

WizMag™ Total RNA

User Manual

Ver 2.0

REF W7070 | W7071 | W7072 | W7073

For in vitro diagnostic use



INTENDED USE

The WizMag™ Total RNA kit is designed to be used on the CLEO™ AP16 Nucleic Acid Extractor System. This kit provides a fast and easy method for the purification of total RNA from cultured cells, body fluids, and liquid biological samples. This kit utilizes advanced magnetic-silica technology to purify high-quality total RNA. Co-purified DNA can be easily removed by DNase I. Purified RNA is highly suited for downstream applications such as PCR-based or enzyme-based reactions.

KIT CONTENTS

Contents	W7070	W7071	W7072	W7073	Storage	
No. of preparation	64	192	32	96		
Pre-packed 96-well Plate	4 ea	12 ea	-	-	Room	
Pre-packed 6-well Strip	-	-	32 ea	96 ea	Temperature	
Plunger	8 ea	24 ea	8 ea	24 ea	(15-25°C)	
Blank solution N	500 uL	500 uL	500 uL	500 uL		

This kit is delivered under ambient conditions.

Long exposure to heat sources can deteriorate the performance of the kit significantly.

QUALITY CONTROL ANALYSIS

In accordance with Wizbiosolutions Inc. ISO 13485-certified Quality Management System, each lot WizMag™ Total RNA kit is tested against predetermined specifications to ensure consistent product quality.

PREVENTING RNASE CONTAMINATION

RNase can be introduced accidentally during RNA preparation. Various factors such as reagents, air, dust, and human hands or skin, can be the sources of RNase contamination. Always wear disposable gloves and use sterile, disposable plastic wares. Use the pipette reserved for RNA work. Maintain a separate area for RNA work. Carefully clean the surfaces.

PRECAUTIONS

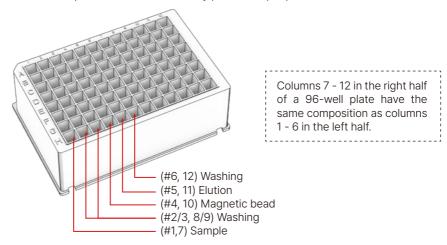


- This product is reserved exclusively for *in vitro* diagnostic purposes.
- · Intended for single use only. Do not reuse.
- Check the expiration date on the box. Do not use it after the expiration date.
- · Wear protective clothing, and use disposable gloves, goggles, and a mask.
- Do not eat, drink or smoke in areas where samples or test reagents are being used.
 Once you finish the test wash your hands.
- Specimens must be treated as potentially infectious as well as all reagents and materials that have been exposed to the samples and handled in the same manner as an infectious agent.
- Regular decontamination of commonly used equipment is recommended, especially micropipettes and work surfaces.
- This product contains irritants that are harmful when in contact with skin or eyes, or inhaled or swallowed. Care should be taken when handling this product.
- Some of the reagents in the 96-well Plate contain chaotropic which can form highly reactive compounds when combined with bleach. Do NOT add bleach or acidic

- solutions directly to the sample-preparation waste.
- Any significant incidents related to the product should be notified to the competent authorities and manufacturers.
- · Do not use it if the package is damaged.
- RNase can be introduced accidentally during RNA preparation. Various factors such as reagents, air, dust, and human hands or skin, can be the sources of RNase contamination. Always wear disposable gloves and use sterile, disposable plastic wares. Use the pipette reserved for RNA work. Maintain a separate area for RNA work. Carefully clean the surfaces.

COMPOSITION OF THE PRE-PACKED 96-WELL PLATE (W7070 | W7071)

A total of 16 samples can be simultaneously processed per plate.



COMPOSITION OF THE PRE-PACKED 6-WELL STRIP (W7072 | W7074)



- (#1) Sample
- (#2,3) Washing
- (#4) Magnetic bead
- (#5) Elution
- (#6) Washing

PROTOCOL

A. Setup of program (For CLEO™ AP16 & AP48 devices, preset program can be used.)

Edit and run the experiment program as follows:

No.	1	2	3	4	5	6	7	8
Well No.	4	1	2	3	4	6	5	4
Step	Bead	Bind	Wash	Wash	Wash	Wash	Elute	Discard
Wait time	-	-	-	-	-	-	03:00	-
Mix time	00:20	10:00	02:00	01:00	01:00	01:00	05:00	00:30
Collect time	00:25	00:30	00:25	00:25	00:25	00:25	00:30	-
Volume(µL)	750	800	750	750	750	750	100	750
Mixing speed	Medium	Fast	Fast	Fast	Fast	Fast	Bottom	Medium
Collect speed	Strong	Normal						
Temperature		Off	·				Off	

B. Sample Preparation

Required material: 1x PBS or DPBS

1. Cultured cells

- Pellet the cultured cells by centrifugation and resuspend the cell pellet in 200 µL of 1x PBS.
- Do not exceed 5 x 106 cells per prep.

