

Manual Total RNA Midi

Kit for total RNA purification.

catalog #	size
032-20	20 isolations

For research use only.

Guarantee

A&A Biotechnology provides a guarantee on this product.

The company does not guarantee the correct performance of this kit in the event of:

- not adhering to the supplied protocol
- use of not recommended equipment or materials
- use of other reagents than recommended or which are not a component of the product
- use of expired or improperly stored product or its components



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Specification

form	midicolumn	
binding capacity	500 μg of RNA	
sample size	 up to 10 ml of bacterial culture up to 10 ml of yeast culture up to 10 ml of blood up to 1 x 10⁷ of cell culture up to 150 mg of plant or animal tissue 	
elution volume	from 400 µl	
elution solution	ultrapure water	

Contents

component	20 isolations	storage
Midicolumns	20 pcs	15-25 °C
Counterweight column	1 pcs	15-25 °C
15 ml tubes	40 pcs	15-25 °C
A1 wash solution	200 ml	15-25 °C
Fenozol	50 ml	2-8 °C
Isopropanol	20 ml	15-25 °C
Ultrapure water	15 ml	-20-25 °C

Additional equipment and reagents

Necessary

- 15 ml sterile Falcon tubes
- Chloroform
- Centrifuge
- Heatblock or incubator set to 50 °C

Optional

- RBCL (cat. # 213-100, 213-250)
- DNAse (cat. # 1009-10, 1009-100)
- Clean-Up RNA Concentrator (cat. # 039-25C, 039-100C)

Important notes

When working with RNA, use RNAse-free consumables. Work sterile, use disposable gloves and change them whenever good laboratory practice requires it.

Material preparation

Bacterial / yeast culture

- 1. Centrifuge 10 ml of overnight bacterial culture. Discard supernatants.
- 2. Follow point 1. of the protocol.

Cell culture

- 1. Centrifuge cell culture containing up to 1 x 10⁷ of cells. Discard supernatants.
- 2. Follow point 1. of the protocol.

Plant / animal tissue

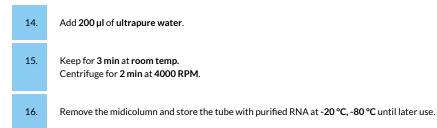
- 1. Homogenize tissue sample (50-150 mg) in liquid nitrogen.
- 2. Transfer the sample to a 15 ml Falcon tube (not included).
- 3. Follow point 1. of the protocol.

Fresh blood (not frozen)

- Add the appropriate amount of RBCL (not included) to 5-10 ml of blood sample. We recommend using 5 volumes of RBCL to 1 volume of blood sample.
- Mix and incubate on ice for 15 min. Note the changing appearance of the sample during the incubation. The initially opaque solution should turn to a completely transparent ruby-red at the incubation end.
- 3. Centrifuge for 5 min at 3000 x g. Carefully discard supernatants.
- 4. Follow point 1. of the protocol.

Isolation protocol

1.	Add 2,4 ml of fenozol and lyse cells by repetitive pipetting.
	 Fenozol deactivates endogenous RNAses. Sample suspended in fenozol can be stored: at -20 °C, -80 °C up to one year from +2 °C to +8 °C up to one week in room temperature up to 24 hours
	Fenozol contains phenol. Avoid contact with skin. Wear suitable protective gloves.
2.	Incubate sample for 5 min at 50 °C .
3.	Add 600 µl of chloroform (not included) and gently mix by inverting the tube a few times.
4.	Keep the sample for 3 min at room temp . Centrifuge the sample for 5 min at 5000 RPM .
	Note: If you have an odd number of samples, please remember about counterweight columns before centrifugation.
5.	Transfer the supernatant to a new 15 ml tube (not included). Add 800 μl of isopropanol .
6.	Thoroughly mix and apply onto the midicolumn.
7.	Centrifuge for 2 min at 4000 RPM .
8.	Add 3.5 ml of A1 wash solution.
9.	Centrifuge for 2 min at 4000 RPM .
10.	Transfer the midicolumn to a new 15 ml tube (included). Add 3.5 ml of A1 wash solution.
11.	Centrifuge for 10 min at 5000 RPM .
12.	Transfer the midicolumn to a new 15 ml tube (included). Add 200 μl of ultrapure water directly onto the minicolumn resin.
13.	Keep for 3 min at room temp. Centrifuge for 2 min at 4000 RPM .



Additional clean-up / concentration of isolated RNA sample (optional)

Total RNA Midi kit effectively isolates and purifies RNA for most downstream applications.

In case of the highest possible RNA sample purity being required, as for example supreme DNA removal, we recommend to additionally process RNA sample, as follows:

Use of the DNAse (cat. # 1009-10, 1009-100)

1. To 400 µl of RNA eluate add:

4 μl of DNAse (10 U/μl) 40 μl of 10x reaction buffer (included with DNAse)

- 2. Incubate for 15 min at 37 °C.
- 3. Incubate for 10 min at 65 °C inactivation of DNAse.

Use of Clean-Up RNA Concentration Kit (cat. # 039-25C, 039-100C)

Kit for removal and concentration of RNA samples. Elution from 15 µl. Microcolums (included with the kit) effectively bind RNA. Most contaminations flow through the microcolumns.

Safety information

	Fenozol
DANGER	H301+H311+H331 Toxic if swallowed, in contact with skin or if inhaled. H314 Causes severe skin burns and eye damage. H341 Suspected of causing genetic defects. H373 May cause damage to organs through prolonged or repeated exposure. H411 Toxic to aquatic life with long-lasting effects. P261 Avoid breathing dust. P273 Avoid release to the environment. P2804 Wear protective gloves, protective clothing, eye protection, face protection. P301+P301 F svallowed: Immediately call a Poison Center or doctor/physician. P305+P351+P338 If in eyes: rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P310 Immediately call a Poison Center or doctor/physician.
	Isopropanol
DANGER	H225 Highly flammable liquid and vapor. H319 Causes serious eye irritation. H336 May cause drowsiness or dizziness. P210 Keep away from heat, sparks, open flames, hot surfaces. No smoking. P261 Avoid breathing vapors. P305+P351+P338 If in eyes: rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
	A1 wash solution
DANGER	H225 Highly flammable liquid and vapor. H319 Causes serious eye irritation. H336 May cause drowsiness or dizziness. P210 Keep away from heat, sparks, open flames, hot surfaces. No smoking. P261 Avoid breathing vapors. P305+P351+P338 If in eyes: rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.



A&A Biotechnology, ul. Strzelca 40, 80-299 Gdańsk, Poland phone +48 883 323 761, +48 600 776 268 info@aabiot.com, www.aabiot.com

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