

BD Accuri™ C6

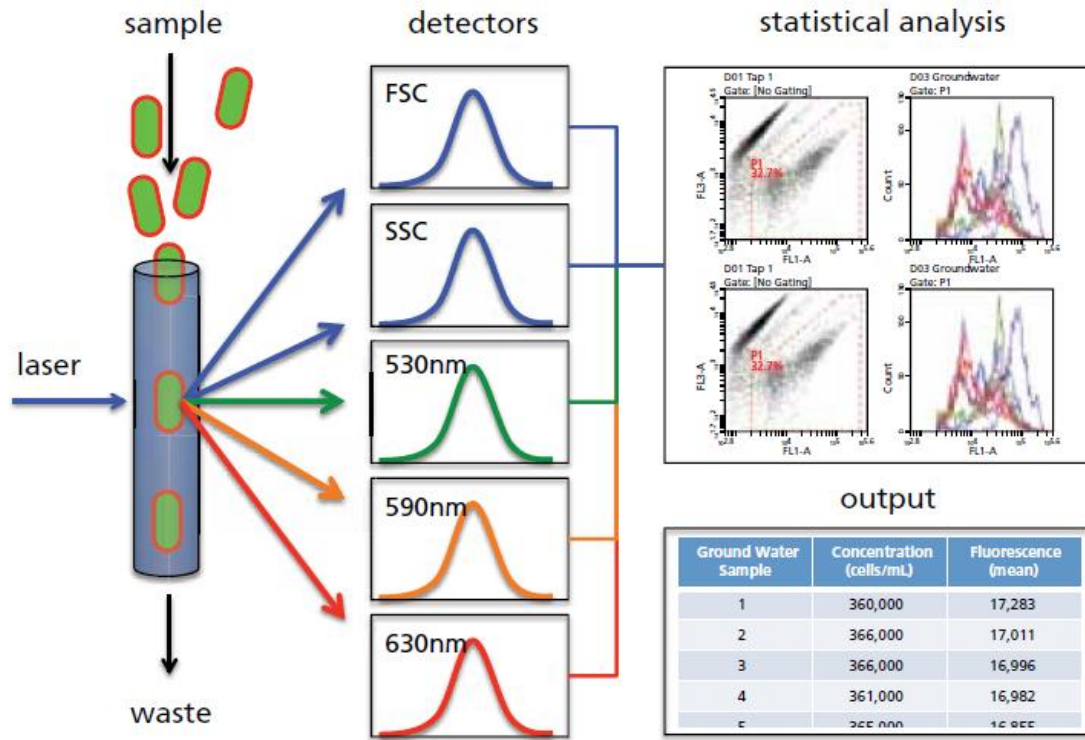
Flow cytometry within reach.™



Helping all people
live healthy lives

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V1.0 July 2012

How it works



BD Accuri™ C6

- Affordable, easy- to- use Flow Cytometer
- 2 lasers and 6 detectors
- A flow cytometer for every lab

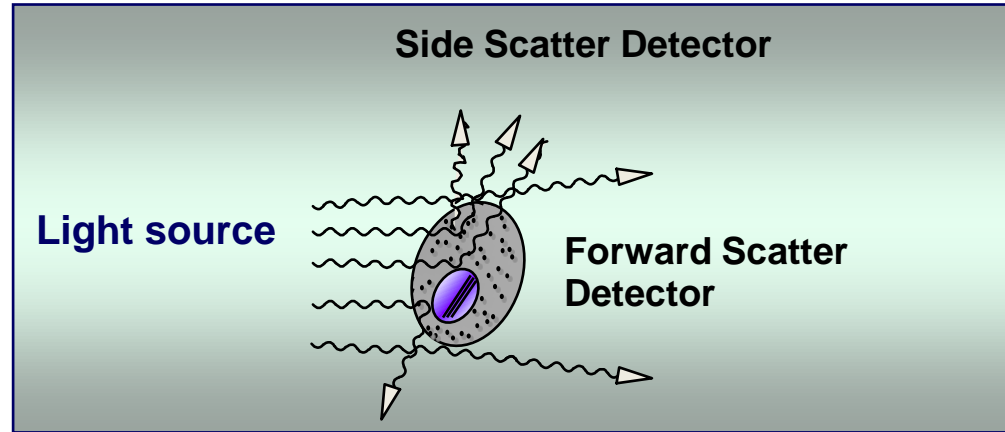


Flow Cytometric measurements

- Relative size (Forward Scatter-FSC)
- Relative internal complexity (Side Scatter-SSC)
- Relative fluorescence intensity

