

1. Ensure the SD card is properly plugged in the socket (Fig. 1 and 2).

*SD card needs to be plugged-in before power on. The SD card CAN NOT be initialized with hot plugging.

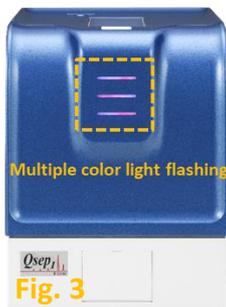


2. Connect the air tube.

3. Plug the power cord (labeled *Qsep₁*)

4. Switch on *Qsep₁*, the LED lights (multiple color light) (Fig. 3). Wait 25 seconds for the instrument and Wi-Fi initialization, and the green LED will start flashing (Fig. 4).

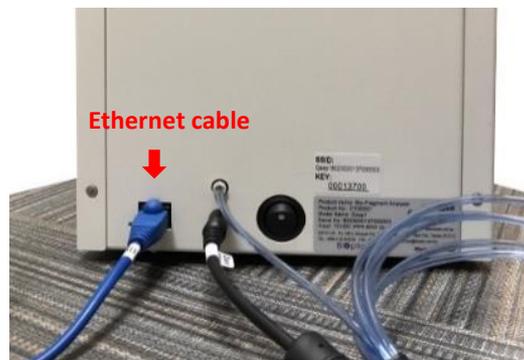
*DO NOT switch on the instrument immediately after powering off. Wait at least 5 seconds



*The green flashing LED shows the instrument is successfully ready for connection. If not, please repeat the step 4.

Ethernet cable connection:

refer to step 5 on Page 2



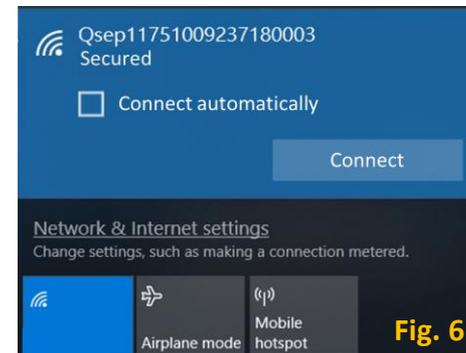
Wi-Fi connection: follow the steps below to connect with *Qsep₁*



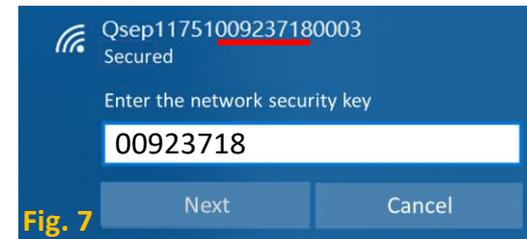
- Check if the AP source is available from your Wi-Fi network in the computer (Fig. 5).



- Find the SSID named with “*Qsep₁*+instrument ID (16 digits)” and click “Connect” (Fig. 6)



- Enter the password:
The password will show in the sticker which is behind the instrument. It is the same as the **middle 8 digits** of instrument serial number (same as SSID name) by blinding the prefixal “*Qsep₁*” (Fig. 7).



*If your computer can't find the SSID of *Qsep₁*, please keep the distance between *Qsep₁* and the computer within 10 meters and try again from the step 4.

*If you can not connect with *Qsep₁*

- Please check the wi-fi IP assignment is “Automatic (DHCP)”
- Allow Q-Analyzer pass through firewall or turn off the firewall

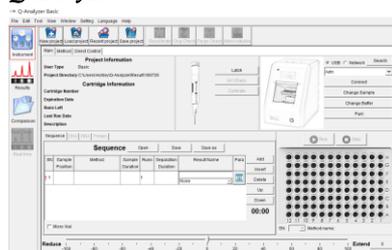
5. Check the LED status.



