

EXTRAClean Cell Culture Media Exosome Purification and RNA Isolation Maxi Kit Product Insert

Product # 73320

Exosomes are 40 - 150 nm membrane vesicles which are secreted by most cell types. Exosomes can be found in cell culture media, plasma, serum, saliva, urine, amniotic fluid and malignant ascite fluids, among other biological fluids. Evidence has been accumulating recently that these vesicles act as cellular messengers, conveying information to distant cells and tissues within the body. The exosomes contain cell-specific proteins, lipids and RNAs, which are transported to other cells, where they can alter function and/or physiology. These exosomes may play a functional role in mediating adaptive immune responses to infectious agents and tumours, tissue repair, neural communication and transfer of pathogenic proteins. Recent work has demonstrated the presence of distinct subsets of microRNAs within exosomes and other extracellular vesicles (EVs) which depend upon the tumour cell type from which they are secreted. For this reason, exosomal RNA may serve as biomarkers for various diseases including cancer. Another subset of RNA that is found in cell culture media is the free-circulating RNA (fc-RNA). These fc-RNA are usually protein-bound RNA that are leaked from cells either during apoptosis or necrosis. As the RNA molecules encapsulated within exosomes or bound to proteins (fc-RNA) are protected from degradation by RNases, they can be efficiently recovered from cell culture media. In general, these two RNA groups contain valuable information for the discovery of biomarkers that can help with early detection of certain cancer types and for monitoring the disease status.

Norgen's Purification Technology

Norgen's EXTRAClean Cell Culture Media Exosome Purification and RNA Isolation Maxi Kit constitutes an all-in-one system for the purification of exosomes and the subsequent isolation of RNA from different cell culture media sample volumes ranging from 20 mL to 35 mL. The purification is based on spin column chromatography that employs Norgen's proprietary resin. The EXTRAClean columns undergo stringent processing and rigorous quality control measures to minimize contamination traces, ensuring optimal results for sensitive applications such as NGS. The kit is designed to isolate all sizes of RNA, including microRNA. The kit provides a clear advantage over other available kits in that it does not require any special instrumentation, protein precipitation reagents, extension tubes, phenol/chloroform or protease treatments. Moreover, the kit allows the user to elute into a flexible elution volume ranging from 50 μ L to 100 μ L. The purified RNA is free from any protein-bound circulating RNA and of the highest integrity. The purified RNA can be used in a number of downstream applications including real time PCR, reverse transcription PCR, Northern blotting, RNase protection and primer extension, and expression array assays.

Advantages

- Purification and enrichment of intact cell culture media exosomes for functional studies.
- No phenol extractions, Proteinase K treatment, nor carrier RNA required
- No time-consuming ultracentrifugation, filtration nor special syringes required.
- No precipitation reagents nor overnight incubation required.
- Pure exosomes are purified and are free-from any other RNA-binding proteins.
- Purification is based on EXTRAClean spin column chromatography that uses Norgen's proprietary resin separation matrix.

Kit Components

Component	Product # 73320 (15 preps)
Slurry E	12.5 mL
ExoC Buffer	1.5 mL
ExoR Buffer	12 mL
Lysis Buffer A	20 mL
Lysis Additive B	2 mL
Wash Solution A	18 mL
Elution Solution A	6 mL
Mini Filter Column	15
EXTRAClean Mini Spin Columns	15
Collection Tubes	15
Elution tubes (1.7 mL)	30
Product Insert	1

Specifications

Kit Specifications	
Sample Type	Cell-Free Cell Culture Media
Sample Volume Range	20 mL to 35 mL
Size of RNA Purified	All sizes, including small RNA (< 200 nt)
Elution Volume	50-100 μ L
Time to Complete 10 Purifications	40-45 minutes
Average Yields*	Variable depending on specimen

* Please check page 6 for Average Yields and Common RNA Quantification Methods

Storage Conditions and Product Stability

All buffers should be kept tightly sealed and stored at room temperature. This kit is stable for 2 years after the date of shipment. It is recommended to warm **Lysis Buffer A** for 20 minutes at 60°C if any salt precipitation is observed.

Precautions and Disclaimers

This kit is designed for research purposes only. It is not intended for human or diagnostic use. Ensure that a suitable lab coat, disposable gloves and protective goggles are worn when working with chemicals. For more information, please consult the appropriate Material Safety Data Sheets (MSDSs). These are available as convenient PDF files online at www.norgenbiotek.com.