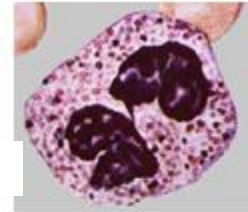
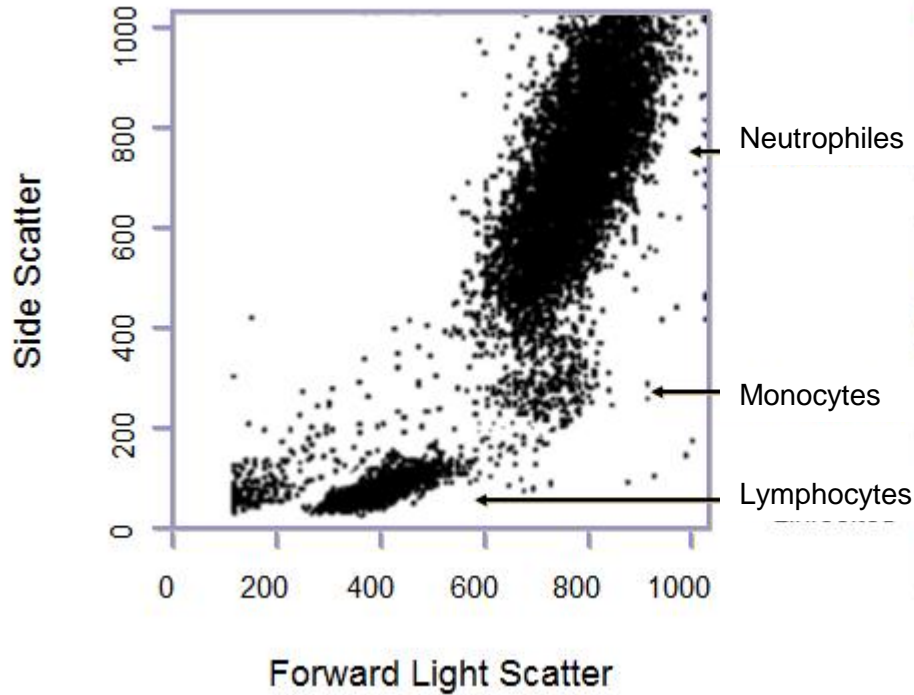
Introduction to Flow Cytometry

Size and granularity



- FSC:
Related to cellular size
- SSC:
Related to internal complexity
Measured at 90°