6. Double click the Q-Analyzer.



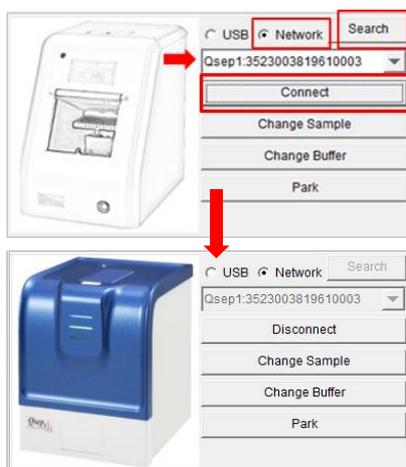
7. Q-Analyzer user interface:



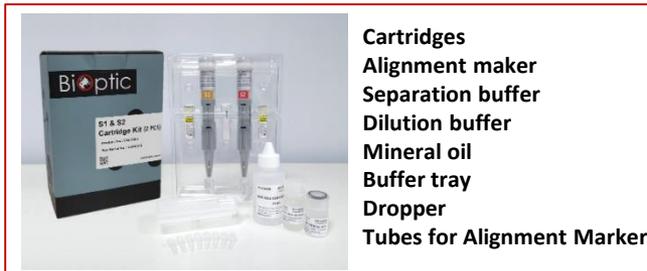
7-1. Select "Network" mode and click "Search"
*After "Search" the instrument ID will display

*Please confirm the instrument ID is the connected instrument

7-2. Click "Connect", the image will switch to Qsep₁ when it's connected

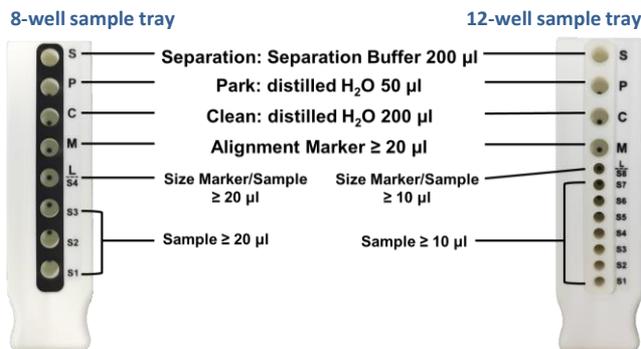


Packing List of Cartridge Kit (Cat. C105200)



- Cartridges
- Alignment maker
- Separation buffer
- Dilution buffer
- Mineral oil
- Buffer tray
- Dropper
- Tubes for Alignment Marker

8. Prepare buffer and alignment marker in 0.2ml PCR tubes and allocate at the corresponding position on the sample tray



*For 12-well sample tray, user needs to use 0.1ml tube (C104252) for sample and size marker

*S/P/C well can only place 0.2ml PCR tubes

*Make sure no air bubble appears in each tube

9. Click "Change Sample" or "Change Buffer", the sample door will open automatically.

10. Slide the buffer tray into the instrument. Make sure sample tray is pushed to the end



11. Click "Park", the sample tray move into instrument

12. Open the cartridge door by pressing the white button and insert cartridge (guiding groove facing front).

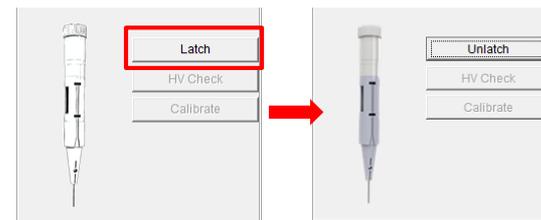
*Please follow cartridge unpacking guide to unpack the cartridge before using



Guiding groove



13. Close the cartridge door and click "Latch".



*The cartridge information will be displayed after latched.