Lysis Buffer A contains guanidine thiocyanate, and should be handled with care. Guanidine thiocyanate forms highly reactive compounds when combined with bleach, thus care must be taken to properly dispose of any of this solution.

If liquid containing these buffers is spilled, clean with suitable laboratory detergent and water. If the spilled liquid contains potentially infectious agents, clean the affected area first with laboratory detergent and water, and then with 1% (v/v) sodium hypochlorite.

Caution: DO NOT add bleach or acidic solutions directly to the sample-preparation waste.

Customer-Supplied Reagents and Equipment

- Disposable powder-free gloves
- Benchtop microcentrifuge
- Micropipettors
- Sterile pipette tips with filters
- Vortex
- 96-100% Ethanol
- Nuclease-Free Water
- 50 mL Tube

Procedures

All centrifugation steps are carried out in a benchtop microcentrifuge. Various speeds are required for different steps, so please check your microcentrifuge specifications to ensure that it is capable of the proper speeds. All centrifugation steps are performed at room temperature. The correct rpm can be calculated using the formula:

$$RPM = \sqrt{\frac{RCF}{(1.118 \times 10^{-5}) (r)}}$$

where *RCF* = required gravitational acceleration (relative centrifugal force in units of g); *r* = radius of the rotor in cm; and *RPM* = the number of revolutions per minute required to achieve the necessary *g*-force.

Working with RNA

RNases are very stable and robust enzymes that degrade RNA. Autoclaving solutions and glassware are not always sufficient to actively remove these enzymes. The first step when preparing to work with RNA is to create an RNase-free environment. The following precautions are recommended as your best defense against these enzymes.

- The RNA area should be located away from microbiological work stations.
- Clean, disposable gloves should be worn at all times when handling reagents, samples, pipettes, disposable tubes, etc. It is recommended that gloves are changed frequently to avoid contamination.
- There should be designated solutions, tips, tubes, lab coats, pipettes, etc. for RNA only.
- All RNA solutions should be prepared using at least 0.05% DEPC-treated autoclaved water or molecular biology grade nuclease-free water.
- Clean all surfaces with commercially available RNase decontamination solutions.
- When working with purified RNA, ensure that they remain on ice during downstream applications.

Notes Prior to Use

- All centrifugation steps are performed at room temperature.
- Ensure that centrifuge tubes used are capable of withstanding the centrifugal forces required.
- The provided EXTRAClean Mini Spin Columns are optimized to be used with a benchtop centrifuge and not to be used on a vacuum apparatus.
- Most standard benchtop microcentrifuges will accommodate Norgen's EXTRAClean Mini Spin Columns.
- Centrifuging Norgen's EXTRAClean Mini Spin Columns at a speed higher than recommended may affect RNA yield.
- Centrifuging Norgen's EXTRAClean Mini Spin Columns at a speed lower than recommended will not affect RNA yield. However, centrifugation at a lower speed may require longer time for the solutions to pass through the spin column.
- Ensure that all solutions are at room temperature prior to use.
- It is highly recommended to warm up Lysis Buffer A at 60°C for 20 minutes and mix well until the solutions become clear again if precipitates are present.

- Prepare a working concentration of the Wash Solution A by adding 42 mL of 96 - 100% ethanol (provided by the user) to the supplied bottle containing 18 mL of concentrated Wash Solution A. This will give a final volume of 60 mL. The label on the bottle has a box that may be checked to indicate that the ethanol has been added.
- If any of the solutions do not go through the EXTRAClean Mini Spin Columns within the specified centrifugation time, spin for an additional 1-2 minutes until the solution completely passes through the Column. Do NOT exceed the centrifugation speed as this may affect RNA yield.

Preparation of Cell-free Cell Culture Media Sample

1. Harvest and transfer the required cell culture media volume into a conical tube and centrifuge at 200 x g (~1,000 RPM) for 15 minutes to remove any cells and debris.
2. Transfer cell-free media into a fresh 15-50 mL conical tube.
 - Cell-Free Cell Culture Media is now ready for exosome purification.

Section 1. Exosome Purification from 20 mL - 35 mL Cell Culture Media

Note: The procedure outlined below is for 35 mL inputs of cell-free media. If processing a sample volume in the range of 20 mL - 35 mL media, simply add 2.5 μ L ExoC Buffer for every 1mL of cell-free media. The volume of Slurry E and ExoR Buffer is constant for any volume to be processed.

