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RNA Isolation from Plant Tissue Protocol 11: Tri Reagent Method

1 Works for me dx.doi.org/10.17504/protocols.io.4ydgxs6



Scott C. Edmunds
GigaScience/BGI Hong Kong/Bauhinia Genome



ABSTRACT

Implemented by: Michael Melkonian and Barbara Surek

The protocol for RNA isolation using [TRI Reagent](#) (Molecular Research Center, Cincinnati OH) is analogous to the TRIzol LS Reagent protocol except that 1 ml of TRI Reagent is added to each 250 µl of homogenized material (= approx. 50–100 mg packed cell volume) (step 3, [Protocol 10](#)).

For background, TRI Reagent combines phenol and guanidine thiocyanate in a monophasic solution to inhibit RNase activity during RNA isolation.

This protocol is part of a collection of eighteen protocols used to isolate total RNA from plant tissue. (RNA Isolation from Plant Tissue Collection: <https://www.protocols.io/view/rna-isolation-from-plant-tissue-439gyr6>)

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GUIDELINES

In cases of high RNA concentration (> 1000 ng/µl), but low 260/230 ratio (less than 1), RNA can be extracted and precipitated again.

For this, steps 8–13 of the TRIzol LS Reagent Method are repeated.

Protocol 10: TRIzol LS Reagent Method
by Julia Roßmanith

PREVIEW

RUN

The RNA phase is mixed with 500 µl 8 M LiCl by vortexing and incubating for 01:00:00 at 4 °C. Subsequently, steps 16–23 of the TRIzol LS Reagent protocol are performed.

Frequently, repetition of extraction decreases previous RNA concentrations up to tenfold, but increases the 260/230 ratio remarkably attaining values of 1.8 or more.

Neither using isopropanol for the second precipitation process nor replacing isopropanol by LiCl as a main precipitator is as efficient as the method described above.

MATERIALS

NAME ▾

[TRI Reagent®](#)

CATALOG # ▾

[View](#)

VENDOR ▾

[Molecular Research Center, Inc.](#)

SAFETY WARNINGS

Please see SDS (Safety Data Sheet) for hazards and safety warnings.



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