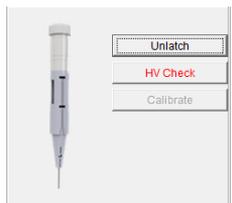
Project Information	
User Type	Basic
Project Directory	C:\Users\Ashley\Q-Analyzer\Results\190726
Cartridge Information	
Cartridge Number	S1-O-190712-4
Expiration Date	2020-Jan-08
Runs Left	37
Last Run Date	2019-Jul-25
Description	High resolution

14. Cartridge Calibration:

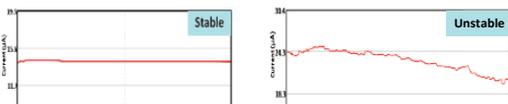
New cartridge needS to be calibrated before using.
Please proceed with the following steps.



14-1. Click "HV Check".



*** The storage and transportation condition may influence the Gel-matrix and cause unstable current. During HV Check, check current (gray line) and see if it's stable. if current is unstable, please repeat this step 2-3 times.**

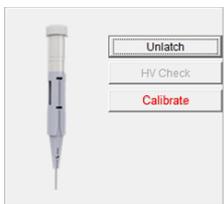


During the "HV Check", the last LED will show red light

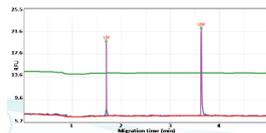
Blue → SD card
Green → WiFi connectable
Red → HV check



14-2. Click "Calibrate".



***The "Calibrate" button will only display after HV check passed.**

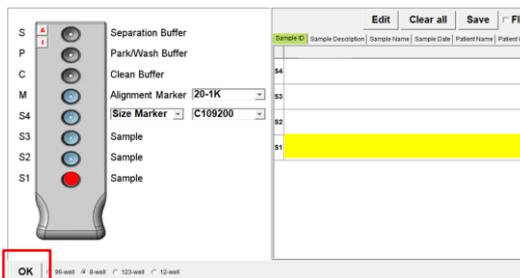


*** For troubleshooting, please refer to cartridge unpacking guide for details.**

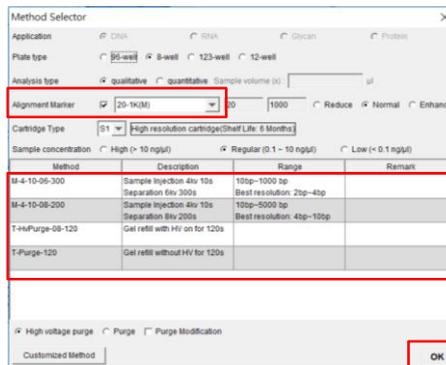
15. To analyze samples, click on the blank column and designate the sample locations, test method, runs and result name followed steps 15-1 to 15-4

SN	Sample Position	Method	Sample Duration	Runs	Separation Duration	Result Name	Para	Add
1				1		None		Insert Delete Up Down 00:00

15-1. Click the "Sample Position", mark the position of sample on the plate and then click "OK".



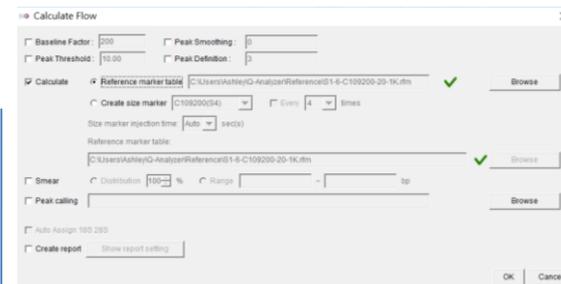
15-2. Click on "Method" to select a proper method for your application.



15-3. Click on "Result Name" and enter the name of result files.

SN	Sample Position	Method	Sample Duration	Runs	Separation Duration	Result Name	Para
1	S1	M-4-10-06-300	10	1	300	Test None	

15-4. Click the icon "Para" and set the parameters. (Baseline Factor, Peak Threshold, Calculate...etc.).



16. Click "Run" to start the process.



During Calibration

Please check the Alignment Marker has been located at correct position. Software will recognize two peaks from Alignment Marker signal. **DO NOT** use Size Marker or DNA sample doing "Calibrate".

Brief introduction of the Signal Chart:

