce RNA yield.

     - To trypsinize and collect cells:

       1. Determine the number of cells.
       2. Aspirate and discard the cell-culture medium and wash the cells with PBS.

          Note: Incomplete removal of the cell-culture medium will inhibit trypsin. Multiple washes may be necessary for cells that are difficult to detach.

       3. Add 0.1-0.25% Trypsin in a balanced salt solution.

4. Incubate for 3-5 minutes to allow cells to detach. Check cells for detachment before proceeding to the next step.

5. Add an equal volume of cell-culture medium containing serum to inactivate the trypsin.

6. Transfer cells to an RNase-free glass or polypropylene centrifuge tube (not supplied).

7. Centrifuge at 500 x *g* for 5 minutes.

8. Aspirate the supernatant.

9. Proceed to Step 3 below.

   Note: Incomplete removal of the cell-culture medium will inhibit lysis and dilute the lysate. This will affect the conditions for binding of RNA to the HiBind® RNA Mini Column and may reduce RNA yield.

3. Disrupt cells (do not use more than 1 x 10^7 cells) with GTC Lysis Buffer. Vortex or pipet up and down to mix thoroughly.

   Note: Add 20 µL 2-mercaptoethanol per 1 mL GTC Lysis Buffer before use.

   Note: For pelleted cells, loosen the cell pellet thoroughly by flicking the tube before adding the appropriate amount of GTC Lysis Buffer based on the table below. To directly lyse the cells in the culture dish, add the appropriate amount of GTC Lysis Buffer directly to the dish. Collect the cell lysate with a rubber policemen and transfer the cell lysate into a 1.5 mL microcentrifuge tube.

Number of Cells	Amount of GTC Lysis Buffer (µL)
< 5 x 10^6	350 µL
5 x 10^6 - 1 x 10^7	700 µL
Dish Diameter (cm)	**Amount of GTC Lysis Buffer (µL)**
< 6	350 µL
6-10	700 µL

4. Insert a gDNA Removal Column into a 2 mL Collection Tube.

5. Transfer the lysate to the gDNA Removal Column.

6. Centrifuge at 13,000 x *g* for 1 minute.

7. Save the filtrate and discard the gDNA Removal Column.

8. Add 1 volume 70% ethanol. Vortex to mix thoroughly. Do not centrifuge.

   Note: A precipitate may form at this point. This will not interfere with the RNA purification. If any sample volume is lost during homogenization, adjust the volume of ethanol accordingly.

9. Insert a HiBind® RNA Mini Column into a 2 mL Collection Tube.

10. Transfer 700 µL sample (including any precipitate that may have formed) to the HiBind® RNA Mini Column.

11. Centrifuge at 10,000 x *g* for 1 minute.

12. Discard the filtrate and reuse the Collection Tube.

13. Repeat Steps 10-12 until all of the sample has been transferred to the column.

Optional: This the starting point of the optional on-membrane DNase I Digestion Protocol. Since the HiBind® matrix of the RNA Mini Column eliminates most DNA, DNase I digestion is not necessary for most downstream applications. However, certain sensitive RNA applications may require further DNA removal. If an additional RNA removal step is required, please continue to the DNase I Digestion Protocol found on Page 23. (See DNase I Digestion Set, (E1091) for more information). If DNase I digestion is not required, proceed to Step 14.

14. Add 500 µL RNA Wash Buffer I.

15. Centrifuge at 10,000 x *g* for 30 seconds.

16. Discard the filtrate and reuse the Collection Tube.

17. Add 500 µL RNA Wash Buffer II.

    Note: RNA Wash Buffer II must be diluted with 100% ethanol before use. Please see Page 8 for instructions.

18. Centrifuge at 10,000 x *g* for 1 minute.

19. Discard the filtrate and reuse the Collection Tube.

20. Repeat Steps 17-19 for a second RNA Wash Buffer II wash step.

21. Centrifuge at maximum speed for 2 minutes to completely dry the HiBind® RNA Mini Column.

    Note: It is important to dry the HiBind® RNA Mini Column matrix before elution. Residual ethanol may interfere with downstream applications.

