

Manual

Total RNA Mini Plus D

Kit for total RNA purification with DNA removal on the column.

catalog#	size
042-25	25 isolations
042-100	100 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
- $\bullet \qquad \text{use of other reagents than recommended or which are not a component of the product} \\$
- use of expired or improperly stored product or its components

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Specification

form	microcolumn		
binding capacity	10 µg of RNA		
sample size	 bacterial culture: up to 500 μl yeast culture: up to 500 μl blood: up to 1 ml cell culture: up to 5 x 10⁵ plant or animal tissue: up to 10 mg 		
elution volume	from 15 µl		
elution solution	ultrapure water		

Contents

	25 isolations		100 isolations		
component	quantity	cat#	quantity	cat#	storage
microcolumns	25 pcs	K-C02-25	100 pcs	K-C02-100	15-25 ℃
1.5 ml tubes	25 pcs	K-C01-25P	100 pcs	K-C01-100P	15-25 ℃
2 ml tubes	50 pcs	K-PC-50	200 pcs	K-PC-200	15-25 ℃
DNAse U/μl	220 µl	K-DNA-220B	650 µI	K-DNA-650B	-20 ℃
10x DNAase buffer	1.5 ml	K-BDNA-15A	1.5 ml	K-BDNA-15A	-20℃
A2WE wash solution	45 ml	K-A2WE-45	180 ml	K-A2WE-180	15-25 ℃
R8I wash solution	20 ml	K-R8I-20	80 ml	K-R8I-80	15-25 ℃
Fenozol Plus	15 ml	K-FENP-15	50 ml	K-FENP-50	4℃
Isopropanol	15 ml	K-IZO-15	50 ml	K-IZO-50	15-25 ℃
ultrapure water	8 ml	K-WUP-8	30 ml	K-WUP-30	15-25 ℃

Additional equipment and reagents

Necessary

- 1.5 ml sterile tubes
- microcentrifuge
- heatblock or incubator set to 50 °C

Optional

• RBCL (cat. # 213-100, 213-250)

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 100-500 µl of overnight bacterial culture or yeast culture. Discard supernatants.
- 2. Follow point <u>1. of the protocol</u>.

Cell culture

- 1. Centrifuge cell culture containing 1 x 10⁵-5 x 10⁵ of cells.

 Discard supernatants.
- 2. Follow point <u>1. of the protocol</u>.

Plant, animal tissue

- 1. Homogenize tissue sample (1-10 mg) in liquid nitrogen.
- 2. Transfer the sample into 1.5 ml tube (not included).
- 3. Follow point <u>1. of the protocol</u>.

Fresh blood (not frozen)

Additional reagents you will need:

• RBCL (max. 5 ml per sample), nr kat. 213-100

Add the appropriate amount of RBCL to 1 ml of blood sample.

Attention. We recommend using 5 volumes of RBCL to 1 volume of blood sample.

Mix and incubate on ice for 15 min.

Note. 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 10 min at 3000 x g. Carefully discard supernatants.

4. Follow point <u>1. of the protocol</u>.

DNA digestion solution preparation

Before proceeding to RNA isolation, prepare the appropriate amount of DNA digestion solution according to the following formula (50 µl of solution per sample):

component	volume per 1 sample	volume per 25 samples	volume per 100 samples	n - sample quantity
10x DNAase buffer	5 μΙ	26 x 5 µl	101 x 5 μl	(n+1) x 5 μl
	+	+	+	+
DNAse	6 μΙ	26 x 6 µl	101 x 6 μl	(n+1) x 6 μl
	+	+	+	+
ultrapure water	39 µl	26 x 39 µl	101 x 39 μl	(n+1) x 39 μl
	=	=	=	
DNA digestion solution	50 μΙ	1,3 ml	5,05 ml	

Mix and describe "DNA digestion solution"

Isolation protocol

1. Add 400 µl of Fenozol Plus and pipette until complete lysis cells occurs. $\textbf{Attention.} \ Fenozol\ Plus\ deactivates\ endogenous\ RNAses.}\ Sample\ suspended\ in\ Fenozol\ Plus\ can\ be\ stored:$ at -20 °C, -80 °C up to one year from +4 °C to +8 °C up to one week at room temperature up to 24 hours Fenozol Plus contains phenol. Avoid contact with skin. Wear suitable protective gloves. 2. Incubate sample for 5 min at 50 °C. 3. Add 150 µl of ultrapure water. Intensively vortex for 15 s. 4. Keep the sample for 5 min at room temp. Centrifuge the sample for 10 min at 10 000 RPM. Note. During the centrifugation step, the DNA and proteins are collected at the bottom of the tube while RNA stays dissolved in the supernatant. 5. Transfer 400 µl of the supernatant to a new 1.5 ml tube (not included). Add 400 µl of isopropanol. 6. Thoroughly mix and apply onto the microcolumn. 7. Centrifuge for 1 min at 12 000 RPM. 8. Transfer the microcolumn to a new 2 ml tube (included). Add 700 ul of A2WE wash solution. 9. Centrifuge for 2 min at 12 000 RPM.

Attention. It is necessary to centrifuge min. 2 min to remove ethanol from the microcolumn membrane.

On-column DNAse digestion		
10.	Transfer the microcolumn to a new 2 ml tube (included).	
11.	Add $50\mu l$ of DNA digestion solution directly onto the microcolumn membrane, ensuring no droplets remain on the column inner walls or membrane-securing ring.	
12.	Incubate for 30 min at 37 °C.	
13.	Add 700 μI of R8I wash solution.	
14.	Centrifuge for 1 min at 12 000 RPM.	
15.	Collect the filtrate from the tube and apply it again onto the microcolumn. Place the microcolumn into the same tube.	
16.	Centrifuge for 1 min at 12 000 RPM.	
17.	Add 700 μI of A2WE wash solution.	
18.	Centrifuge for 1 min at 12 000 RPM.	
19.	Remove the microcolumn from the tube and discard the filtrate. Place the minicolumn into the same tube. Add 200 μI of A2WE wash solution.	
20.	Centrifuge for 2 min at 12 000 RPM.	
	$\textbf{Attention}. \ \textbf{It is necessary to centrifuge min. 2 min to remove ethanol from the microcolumn membrane.}$	
21.	Transfer the dry microcolumn to a new 1.5 ml elution tube (included). Add 15-40 µl of ultrapure water directly onto the minicolumn resin.	
22.	Keep for 3 min at room temp. Centrifuge for 1 min at 10 000-12 000 RPM.	
23.	Remove the micrcolumn and store the tube with purified RNA at -20 °C, -80 °C until later use. Note. Elution tube has a long, elastic cap connector. Start closing the tube by careful pressing the cap on the connector side. A "click" sound confirms proper closure. Different ways of closing may cause opening of the tube during storage.	

Safety information





DANGER



Fenozol Plus

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.

P280 Wear protective gloves, protective clothing, eye protection, face protection.

P301+P310 If swallowed: 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

P310 Immediately call a Poison Center or doctor/physician.

Isopropanol





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.

R8I wash solution





H225 Highly flammable liquid and vapor. H319 Causes serious eye irritation.

 $H336\,May\,cause\,drows in ess\,or\,dizziness.$

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.

A2WE wash solution



DANGER

DANGER

H225 Highly flammable liquid and vapor.

H319 Causes serious eye irritation.

P210 Keep away from heat, sparks, open flames, hot surfaces. No smoking.

P233 Keep container tightly closed.

P305+P351+P338 If in eyes: rinse cautiously with water for several minutes. Remove contact lenses,

if present and easy to do. Continue rinsing.



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