1. To 35 mL cell-free media add 87.5 μ L of ExoC Buffer followed by the addition of 600 μ L of Slurry E. **(Note: Mix Slurry E well prior to use. For optimal performance ensure that resin is completely resuspended).**
2. Mix well by vortexing for 10 seconds and let stand at room temperature for 10 minutes.
3. Mix well by vortexing for 10 seconds. Centrifuge for 2 minutes at 2,000 RPM. Discard the supernatant.
4. Apply 600 μ L ExoR Buffer to the slurry pellet and mix well by vortexing for 10 seconds.
5. Incubate the slurry pellet resuspended in the 600 μ L ExoR Buffer at room temperature for 15 minutes.
6. After incubation, mix well by vortexing for 10 seconds then centrifuge for 2 minutes at 500 RPM.
7. Transfer the supernatant to a Mini Filter column assembled with an elution tube and centrifuge for 1 minute at 6,000 RPM. **Do not discard the flowthrough which contains your purified exosomes.**
 - Your exosomes are now ready for RNA isolation (Section 2) or any other downstream applications.

Section 2. Exosomal RNA Isolation

1. Add 900 μ L of Lysis Buffer A and 125 μ L of Lysis Additive B to the 600 μ L ExoR Buffer containing the purified Exosomes (Step 7, Section 1).
2. Mix well by vortexing for 10 seconds then incubate at room temperature for 20 minutes.
3. After incubation add 1.5 mL of 96-100% Ethanol to the mixture from Step 2 and mix well by vortexing for 10 seconds.
4. Transfer 750 μ L of the mixture from Step 3 into an EXTRAClean Mini Spin Column. Centrifuge for 1 minute at 3,300 x g (~6,000 RPM). Discard the flowthrough and reassemble the spin column with its collection tube.
5. Repeat Step 4 three more times to transfer the remaining mixture from Step 3 into the EXTRAClean Mini Spin Column.

6. Apply 600 μ L of Wash Solution A to the column and centrifuge for 30 seconds at 3,300 \times g (~6,000 RPM). Discard the flowthrough and reassemble the spin column with its collection tube.
7. Repeat step 6 one more time, for a total of two washes.
8. Spin the column, empty, for 1 minute at 13,000 \times g (~14,000 RPM). Discard the collection tube.
9. Transfer the EXTRAClean Mini Spin Column to a fresh 1.7 mL Elution tube. Apply 50 μ L of Elution Solution A to the column and centrifuge for 1 minute at 2,000 RPM, followed by 2 minutes at 8,000 RPM.
10. For maximum recovery, transfer the eluted buffer back to the column and let stand at room temperature for 2 minutes. Centrifuge for 1 minute at 400 \times g (~2,000 RPM), followed by 2 minutes at 5,800 \times g (~8,000 RPM).

Appendix A

Exosomal RNA Yield

Exosomal RNA from conditioned cell culture media is normally found in very low amounts (1 - 100 pg/ μ L), therefore measuring exosomal RNA concentration using common quantification methods is very difficult and challenging. Typical yields of exosomal RNA vary significantly depending on the number of exosomes that are shed from the cultured cells, as well as the kind of treatment the cultured cells are receiving, and therefore there is no absolute yield for RNA purified from conditioned cell culture media.

Below is a list of the most common RNA quantification methods, as well as the limit of detection for each of these methods. **Unfortunately, none of these methods can be used reliably for measuring the concentration of RNA purified from exosomes unless large volumes have been processed.** This would only be applicable if exosomes contain the maximum amount of RNA that can fit within the specification range of these quantification tools. It should be noted that the specifications outlined below are based on measuring a pure RNA sample, which will not be the case for the exosomal RNA purified from media. Exosomal RNA is short fragmented RNA which is usually present in less than 1000 bp. Purified Exosomal RNA usually contains traces of proteins which will interfere with most quantification methods, leading to the overestimation of the purified RNA concentration. Therefore, purified RNA contaminated with more proteins will be presented at a higher concentration as compared to RNA purified with less protein contaminants, which in this case will depend on the method used for RNA purification. ***The only reliable method that can assess the quality and the relative quantity of the exosomal RNA is RT-qPCR amplification of a standard RNA using a small RNA amplicon such as the 5S rRNA housekeeping gene.***

Common RNA Quantification Methods

1) Bioanalyzer RNA Quantification kits

	RNA 6000 Nano Kit		RNA 6000 Pico Kit		Small RNA kit
	Total RNA	mRNA	Total RNA	mRNA	Total RNA
Quantitative range	25 - 500 ng/ μ L	25 - 250 ng/ μ L	----	----	50-2000 pg/ μ L
Qualitative range	5 - 500 ng/ μ L	5 - 250 ng/ μ L	50 - 5000 pg/ μ L	250 - 5000 pg/ μ L	50-2000 pg/ μ L
Quantitation accuracy	20% CV	20% CV	30% CV	----	-----