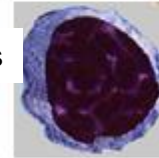
Introduction to Flow Cytometry



Neutrophils

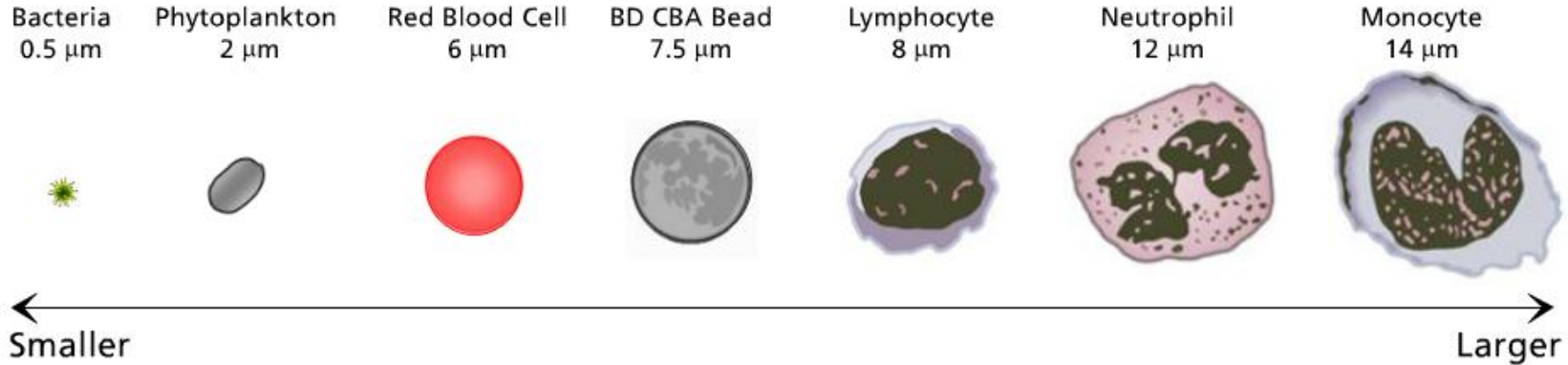


Monocytes



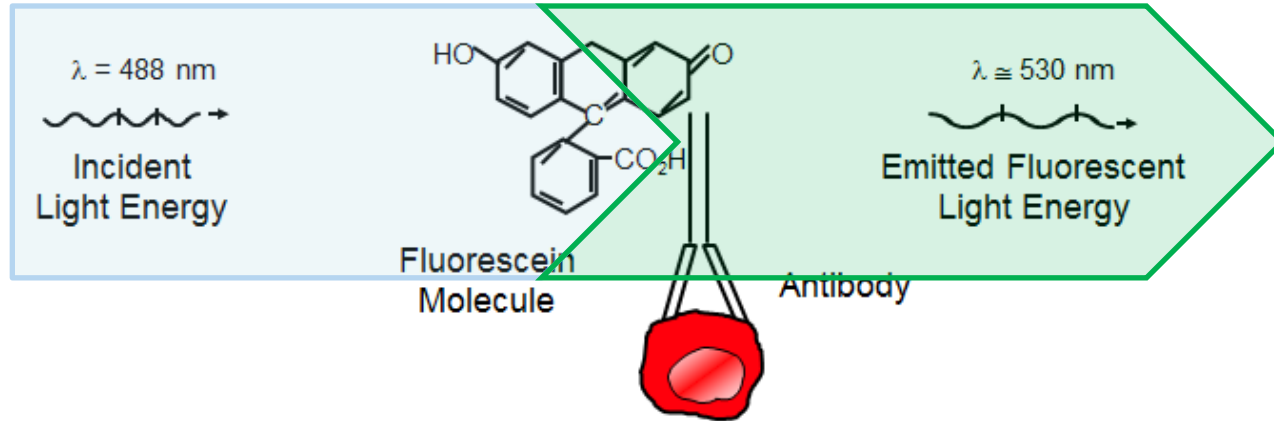
Lymphocytes

What can we detect by Flow Cytometry?



Introduction to Flow Cytometry

- Fluorescence

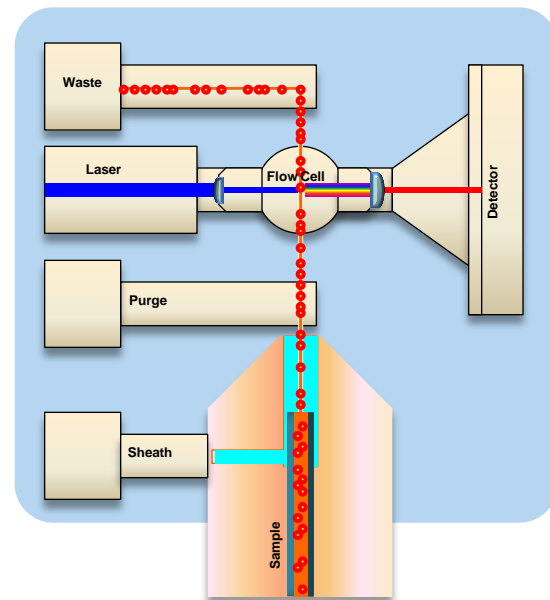


Introduction to Flow Cytometry

- Flow cytometer needs a combined system including:
 - Fluidics: Introduce and focus the sample in the flow cell
 - Optics: Illuminate the cells and direct the resulting light and fluorescence to the detectors
 - Electronics: Convert light to electronic signals that can be analysed in a computer

Fluidics

- Non-pressurized, peristaltic pump-driven system
- User adjustable flow rate and core diameter for a variety of applications
- Volume reported for absolute counts
- Minimum sample volume 50 μL
- Up to 10,000 events/second



Optical System

- Excitation Optics
 - Lasers
- Collection Optics:
 - Filters that guide signal wavelengths to a specific detector

