

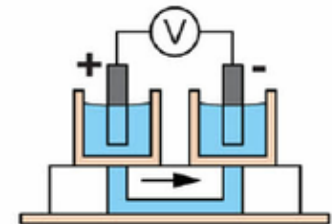
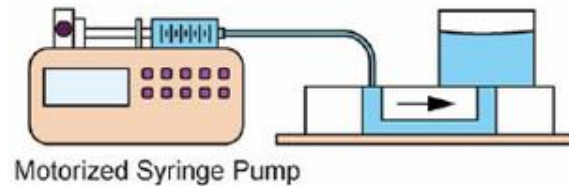
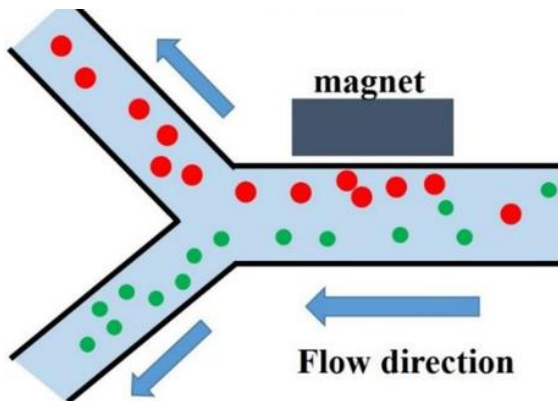
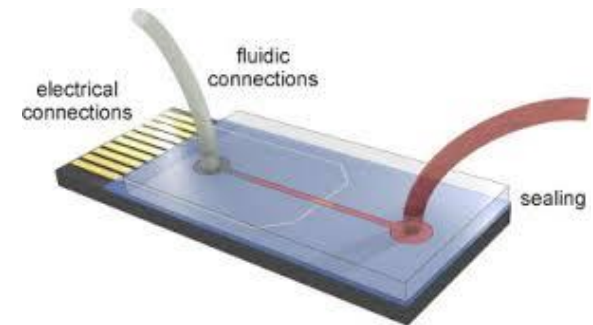


Enhanced capture of magnetic microbeads in a microfluidic channel for water treatment

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- **μ TAS**: micro total analysis systems or lab-on-chip devices^{1,2}
 - Small, fast response, reliable
- **Magnetophoretic** separation³
 - Adaptable, reliable



¹Temiz et al (2015)

²Byun et al (2014)

³Jiang et al (2014)



Objective & Hypothesis

- *Objective*

To develop an efficient **lab-on-chip device** by increasing **capture efficiency** of magnetic microbeads

- *Hypothesis*

The **flow switching** protocol will significantly increase the **capture efficiency** of microbeads, greatly increasing the reliability and accuracy of the device

- *Specific Aim 1*

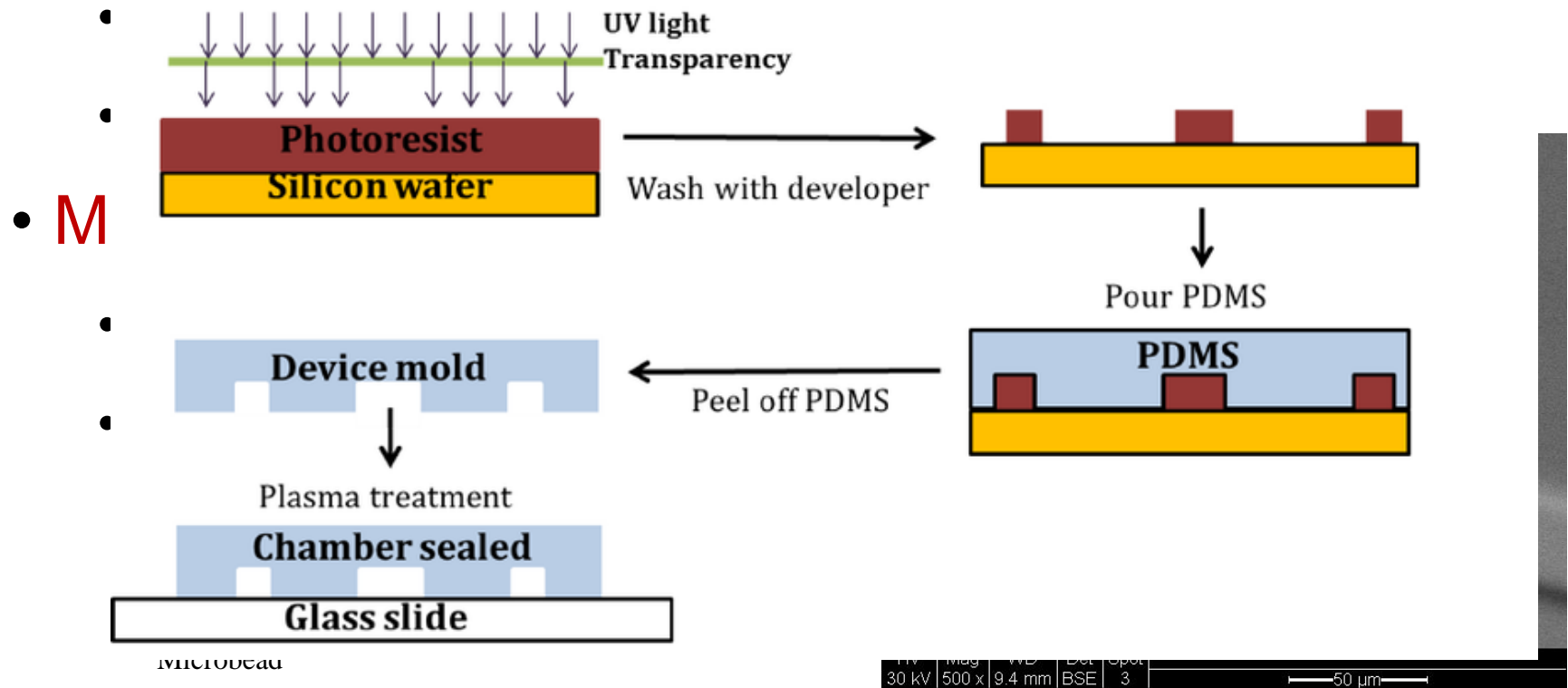
- Examine the effect of **electroosmotic flow switching** on capture efficiency in microfluidic channels