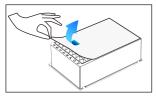
2. Body fluids

- Use up to 200 µL of specimen.
- Adjust to 200 µL with 1x PBS if less.

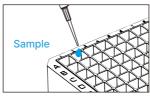
3. Swabs

- Scrape the swab to the specimen and place the swab into a 2 mL centrifuge tube by cutting out the stem part with sterile scissors.
- Apply 400 µL of 1x PBS into the tube and vortex vigorously for 15 seconds. The preserved swab sample in PBS or other commercial solution can be directly used. Use 200 µL of the preserved sample.

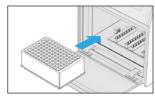
C-1. RNA extraction procedure (W7070, W7071)



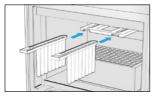
 Carefully peel off the film of the 96-well Plate not to cross-contaminate



2. Add 200 µL of the sample into the each first well (#1, 7)



3. Mount the 96-well Plate on the CLEO™ AP16 carefully.



4. Insert a Plunger all the way into the socket above the 96-well Plate.

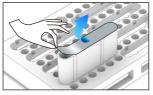


5. Close the front door of the instrument

C-2. RNA extraction procedure (W7072, W7073)



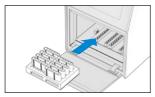
the Strip Adapter Plate.



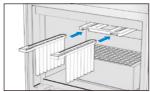
1. Mount the 6-well Strip onto 2. Carefully peel off the film of the 6-well Strip not to cross-contaminate.



3. Add 200 uL of the sample into the each first well (#1).



4. Mount the 6-well Strip Adapter Plate on the CLEO™ AP16 carefully.



5. Insert a Plunger all the way into the socket above the Plate

- 6. Close the front door of the instrument.
- 7. Select **MENU DNA Total RNA** on the screen.







- 8. Press 'RUN' button on the screen.
- 9. After the alarm finishes, open the door and carefully remove the Plunger.
- 10. Detach the 96-well Plate (or the Strip Adapter Plate) from the machine carefully.
- 11. Transfer the 70 90 µL RNA eluate of each fifth well (#5, 11) into a new 1.5 mL centrifuge tube. **NOTE**: The volume of eluate can be decreased slightly during the process.
- 12. Dispose of 96-well Plates (or 6-well Strip) and Plunger used in the test according to local or national waste disposal methods.

TROUBLE SHOOTING GUIDE

Problem	Possible causes	Recommendations		
Low recovery of RNA	Too much starting materials	Too much starting materials will bring about inefficient lysis, followed by poor RNA yields. Keep the maximum amount of starting material asdescribed in the procedure.		
	Poor quality of starting material	Use a freshly harvested sample if possible. Sample should always be handled quickly under low temperature.		
RNA degraded	Inappropriate handling of starting materials	The starting sample should be quickly treated under low temperatures. Long exposure to high temperatures or retarded processing would be a cause of degradation.		
	Poor quality of starting material	Refer to the same item in 'Low recovery of RNA' section.		
	RNase contamination	RNase can be introduced accidentally into a preparation at any step. Always wear disposable gloves and use RNase-free plasticwares. Do not use shared equipment if possible.		
DNA contamination	DNase not treated	This kit is designed to prepare RNA, but DNA can be co-purified depending on the sample. To remove DNA from the eluate, treat the eluate with DNase I.		
Inconsisten recovery of DNA	Contamination between reagent wells	The reagent in the well may evaporate and form a deposit on the film during storage, which may cause contamination between wells when the film is removed. Prepacked plate or tube always should be stored at proper condition. Before removing the film of the plate or the tube, it is recommended to shake off the deposit on the film with holding the plate or the tube tightly.		

SYMBOL GLOSSARY

REF	Catalogue number	—	Manufacturer	Ω	Use-by date
LOT	Batch code	2	Do not re-use	1	Temperature limitation
IVD	in-vitro diagnostic use	[]i	Instructions for use	漛	Keep away from sunlight
Σ	Contents sufficient for <n> tests</n>	<u> </u>	Caution	**	Keep dry
®	Do not use if package is damaged	UDI	Unique Device Identification		

ORDERING INFORMATION

Product	Cat No.	Package	Note		
	W7070	64 Prep	16 prop/rup		
 WizMag™ Total RNA	W7071	192 Prep	16 prep/run		
	W7072	32 Prep	Single prep		
	W7073	96 Prep	Sirigle prep		
CLEO™ AP16 Nucleic acid Extractor	CL2016	1 system	1-16 sample		
CLEO™ AP48 Nucleic acid Extractor	CL2048	1 system	1-48 sample		

MFDS License No.: IVD-23-843

WizMag[™] Total RNA 6



Technical Support



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