





A Geno Technology, Inc. (USA) brand name

Arrest[™] Extraction Buffer

(Cat. #786-133)



INTRODUCTION

Arrest[™] Extraction Buffer is an optimized combination of various chaotropic agents and RNase inhibitors, which inhibits RNase in 5-10 minutes. Arrest[™] Extraction Buffer may be used in conjunction with any RNA extraction method, including extractions based on phenol, chloroform and other organic solvents and detergents. A quick 10 minutes single step protocol isolates high quality, total RNA from any tissue and is conveniently packed as two 50ml bottles. Enough reagents are provided to isolate RNA from a total of 10 grams of tissues.

ITEM(S) SUPPLIED (Cat# 786-133)

Description	Size
Arrest [™] Extraction Buffer	100ml

STORAGE CONDITION

It is shipped at ambient temperature. Upon arrival, store Refrigerated at 4⁰C. When stored and used properly the reagent is stable for 1 year.

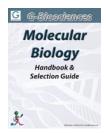
PROTOCOL

Preparation before Use:

- 1. Add 0.20ml beta-mercaptoethanol into the Arrest $^{\infty}$ Extraction Buffer. Prepare only one bottle at a time and place the date of use on the bottle label.
- Use 0.5ml Arrest[™] Extraction Buffer for each 10-50mg tissue. For cell culture, add 0.5ml per 1-2 million cells.
- Use any grinding method. After grinding, add 0.5ml chloroform and invert the tube several times to mix.
- 4. Centrifuge and collect aqueous phase supernatant for extraction of RNA. Take care not to disturb the debris in the inter-phase.
- 5. Add equal volume ice-cold isopropanol to precipitate RNA. Mix and incubate on ice for 5 minutes. Centrifuge 5 minutes at 4°C to collect the RNA pellet.
- 6. Wash the RNA pellet with 1ml ice-cold 70% ethanol, and centrifuge 3 minutes to collect the RNA pellet. Remove ethanol and invert the tube on a clean absorbent tissue for several minutes to allow the excess ethanol to drain away.
- Rehydrate RNA pellet with a minimal volume (2-40ul) of water or other buffer.
 <u>NOTE</u>: After grinding the tissue in Arrest[™] Extraction Buffer, the protocol can be modified for extraction of RNA, such as using different combination of chloroform and other organic solvents.

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