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Total nucleic acid extraction - NucleoMag® DNA/RNA Water Kit (Macherey-Nagel)

 Forked from [Total nucleic acid extraction - Maxwell\(R\) HT Environmental TNA Kit, custom \(Promega\)](#).

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Protocol status: Working

We use this protocol and it's working

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Abstract

Total nucleic acid extraction from wastewater using NucleoMag® DNA/RNA Water Kit (Macherey-Nagel)

Guidelines

When work is completed, remove equipment and supplies from the cabinet. Wipe the work area with 10% bleach, let stand for 10 min, rinse with water, then with 70% ethanol, and finally with RNAase AWAY.

Materials

MATERIALS

 NucleoMag® DNA/RNA Water Kit **Macherey-Nagal Catalog #744220.1**

- Molecular biology grade water
- 6x KingFisher 96-well plates** (Cat. no.: 95040460)
- 1x KingFisher 96-tip comb well plate**(Cat. no.: 97002534)
- Screw cap** microcentrifuge tubes

Equipment

Mini-Beadbeater-16	NAME
high-energy cell disrupter	TYPE
BioSpec	BRAND
607	SKU
https://biospec.com/product/mini-beadbeater-16	LINK
1 speed	SPECIFICATIONS

Equipment

Kingfisher Flex	NAME
Automated Extraction System	TYPE
ThermoFisher	BRAND
5400630	SKU

 Promega_Maxwell_HT_RNA_Waste...

 MN_96_Flex.bdz



Before start

1. Clean the working area and all equipment: wipe down with 10% bleach and let dry. Wipe down with 70% ethanol and let dry. Then, wipe down using RNase AWAY and let dry.
2. Prepare the 50% ethanol solution (it must be fresh!)
3. Prepare the 6 purification plates:
 - **Wash 1 plate:** Add 100 μ l of 50% ethanol and 900 μ l of wash buffer (WBA) to each well required for purification.
 - **Wash 2 plate** (same as plate Wash 1): Add 100 μ l of 50% ethanol and 900 μ l of wash buffer (WBA) to each well required for purification.
 - **Ethanol Wash plate:** Add 450 μ l of 50% ethanol to each well required for purification.
 - **Elution plate:** Add 100 μ l of 25 mM Tris-HCl (pH 8.0) to each well required for purification.
 - **Tip plate:** Place KingFisher 96-tip comb into an empty KingFisher 96-well plate. *While opening the 96-tip comb plate, pay attention to not touch the tips.*
 - **Lysis and Bind plate:** Add 35 μ L of Resin to each well required for purification (*vortex the bottle at max speed before use*). Add 50 μ l of Alkaline Protease Solution custom (APA) to each well required for purification. Add 250 μ l of cell lysis solution (CLD) to each well required for purification. Add 400 μ l of Isopropanol (100%) to each well required for purification.

Total nucleic acid extraction

2h

- 1 For **HA filter** extraction, let the sample thaw on ice and go to **step 2**.

5m

For **BCoV/BRSV** extraction (in duplicate), add 5 µL of BCoV/BRSV solution to the 2-mL tube containing 500 µL MWA1. Vortex for 15 seconds (speed 7 out of 10) and flash freeze the tube. Go to step 4.

For **Direct extraction**, add 150 µL of wastewater to the 2-mL tube containing 100 µL CTAB. Vortex for 15 seconds (speed 7 out of 10) and flash freeze the tube. Go to step 4.

- 2

For **HA filter** extraction, place the 2-mL tubes in the bead beater.

Equipment

Mini-Beadbeater-16

NAME

high-energy cell disrupter

TYPE

BioSpec

BRAND

607

SKU

<https://biospec.com/product/mini-beadbeater-16>

LINK

1 speed

SPECIFICATIONS

- 2.1 Bead beat for  00:02:30

2m 30s

Safety information

Start the bead beating when the beads start to be loose in the tubes.

- 2.2 Cooldown the samples on ice for  00:05:00 .

5m



2.3 Repeat Steps 9.1 and 9.2 once  . 7m 30s

3 Centrifuge at maximum speed for 1 min at room temperature. 1m
 150000 rpm, Room temperature, 00:01:00

4 For **HA filter** extraction, transfer 400 µL of supernatant to the **Lysis/Bind plate**. 10m
For **BCoV/BRSV/Direct extraction**, transfer all supernatant to the **Lysis/Bind plate**.

5 Start the protocol MN_96_Flex.bdz on the KingFisher Flex  01:14:00 1h 14m

Equipment

Kingfisher Flex	NAME
Automated Extraction System	TYPE
ThermoFisher	BRAND
5400630	SKU

6 Transfer the purified sample from the **Elution plate** to the **microcentrifuge tubes**. 10m

Note

The DNA/RNA is now ready for downstream applications. RNA extract may be stored in RNase-free water at -80°C for 1 year.

RT-ddPCR

7 Quantification by Droplet Digital PCR (ddPCR)



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