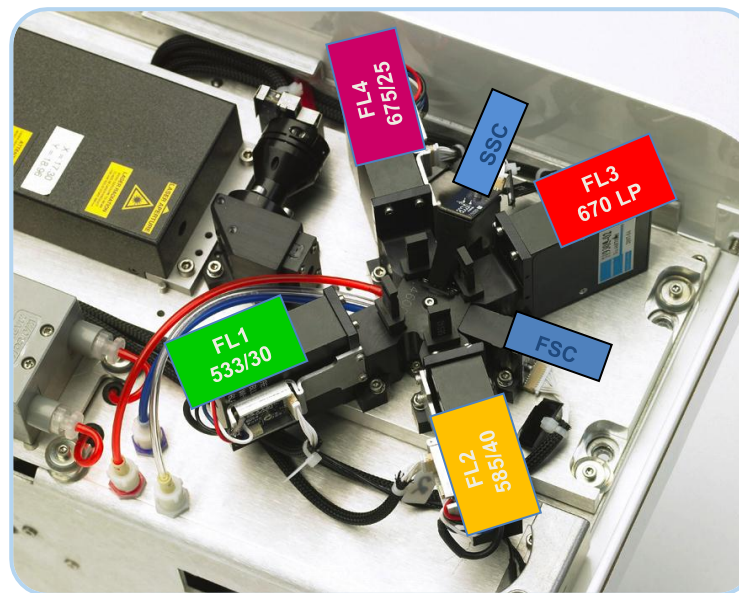
Optical system

488 nm
solid state laser

640 nm diode laser

PMTs for fluorescence
detection

Optical path



BD Accuri™ C6

The BD Accuri™ C6 is Portable!