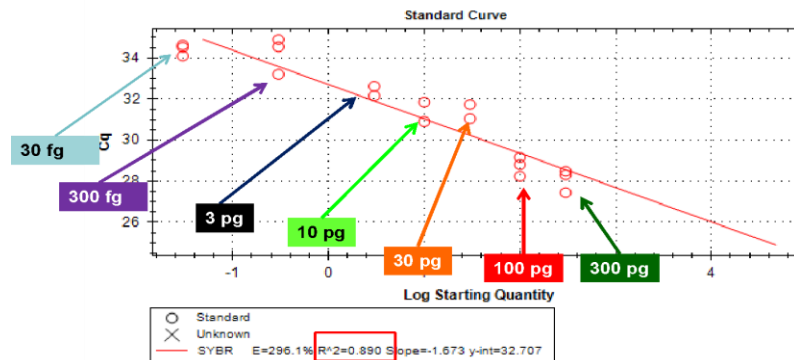
2) NanoDrop 2000

- Detection Limit: 2 ng/ μ l (dsDNA)

3) Quant-iT™ RiboGreen® RNA Assay Kit

- Quantitation Range: 1-200 ng

4) qPCR Standard Curve (generated by Norgen)



Frequently Asked Questions

1. What if a variable speed centrifuge is not available?

A fixed speed centrifuge can be used, however reduced yields may be observed.

2. At what temperature should I centrifuge my samples?

All centrifugation **steps** are performed at room temperature. Centrifugation at 4°C will not adversely affect kit performance.

3. What if I added more or less of the specified reagents' volume?

Adding more or less than the specified volumes may reduce both the quality and the quantity of the purified RNA. Eluting your RNA in high volumes will increase the yield but will lower the concentration. Eluting in small volumes will increase the concentration but will lower the overall yield.

4. What if I forgot to do a dry spin before my final elution step?

Your purified RNA will be contaminated with the Wash Solution A. This may reduce the quality of your purified RNA and will interfere with your downstream applications.

5. Can I perform a second elution?

Yes, but it is recommended that the 2nd elution be in a smaller volume (50% of 1st Elution). It is also recommended to perform the 2nd elution into a separate elution tube to avoid diluting the 1st elution.

6. Why do my samples show low RNA yield?

Exosomes contain very little RNA. This varies significantly depending on the number of exosomes that are shed from the cultured cells, as well as the kind of treatment the cultured cells are receiving. In order to increase the yield, the amount of media could be increased.

7. Why is the A260/280 ratio of the purified RNA lower than 2.0?

Most of the Exosomal RNA is short RNA fragments with a very low concentration where the A260/280 ratio tends to decrease with the decrease in the RNA concentration. The A260/280 ratio is normally between 1 – 1.6. This low A260/280 ratio will not affect any downstream application.

8. Why does my isolated RNA not perform well in downstream applications?

If a different Elution Buffer was used other than the one provided in the kit, the buffer should be checked for any components that may interfere with the application. Common components that are known to interfere are high salts (including EDTA), detergents and other denaturants. Check the compatibility of your elution buffer with the intended use.

9. What should I do if some of the grey resin is transferred out when I am decanting the media supernatant?

Simply remix and recentrifuge. After centrifuging decant the supernatant.

10. What if I added more or less of Slurry E?

Adding less volume may reduce the amount of the purified exosomes. Adding more may not affect the exosome capture but may affect the release of the purified exosomes in the ExoR Buffer

11. What if I added more or less of ExoC Buffer?

Adding a different volume from the specified optimum volume will significantly reduce the amount of the purified exosomes

12. What if I added more or less of ExoR Buffer?

Adding less volume will reduce the release of the captured exosomes in the ExoR Buffer. Adding more will not affect the release of the captured exosomes but it will be more diluted

13. What will happen if accidentally some of the grey resin was transferred with the ExoR buffer?

Any grey resin will be filtered through the Mini Filter Column and the flowthrough which contains the purified exosomes should not contain any grey resin.

Technical Support

Contact our Technical Support Team between the hours of 9:00 and 5:30 (Eastern Standard Time) at (905) 227-8848 or Toll Free at 1-866-667-4362.

Technical support can also be obtained from our website (www.norgenbiotek.com) or through email at support@norgenbiotek.com.

Norgen's purification technology is patented and/or patent pending. See www.norgenbiotek.com/patents

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