22. Transfer the HiBind® RNA Mini Column to a clean 1.5 mL microcentrifuge tube (not provided).

23. Add 40-70 µL Nuclease-free Water.

    Note: Make sure to add water directly onto the HiBind® RNA Mini Column matrix.

24. Centrifuge at maximum speed for 2 minutes and store eluted RNA at -70°C.

    Note: Any combination of the following steps can be used to help increase RNA yield.

    - Heat the Nuclease-free Water to 70°C before adding to the column.
    - Increase the incubation time to 5 minutes.
    - Increase the elution volume.
    - Repeat the elution step with fresh Nuclease-free Water (this may increase the yield, but decrease the concentration).
    - Repeat the elution step using the eluate from the first elution (this may increase yield while maintaining elution volume).

E.Z.N.A.® HP Total RNA Kit - Animal Tissue Protocol

All centrifugation steps used are performed at room temperature.

Materials and Equipment to be Supplied by User:

- Microcentrifuge capable of at least 14,000 x *g*
- Vortexer
- RNase-free pipette tips
- RNase-free 1.5 mL microcentrifuge tubes
- 100% ethanol
- 70% ethanol in sterile DEPC-treated water
- 2-mercaptoethanol (β-mercaptoethanol)
- Disruption equipment
  - Glass beads
  - Rotor-stator homogenizer

Before Starting:

- Prepare GTC Lysis Buffer and RNA Wash Buffer II according to the "Preparing Reagents" section on Page 8.

1. Determine the proper amount of starting material.

   Note: It is critical to use the correct amount of tissue in order to obtain optimal yield and purity with the HiBind® RNA Mini Column. The maximum amount of tissue that can be processed with the Total RNA Protocol is dependent on the tissue type and its RNA content. The maximum binding capacity of the HiBind® RNA Mini Column is 100 µg. The maximum amount of tissue that GTC Lysis Buffer can lyse in the this protocol is 30 mg. Use the table on the following page as a guideline to select the correct amount of starting material. If no information regarding your starting material is available, begin with 10 mg. Based on RNA yield and quality obtained from 10 mg, the starting amount can be adjusted for the next purification.

Average Yield of Total Cellular RNA from Mouse Tissue

Source	Amount of Tissue (mg)	RNA Yield (µg)
Brain	10	10
Kidney	10	30
Liver	10	45
Heart	10	5
Spleen	10	33
Lung	10	12
Pancreas	10	40
Thymus	10	20

2. Disrupt the tissue according to one of the following methods described below:

Amount of GTC Lysis Buffer per Tissue Sample

Amount of Tissue	Amount of GTC Lysis Buffer (µL)
≤ 15 mg	350 µL
20-30 mg	700 µL

Note: For samples stored in RNALater® use 700 µL GTC Lysis Buffer.

A. Rotor-Stator Homogenizer: Disrupt tissue with a rotor-stator homogenizer until the sample is uniform. See Page 9 for details.

B. By using bead milling, cells and tissue can be disrupted by rapid agitation in the presence of glass beads and a lysis buffer. The optimal size of glass beads to use for RNA isolation are 0.5 mm for yeast/unicellular cells and 3-6 mm for animal tissue samples. Disrupt according to manufacturers recommended protocol. See Page 9 for details.

3. Centrifuge at maximum speed for 5 minutes.

Note: In some preparations, a fatty upper layer will form after centrifugation. Transfer of any of the fatty upper layer may reduce RNA yield or clog the column.

4. Insert a gDNA Removal Column into a 2 mL Collection Tube.

5. Transfer the lysate to the gDNA Removal Column.

6. Centrifuge at 13,000 x *g* for 1 minute.

   Note: Make sure that all of the liquid has passed through the gDNA Removal Column after centrifugation. If necessary repeat the centrifugation until all liquid passes through the membrane.

7. Save the filtrate and discard the gDNA Removal Column.

8. Transfer the cleared lysate to a clean 1.5 mL microcentrifuge tube (not provided).

9. Add 1 volume 70% ethanol. Vortex to mix thoroughly. Do not centrifuge.

   Note: A precipitate may form at this point. This will not interfere with the RNA purification. If any sample has lost its volume during homogenization, adjust the volume of ethanol accordingly.

10. Insert a HiBind® RNA Mini Column into a 2 mL Collection Tube.

11. Transfer 700 µL sample (including any precipitate that may have formed) to the HiBind® RNA Mini Column.

12. Centrifuge at 10,000 x *g* for 1 minute.

13. Discard the filtrate and reuse the Collection Tube.

14. Repeat Steps 11-13 until all of the sample has been transferred to the column.

Optional: This the starting point of the optional on-membrane DNase I Digestion Protocol. Since the HiBind® matrix of the RNA Mini Column eliminates most DNA, DNase I digestion is not necessary for most downstream applications. However, certain sensitive RNA applications may require further DNA removal. If an additional RNA removal step is required, please continue to the DNase I Digestion Protocol found on Page 23. (See DNase I Digestion Set, Cat # E1091 for more information). If DNase I digestion is not required, proceed to Step 15 on the next page.

15. Add 500 µL RNA Wash Buffer I.

16. Centrifuge at 10,000 x *g* for 30 seconds.

17. Discard the filtrate and reuse the Collection Tube.

18. Add 500 µL RNA Wash Buffer II.

    Note: RNA Wash Buffer II must be diluted with 100% ethanol before use. Please see Page 8 for instructions.

19. Centrifuge at 10,000 x *g* for 1 minute.

20. Discard the filtrate and reuse the Collection Tube.

21. Repeat Steps 18-20 for a second RNA Wash Buffer II wash step.

22. Centrifuge at maximum speed for 2 minutes to completely dry the HiBind® RNA Mini Column.

    Note: It is important to dry the HiBind® RNA Mini Column matrix before elution. Residual ethanol may interfere with downstream applications.

23. Transfer the HiBind® RNA Mini Column to a clean 1.5 mL microcentrifuge tube (not provided).

24. Add 40-70 µL Nuclease-free Water.

    Note: Make sure to add water directly onto the HiBind® RNA Mini Column matrix.

25. Centrifuge at maximum speed for 2 minutes and store eluted RNA at -70°C.

    Note: Any combination of the following steps can be used to help increase RNA yield.

    - Heat the Nuclease-free Water to 70°C before adding to the column.
    - Increase the incubation time to 5 minutes.
    - Increase the elution volume.
    - Repeat the elution step with fresh Nuclease-free Water (this may increase the yield, but decrease the concentration).
    - Repeat the elution step using the eluate from the first elution (this may increase yield while maintaining elution volume).

E.Z.N.A.® HP Total RNA Kit - Vacuum Protocol

All centrifugation steps used are performed at room temperature.

Materials and Equipment to be Supplied by User:

- Microcentrifuge capable of at least 14,000 x *g*
- Vortexer
- Vacuum manifold
- Vacuum source
- RNase-free pipet tips and 1.5 mL microcentrifuge tubes
- 100% ethanol
- 70% ethanol in sterile DEPC-treated water
- 2-mercaptoethanol (β-mercaptoethanol)
- Disruption Equipment
  - Glass heads
  - Rotor-stator homogenizer

Before Starting:

- Prepare GTC Lysis Buffer and RNA Wash Buffer II according to the "Preparing Reagents" section on Page 8.
- Assemble vacuum manifold. Please see Page 6 for details.

Note: Please read through previous sections of this manual before proceeding with this protocol. Steps 1-8 from the HP Total RNA Animal Cell Kit protocol should be completed or Steps 1-9 from the HP Total RNA Animal Tissue Kit protocol should be completed before loading the sample to the HiBind® RNA Mini Column. Instead of continuing with centrifugation, follow the steps below. Do not use more than $1x10^6$ cells or 10 mg tissue for the vacuum protocol.