2nd Norwich Flow Day:
Institute for Food Research



The Ecosystem Centre:
Research vessel, Palmer Peninsula, Antarctica

FLUIDICS

OPTICS

SOFTWARE

APPLICATIONS

RESOURCES



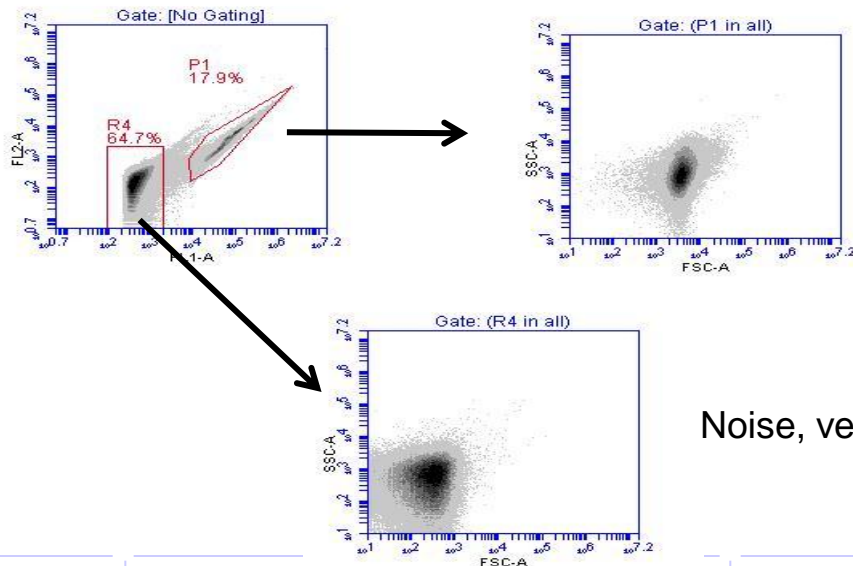


Applications

E. Coli in PBS + Syto BC

PBS spiked with E. Coli, plus Syto BC dye

Trigger on FL1, channel = 200

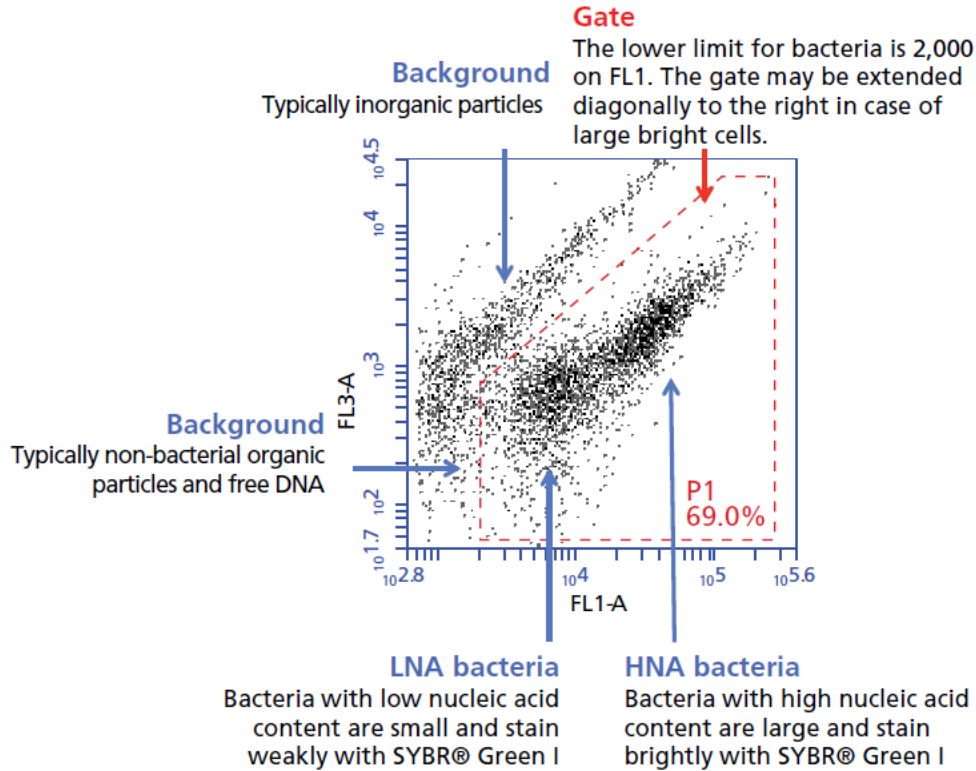


E. Coli per uL sample:
events in P1 / uL collected
 $76,050 / 12.3 \mu\text{L} = 6183 \text{ E. Coli}$

Scatter of Syto BC + e. coli

Noise, very small debris

Water Quality Assessment (Eawag method)



Water Quality Assessment (Eawag method)

