

RAPID USER GUIDE

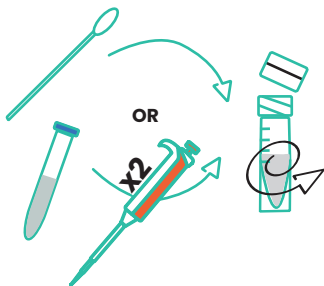


Switch on the analyser and the heating blocks.



Centrifuge the tubes before starting the procedure.

1



Swab: Immerse and rotate the swab for about ten seconds in the buffer tube before wringing it out against the wall of the large tube and vortexing.

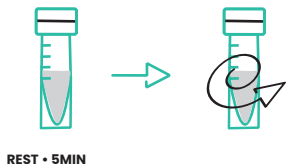
BAL : Withdraw **two 25µL volumes** of BAL using the orange pipette fitted with a filter cone. Place this volume in the large buffer tube and vortex.

2



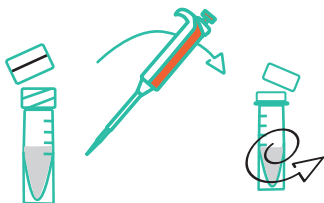
Place the buffer tube in the 95°C incubator for 10 minutes.

3



Leave the tube to stand for 5 minutes at room temperature, then vortex.

4



Using the orange pipette fitted with a filter tip, withdraw 25 μ L from the buffer tube.

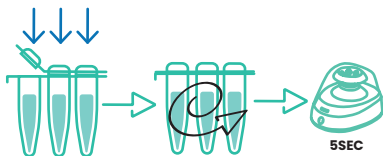
Place this volume in the white buffer tube and vortex.

5



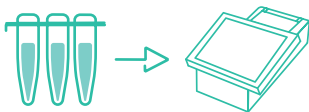
Using the orange pipette fitted with a filter cone, add 25 μ L from the white buffer tube to each minitube.

6



Close the minitubes tightly, vortex them and then place them in the centrifuge for 5 seconds.

7



Insert the minitubes into the instrument (place the end tab on the left) then immediately close the lid of the instrument and press ok to start the analysis.



Never reopen the minitubes once they have been inserted in the instrument, or after the end of the reaction.

Never leave the minitubes inside the instrument after the reaction has ended.

Refer to the user manual for more detailed information:

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