



P Opentrons COVID-19 testing (Zymo, station B, 48 samples)

Forked from Opentrons COVID-19 testing (Randox/qPCR common path, stations A & B, 24 samples)

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Opentrons COVID-19 Testing



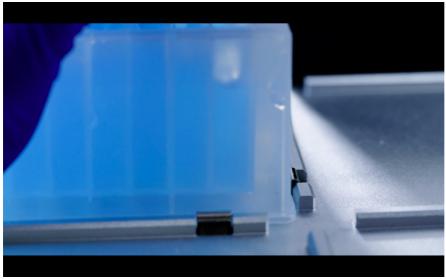
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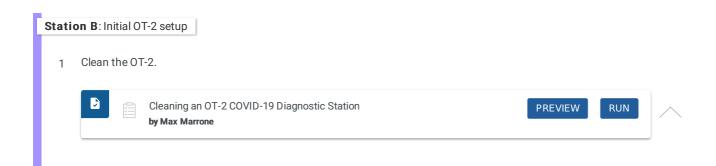
BEFORE STARTING

When placing labware on the OT-2's deck:

- 1. Make sure the labware is properly inserted by pressing the corner into the metal springs (pictured below). You should feel a slight click, and the labware should sit completely flat.
- 2. Make sure the labware is inserted the right way around, with well A1 at the top left.



Closeup of gloved hand pressing reagent trough into the labware holding springs



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- 1.1 Wipe these parts of the OT-2 down with a 1:10 dilution of bleach:
 - 1. The clear polycarbonate windows.
 - 2. The black pipette stems. (Avoid the rest of the pipettes, including the ejectors.)
 - 3. The aluminum deck.
 - 4. The removable black trash bin.
- 1.2 Wait © 00:00:30, then quickly rinse the bleach off with distilled water.



The aluminum on the OT-2 will be discolored if the bleach sits for too long. In the long term, it may also cause more serious corrosion.

1.3 Wipe these parts of the OT-2 down with RNaseZap or RNase AWAY.

The same parts that you wiped down with bleach:

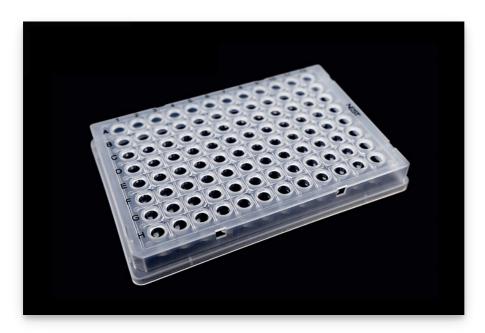
- 1. The clear polycarbonate windows.
- 2. The black pipette stems. (Avoid the rest of the pipettes, including the ejectors.)
- 3. The aluminum deck.
- 4. The removable black trash bin.

Plus these additional parts:

- 1. The bottoms of the pipette ejectors.
- 2. Any Temperature Modules or Magnetic Modules that the OT-2 has on its deck.
- 3. Any 96 well aluminum blocks that are going to be used on the OT-2.
- 1.4 Rinse the RNaseZap or RNase AWAY off with distilled water.
- 1.5 Wipe the OT-2 dry, or let the water evaporate.
 - 2 Place the following labware on the OT-2's deck:

Slot 1: An Opentrons Temperature Module with an Opentrons 96 well aluminum block and an empty, sterile NEST 100 μL PCR plate.





Slots 3, 6, 7, 9, and 10: A full, sterile rack of Opentrons 200 μ L filter tips.



Slot 4: An Opentrons Magnetic Module with nothing on it.



Slot 11: An empty NEST 1-well reservoir, for organic liquid waste.



3 Start pre-cooling the Temperature Module to § 6 °C.

4 Prepare the sample well plate.

The sample well plate is a NEST 96 deep well plate.



The 48 samples (#1-#48) should be laid out like this:

	Col. 1	Col. 2	Col. 3	Col. 4	Col. 5	Col. 6	Col. 7	Col. 8	Col. 9	Col. 10	Col. 11	Col. 12
Row A	#1	empty	#9	empty	#17	empty	#25	empty	#33	empty	#41	empty
Row B	#2	empty	#10	empty	#18	empty	#26	empty	#34	empty	#42	empty
Row C	#3	empty	#11	empty	#19	empty	#27	empty	#35	empty	#43	empty
Row D	#4	empty	#12	empty	#20	empty	#28	empty	#36	empty	#44	empty
Row E	#5	empty	#13	empty	#21	empty	#29	empty	#37	empty	#45	empty
Row F	#6	empty	#14	empty	#22	empty	#30	empty	#38	empty	#46	empty
Row G	#7	empty	#15	empty	#23	empty	#31	empty	#39	empty	#47	empty
Row H	#8	empty	#16	empty	#24	empty	#32	empty	#40	empty	#48	empty

For each sample well...



P

Add $\blacksquare 4 \mu l$ Proteinase K .



4.3 X

Station B: Reagent preparation

5 Prepare 3x **□14.35 ml Viral DNA/RNA Buffer + MagBinding Beads** ...



Vortex the **MagBinding Beads** in their container.



- 5.4 Repeat for a total of 3x 14.35 ml Viral DNA/RNA Buffer + MagBinding Beads.
- 6 (II)
 Wait for the Temperature Module to reach § 6 °C.

7 Prepare the reagent reservoir.



This step is time-sensitive. The MagBinding Beads will slowly settle in the Viral DNA/RNA Buffer. Continue quickly until you start the run on the OT-2. Vortex the Viral DNA/RNA Buffer + MagBinding Beads immediately before adding them to the reservoir.

Into a 15 mL NEST 12 well reservoir, pipette or pour the following reagents:

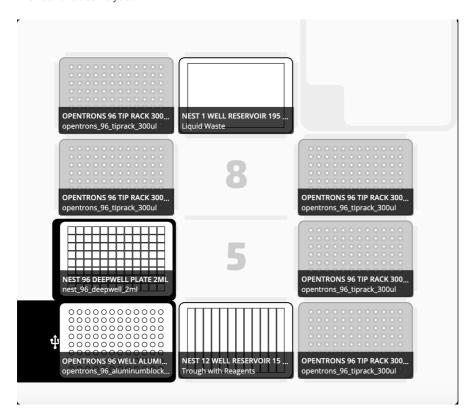
- Well 1: 14.35 ml Viral DNA/RNA Buffer + MagBinding Beads
- Well 2: ■14.35 ml Viral DNA/RNA Buffer + MagBinding Beads
- Well 3: ■14.35 ml Viral DNA/RNA Buffer + MagBinding Beads
- Well 4: ■13 ml MagBead DNA/RNA Wash 1
- Well 5: ■13 ml MagBead DNA/RNA Wash 1
- Well 6: ■13 ml MagBead DNA/RNA Wash 2
- Well 7: 13 ml MagBead DNA/RNA Wash 2
- Well 8: **13** ml 95-100% ethanol
- Well 9: **13** ml 95-100% ethanol
- Well 10: **13 ml** 95-100% ethanol
- Well 11: **13 ml** 95-100% ethanol
- Well 12: **3** ml nuclease-free water



A 15 mL NEST 12 well reservoir.

Station B: Final OT-2 setup

- 8 Place the sample well plate in slot 4.
- 9 Place the reagent reservoir in slot 2.
- 10 Double-check all the labware to make sure it looks correct.
 - Check that labware are inserted the right way around (well A1 at the top-left for the plates, well 1 at the left for the troughs).
 - Check that labware are properly clicked into the deck slots.
 - Check the deck layout:





- 11.2 Ensure you are connected to the robot. In the **Robots** tab, you can try flipping the robot's lights on and off to test the connection.
- 11.3 Go to the **Run** tab.

Double-check the name at the top to make sure the correct protocol is uploaded.

11.4 Click Start run. The OT-2 will home its motors and then begin the protocol.



Do not click **Start run** more than once. If you do, a known bug will make the OT-2 run the protocol back-to-back.



If something goes wrong and you need to abort the protocol:

- 1. Shut down the OT-2 with the power switch on its back left side.
- 2. Turn the OT-2 back on. Wait a couple of minutes for the pipettes to rise.
- 3. Manually remove any tips attached to the pipettes. (This ensures that the pipettes will not aspirate liquid into themselves when they home.)
- 4. Reconnect to the OT-2 in the Opentrons App. Click the Home button to move the gantry out of the way so you can access the labware on the deck.
- 12 Wait for the run to finish. It should take 2.5–3 hours.
- 13 The output is the **NEST 100 µL PCR plate** sitting on the aluminum block atop the Temperature Module in **slot 1**.

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