1. Prepare the vacuum manifold according to manufacturer's instructions.

2. Connect the HiBind® RNA Mini Column to the vacuum manifold.

3. Transfer the sample to the HiBind® RNA Mini Column.

4. Switch on the vacuum source to draw the sample through the column.

5. Turn off the vacuum.

6. Add 500 µL RNA Wash Buffer I.

7. Switch on the vacuum source to draw the RNA Wash Buffer I through the column.

8. Turn off the vacuum.

9. Add 500 µL RNA Wash Buffer II.

   Note: RNA Wash Buffer II must be diluted with 100% ethanol before use. Please see Page 8 for instructions.

10. Switch on the vacuum source to draw the RNA Wash Buffer II through the column.

11. Turn off the vacuum.

12. Repeat Steps 9-11 for a second RNA Wash Buffer II wash step.

13. Transfer HiBind® RNA Mini Column to a 2 mL Collection Tube provided with this kit.

14. Centrifuge at maximum speed for 2 minutes to completely dry the HiBind® RNA Mini Column.

    Note: It is important to dry the HiBind® RNA Mini Column matrix before elution. Residual ethanol may interfere with downstream applications.

15. Transfer the HiBind® RNA Mini Column to a clean 1.5 mL microcentrifuge tube (not provided).

16. Add 40-70 µL Nuclease-free Water.

    Note: Make sure to add water directly onto the HiBind® RNA Mini Column matrix.

17. Centrifuge at maximum speed for 2 minutes and store eluted RNA at -70°C.

    Note: Any combination of the following steps can be used to help increase RNA yield.

    - Heat the Nuclease-free Water to 70°C before adding to the column.
    - Increase the incubation time to 5 minutes.
    - Increase the elution volume.
    - Repeat the elution step with fresh Nuclease-free Water (this may increase the yield, but decrease the concentration).
    - Repeat the elution step using the eluate from the first elution (this may increase yield while maintaining elution volume).

E.Z.N.A.® HP Total RNA Kit - DNase I Digestion Protocol

Since the HiBind® matrix of the RNA Mini Column eliminates most DNA, DNase I digestion is not necessary for most downstream applications. However, certain sensitive RNA applications may require further DNA removal. (See DNase I Digestion Set, Cat# E1091 for further information).

After completing Steps 1-13 of the Animal Cell Protocol (Pages 10-13) or Steps 1-14 of the Animal Tissue Protocol (Pages 15-17), proceed with the following protocol.

User Supplied Material:

- DNase I Digestion Set (E1091)

1. For each HiBind® RNA Mini Column, prepare the DNase I stock solution as follows:

Buffer	Volume per Prep
E.Z.N.A.® DNase I Digestion Buffer	73.5 µL
RNase-free DNase I (20 Kunitz/µL)	1.5 µL
Total Volume	75 µL

Important Notes:

- DNase I is very sensitive and prone to physical denaturing. **Do not vortex the DNase I mixture.** Mix gently by inverting the tube.
- Freshly prepare DNase I stock solution right before RNA isolation.
- Standard DNase buffers are not compatible with on-membrane DNase I digestion. The use of other buffers may affect the binding of RNA to the HiBind® matrix and may reduce RNA yields and purity.
- All steps must be carried out at room temperature. Work quickly, but carefully.

2. Use the HiBind® RNA Mini Column and 2 mL Collection Tube from Step 13 (Animal Cell protocol, Page 13) or from Step 14 (Animal Tissue protocol, Page 17) for Step 3 on the following page.

3. Add 250 µL RNA Wash Buffer I.

4. Centrifuge at 10,000 x *g* for 1 minute.

5. Discard the filtrate and reuse the Collection Tube.

6. Add 75 µL DNase I digestion mixture directly onto the surface of the membrane of the HiBind® RNA Mini Column.

   Note: Pipet the DNase I directly onto the membrane. DNA digestion will not be complete if some of the mixture is retained on the wall of the HiBind® RNA Mini Column.

7. Let sit at room temperature for 15 minutes.

8. Add 250 µL RNA Wash Buffer I.

9. Let sit at room temperature for 2 minutes.

10. Centrifuge at 10,000 x *g* for 1 minute.

11. Discard the filtrate and reuse the Collection Tube.

12. Add 500 µL RNA Wash Buffer II.

    Note: RNA Wash Buffer II must be diluted with 100% ethanol before use. Please see Page 8 for instructions.

