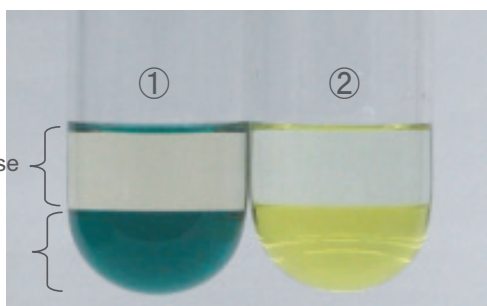


Ready-to-use Reagent for Total RNA Isolation

Sepasol-RNA I Super G

Sepasol-RNA I Super G is ready-to-use, green-phasic solution for isolating total RNA from biological samples such as cell or tissue etc. The green color of dye makes the separation of aqueous and phenol phases easier compared with RNA I Super included in yellow dye.

- » **Easy to use** - Ready-to-use green mono-phasic solution.
- Easy to identify interphase compared to Sepasol RNA I Super.
- » **Fast** - Isolation procedure are completed in less than 1hr.
- » **High purity** - The purified RNA is ready for use in standard downstream application such as RT-PCR.

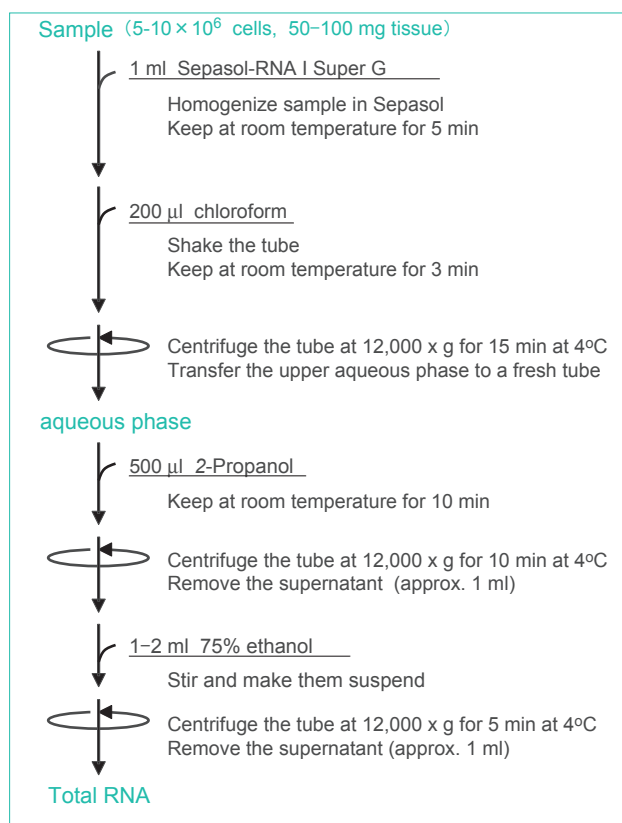


Aqueous phase
Phenol phase

① Sepasol-RNA I Super G
② Sepasol-RNA I Super

Protocol

Isolation procedure is completed in less than one hour.



Yield and purity of the isolated RNA



Data on isolated total RNA from HL-60 cell by Sepasol-RNA I Super G (Left) and by Sepasol-RNA I Super (Right). The isolation was from 5 × 10⁶ cell.

Product Name	Yield (mg)	Purity (A ₂₆₀ /A ₂₈₀)
Sepasol-RNA I Super G	32.4	2.08
Sepasol-RNA I Super	29.8	2.07

Ordering Information

Product Name	Storage	Product Number	PKG Size
Sepasol-RNA I Super G	4°C	09379-84	100 ml
		09379-97	200 ml
		09379-55	500 ml

For RNase Decontamination

RNase Quiet

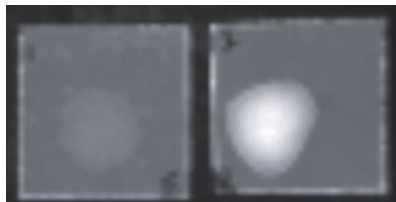


RNase Quiet is a ready-to-use solution for eliminating RNase contamination. It completely removes RNase contamination from glass, plastic equipments and laboratory tables.

- » Removes RNase contamination effectively
- » Easy to use spray type
- » Easy to wipe with no detergent
- » Non-carcinogenic with no DEPC

Decontamination of cover glass

DEPC treated water RNase Quiet

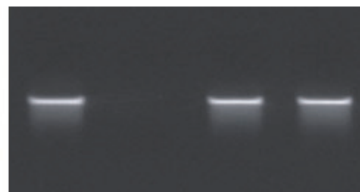


Condition

1. Apply 100 μ l RNase A solution (1mg/ml) on cover glasses and dry them.
2. Spray with DEPC treated water or RNase Quiet and wait for 1 minute. Wipe thoroughly with a clean paper towel and then rinse with RNase-free sterile water.
3. Apply 50 μ l RNA Solution (40 μ g/ml) on the cover glasses and incubate them at 37°C for 30 minutes.
4. Apply 1 μ l ethidium bromide solution (20 μ g/ml) on the cover glasses with a pipette.
5. Observe the cover glasses with UV.

Decontamination of 1.5 ml micro-tubes

DEPC treated water RNase Quiet
RNase A - + - +



Condition

1. (+): Add 10 μ l RNase A solution (10 mg/ml) in 1.5 ml micro-tubes.
(-): Add 10 μ l lysate buffer in 1.5 ml micro-tubes.
2. Add 1 ml DEPC treated water or RNase Quiet and wait for 1 minute. Remove the solution from the tube.
3. Add 25 μ l RNA Solution (40 μ g/ml) and incubate them at 37°C for 30 minutes.
4. Analyze RNA solution by electrophoresis with 1% agarose gel including 100 ng/ml ethidium bromide and then staining gel with 200 ng/ml ethidium bromide solution.

Ordering Information

Product Name	Storage	Product Number	PKG Size
RNase Quiet (with spray nozzle)	Room Temp.	09147-14	475 ml
RNase Quiet for Replacement	Room Temp.	09477-94	475 ml

Related Products

Ordering Information

Product Name	Storage	Product Number	PKG Size
100g/l-Hexadecyltrimethylammonium Bromide Solution, Nuclease tested	Room Temp.	17472-94	100 ml
Phenol:Chloroform 5:1 Mixed, pH4.5	4°C	26729-64 26729-06	100 ml 400 ml
Phenol:Chloroform:Isoamyl Alcohol 25:24:1 Mixed, pH5.2	4°C	26058-54 26058-96	100 ml 400 ml
Water, DEPC treated, RNase tested	Room Temp.	36420-61	10 x 1 ml
		36420-74	50 x 1 ml
		36415-54 36415-41	100 ml 1 L
Hexadecyltrimethylammonium Bromide	Cool and Dark	08897-82	25 g
Guanidine Thiocyanate	Room Temp.	06287-32	25 g
		06287-45	500 g
Ethanol(99.5)	Room Temp.	08948-54	100 ml
		08948-25	500 ml

For research use only, not intended for diagnostic or drug use.

NACALAI TESQUE, INC.

Nijo Karasuma, Nakagyo-ku, Kyoto 604-0855 JAPAN

TEL : +81-(0)75-251-1730

FAX : +81-(0)75-251-1763

Website : www.nacalai.com

E-mail : info.intl@nacalai.com