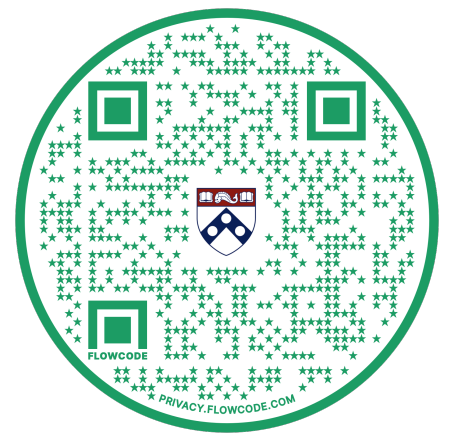
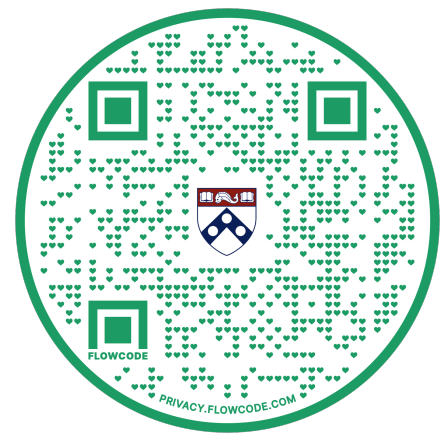


\*Samples should be freshly filtered. After 2 hrs, small clumps/floaties can occur.



No/few events displaying when acquiring

1

2

Tube cracked?

Use new (uncracked) tube.

Is your tube Falcon polystyrene (12x75mm)?

Transfer all specimens to polystyrene tubes. (Polypropylene tubes will not work)

Did you fill sheath/empty waste?

Fill sheath/empty waste. Purge filter of air.

Is there a FFSS (fluidics cart)?

Are your samples filtered AND free of floaties?\*

Filter samples.

Run 2% contrad with arm over for 30 secs. Then, put arm under and run on high for 2 mins. Repeat with diH2O. Prime.

Is it pumping?

Ensure FFSS is ON (green button). Check plenum: if empty, hold down prime button on fluidics cart for 1 min. Purge filter of air.

Is RUN button green with 0 events displaying while acquiring?

Contact the core at 215-898-3528 or try "More Tips" QR code (above)

Turn off cytometer and restart computer. Turn cytometer/lasers back on.

Not fixed?

No

Yes

Not fixed?

See video

See video