13. Centrifuge at 10,000 x *g* for 1 minute.

14. Discard the filtrate and reuse the Collection Tube.

15. Repeat Steps 12-14 for a second RNA Wash Buffer II wash step.

16. Centrifuge at maximum speed for 2 minutes to completely dry the HiBind® RNA Mini Column matrix.

    Note: It is important to dry the HiBind® RNA Mini Column matrix before elution. Residual ethanol may interfere with downstream applications.

17. Place the column in a clean 1.5 mL microcentrifuge tube (not provided).

18. Add 40-70 µL Nuclease-free Water.

    Note: Make sure to add water directly onto the HiBind® RNA Mini Column matrix.

19. Centrifuge at maximum speed for 2 minutes and store eluted RNA at -70°C.

    Note: Any combination of the following steps can be used to help increase RNA yield.

    - Heat the Nuclease-free Water to 70°C before adding to the column.
    - Increase the incubation time to 5 minutes.
    - Increase the elution volume.
    - Repeat the elution step with fresh Nuclease-free Water (this may increase the yield, but decrease the concentration).
    - Repeat the elution step using the eluate from the first elution (this may increase yield while maintaining elution volume).

Troubleshooting Guide

Please use this guide to troubleshoot any problems that may arise. For further assistance, please contact the technical support staff, toll free, at **1-800-832-8896**.

Problem	Cause	Solution
Little or no RNA eluted	RNA remains on the column	Repeat the elution step.
	Column is overloaded	Reduce the amount of starting material.
Problem	Cause	Solution
Clogged column	Incomplete homogenization	Completely homogenize the sample.
		Increase the centrifugation time.
		Reduce the amount of starting material.
Problem	Cause	Solution
Degraded RNA	Starting culture problems	Freeze starting material quickly in liquid nitrogen.
		Do not store tissue culture cells prior to extraction unless they are lysed first.
		Follow protocol closely and work quickly.
	RNase contamination	Ensure not to introduce RNase during the procedure.
		Check buffers for RNase contamination.
Problem	Cause	Solution
Problem in downstream applications	Salt carry-over during elution	Ensure RNA Wash Buffer II has been diluted with 100% ethanol as indicated on bottle.
		RNA Wash Buffer II must be stored and used at room temperature.
		Repeat wash steps with RNA Wash Buffer II.
Problem	Cause	Solution
DNA contamination	DNA contamination	Digest with RNase-free DNase and inactivate DNase by incubation at 65°C for 5 minutes in the presence of EDTA.
Problem	Cause	Solution
Low Abs ratios	RNA diluted in acidic buffer or water	Nuclease-free Water is acidic and can dramatically lower Abs_{260} values. Use TE Buffer to dilute RNA prior to spectrophotometric analysis.

Ordering Information

The following components are available for purchase separately.
(Call Toll Free at 1-800-832-8896)

Product	Part Number
RNA Wash Buffer I, 100 mL	PR030
RNA Wash Buffer II, 25 mL	PR031
2 mL Collection Tubes, 500/pk, 50 pk/cs	AC-1370-00
1.5 mL DNase/RNase-free Microcentrifuge Tubes, 500/pk, 10 pk/cs	SS1-1210-00
RNase-free DNase Set, 50 preps	E1091
RNase-free DNase Set, 200 preps	E1091-02
Proteinase K (>600 mAU/mL, Solution), 2 mL	AC115
Proteinase K (>600 mAU/mL, Solution), 10 mL	AC116

Notes:

For more purification solutions, visit **www.omegabiotek.com**

AVAILABLE FORMATS


Spin Columns

96-Well Silica Plates


Mag Beads

SAMPLE TYPES


Blood / Plasma

Plasmid


Cultured Cells


Plant & Soil


NGS Clean Up


Tissue


FFPE


Fecal Matter

omega BIO-TEK

innovations in nucleic acid isolation

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