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We use this protocol and it's working

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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)

Adélaïde Roguet[1], Melissa Schussman[2]

[1]UWM; [2]University of Wisconsin - Milwaukee

mclellan lab

Melissa Schussman
University of Wisconsin - Milwaukee

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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.

Keywords: wastewater, SARS-CoV-2, total nucleic extraction, RNA, DNA, extraction, purification, Promega, KingFisher

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_Wastewater_V1.bdz MN_96_Flex.bdz

BEFORE START INSTRUCTIONS

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 µl of 50% ethanol and 900 µl of wash buffer (WBA) to each well required for purification.
- **Wash 2 plate** (same as plate Wash 1): Add 100 µl of 50% ethanol and 900 µl of wash buffer (WBA) to each well required for purification.
- **Ethanol Wash plate**: Add 450 µl of 50% ethanol to each well required for purification.
- **Elution plate**: Add 100 µl 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 µL of Resin to each well required for purification (v*ortex the bottle at max speed before use)*. Add 50 µl of Alkaline Protease Solution custom (APA) to each well required for purification. Add 250 µl of cell lysis solution (CLD) to each well required for purification. Add 400 µl 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
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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 -80ºC for 1 year.

RT-ddPCR

7 Quantification by Droplet Digital PCR (ddPCR)

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