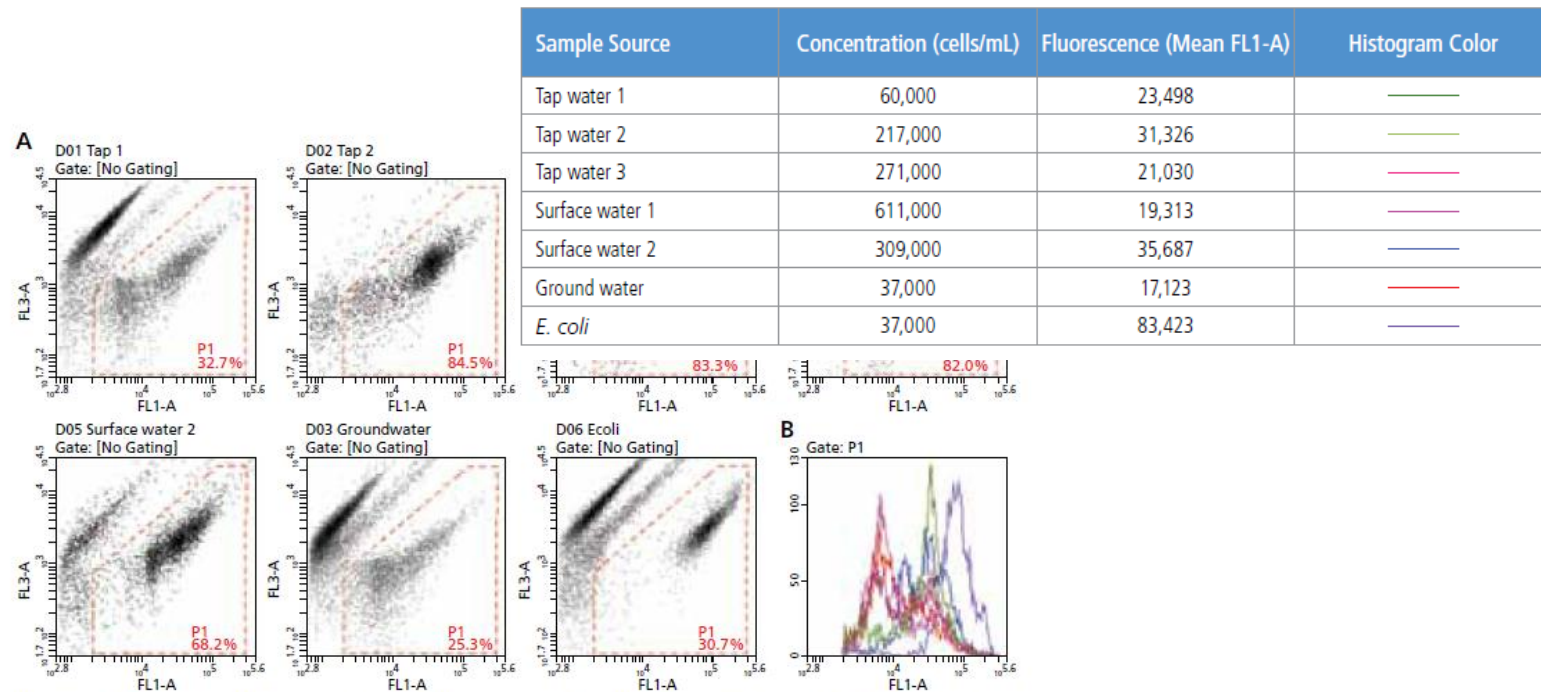
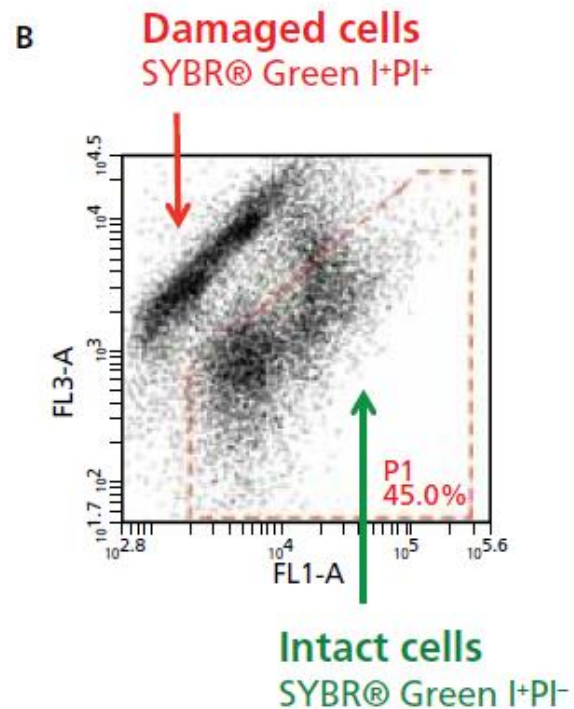
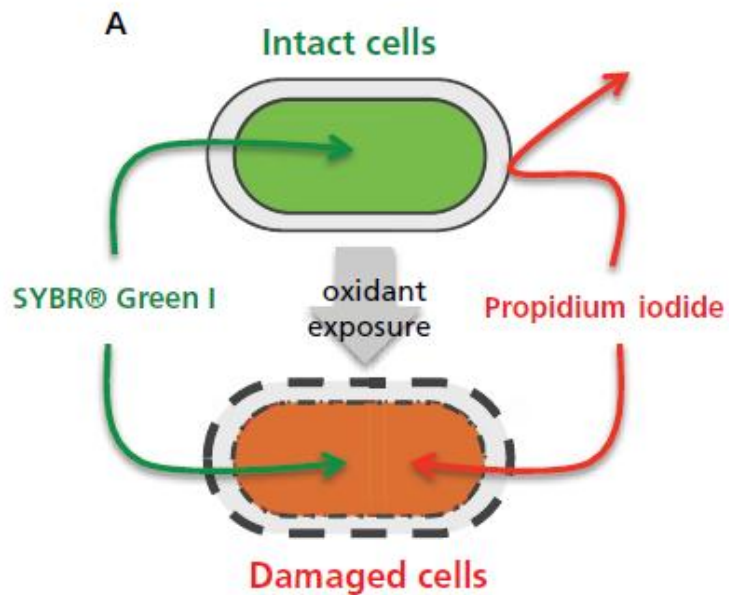
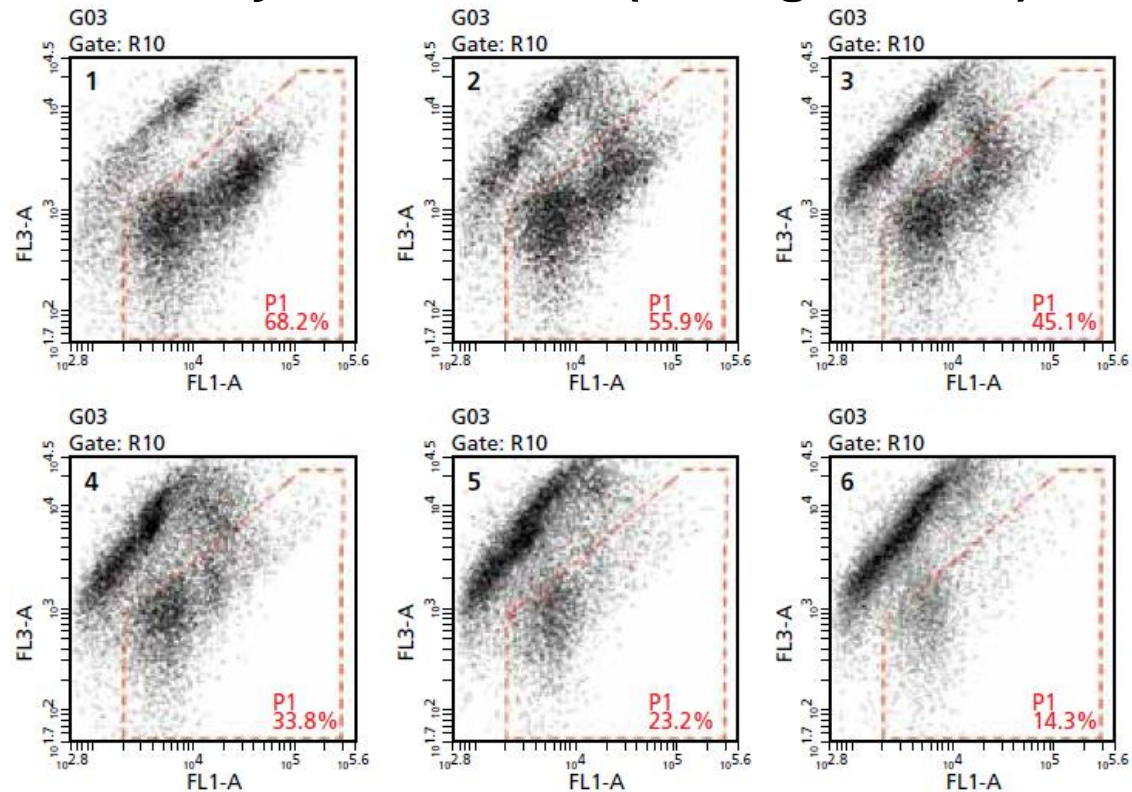


Figure 6. Water samples analyzed using the Eawag method and template on the BD Accuri C6.

Water Quality Assessment (Eawag method)



Water Quality Assessment (Eawag method)

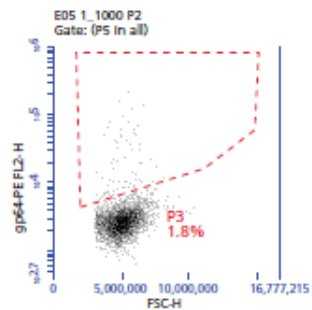
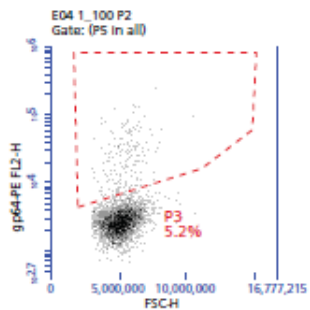
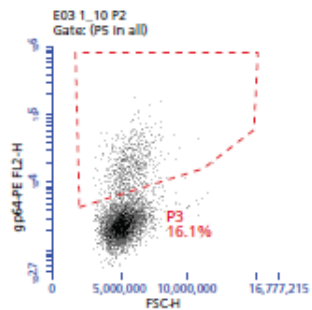
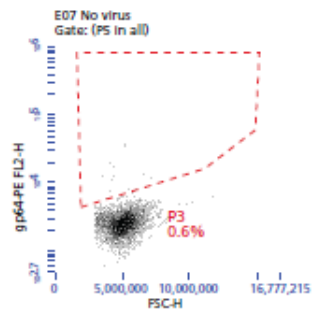


Baculovirus Titration

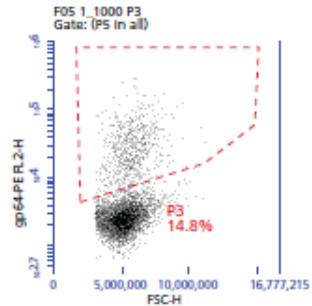
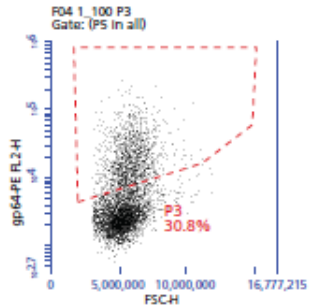
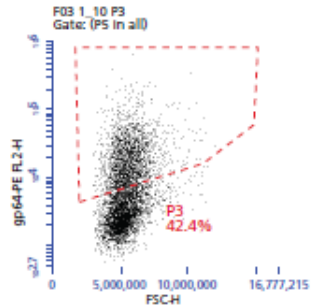
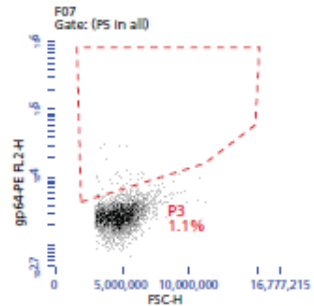


Generation

P2



P3



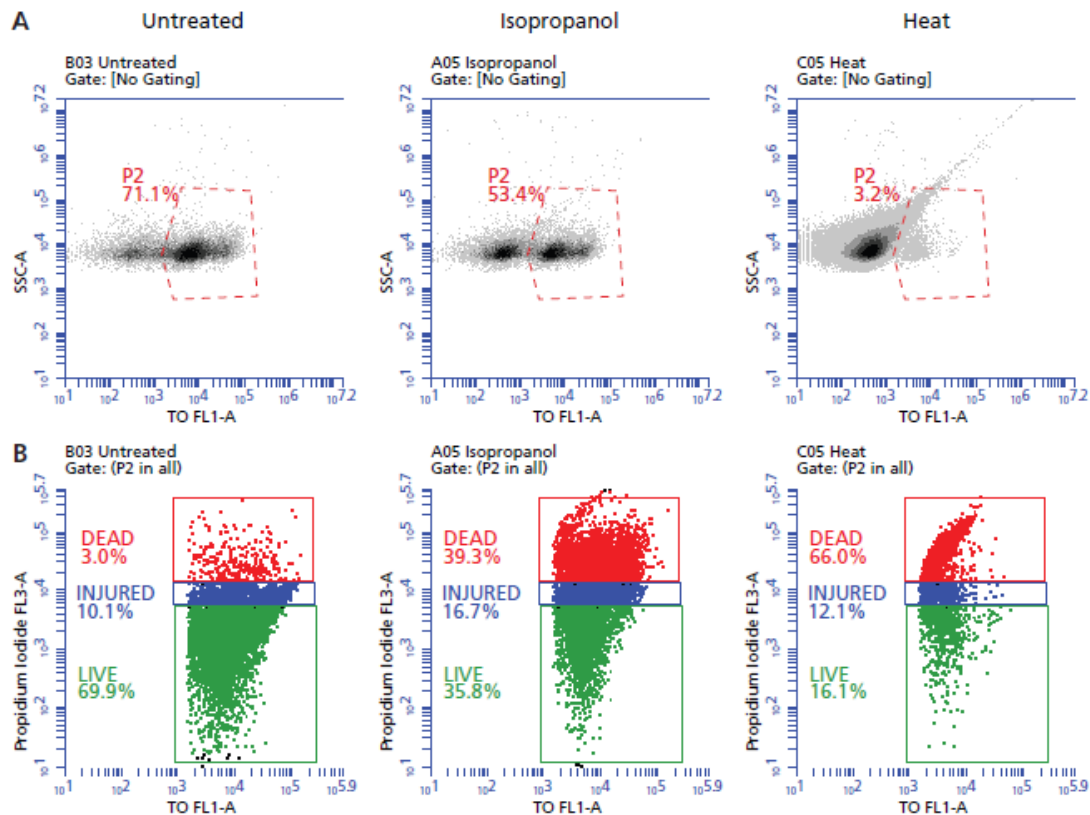
Viral Dilution → No virus

1:10

1:100

1:1000

E.Coli Viability Staining



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