Nov 13, 2018

DNA extraction from whatman filter papers

DOI
dx.doi.org/10.17504/protocols.io.vjme4k6

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OPEN ACCESS

DOI: dx.doi.org/10.17504/protocols.io.vjme4k6

Protocol Citation: Xiang Guo, Xiaoqing Zhang, Xiaohong Zhou 2018. DNA extraction from whatman filter papers. **protocols.io** **https://dx.doi.org/10.17504/protocols.io.vjme4k6**

Manuscript citation:

Protocol status: Working
We use this protocol and it's working

Created: November 12, 2018

Last Modified: November 13, 2018

Protocol Integer ID: 17741

Keywords: DNA extraction, whatman filter papers

Abstract

This protocols is for RNA extraction from Whatman filter paper.

Materials

STEP MATERIALS

Chelex 100 **Merck MilliporeSigma (Sigma-Aldrich)** **Catalog #**C7901-100G

DNAse/RNAse free distilled water **Thermo Fisher Scientific** **Catalog #**10977023

Protocol materials

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Lyse RBC

1 Filter paper disc containing blood cut and placed into 1.5 mL EP tubes.

2 Add 1 mL RNase-free water and vortex to mix.

DNAse/RNAse free distilled water **Thermo Fisher Scientific Catalog #**10977023

3 Incubate at 25 °C Room temperature for 00:15:00 to lyse RBC.

Precipitate the DNA and separate phases

4 Centrifuge at 20,000 × g for 00:05:00, discard the supernatant with a micropipettor.
go to step #2 undefined until the DNA precipitate forms a white or pink gel-like pellet at the bottom of the tube.

5 Resuspend the pellet in 150 µL 10% Chelex-100 solution , vortex the sample briefly for 00:00:30 .

Chelex 100 **Merck MilliporeSigma (Sigma-Aldrich) Catalog #**C7901-100G

6 Incubate the sample at 100 °C for 00:10:00 .

7 Centrifuge for 00:05:00 at 20,000 × g.

8 Carefully transfer the supernatant to a clean 1.5 mL EP tube with a micropipettor. Proceed to downstream applications, or store the DNA at -20 °C .

Determine the DNA yield

9 Measure absorbance at 230nm, 260nm, and 280nm, Calculate the A260/A280 and A230/A260 ratio.