- *Specific Aim 2*

- Evaluate **bacteria binding** and create a **calibration curve** for bacteria capture in the device

• Microchannel

• SU-8 micro lithography⁴



- **Electroosmotic Flow (EOF):** plug flow profile⁵, easy flow manipulation

$$U_{ep} = - \frac{E_z \epsilon_r \epsilon_o \zeta_p}{\mu}$$

U_{ep} : fluid velocity (cm/s)

E_z : applied electric field (V/cm)

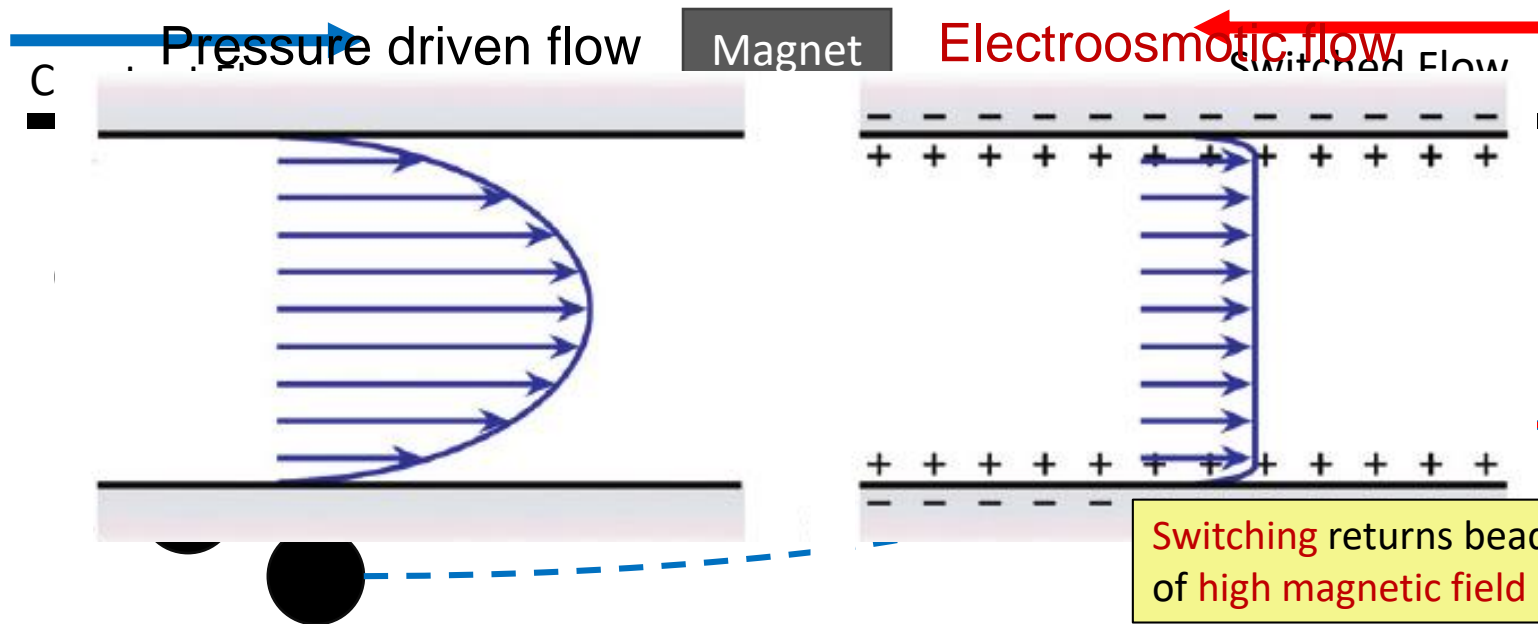
ϵ_r : dielectric constant of medium

ϵ_o : vacuum permittivity (F/m)

ζ_p : zeta potential (V)

μ : dynamic viscosity (Pa·s)

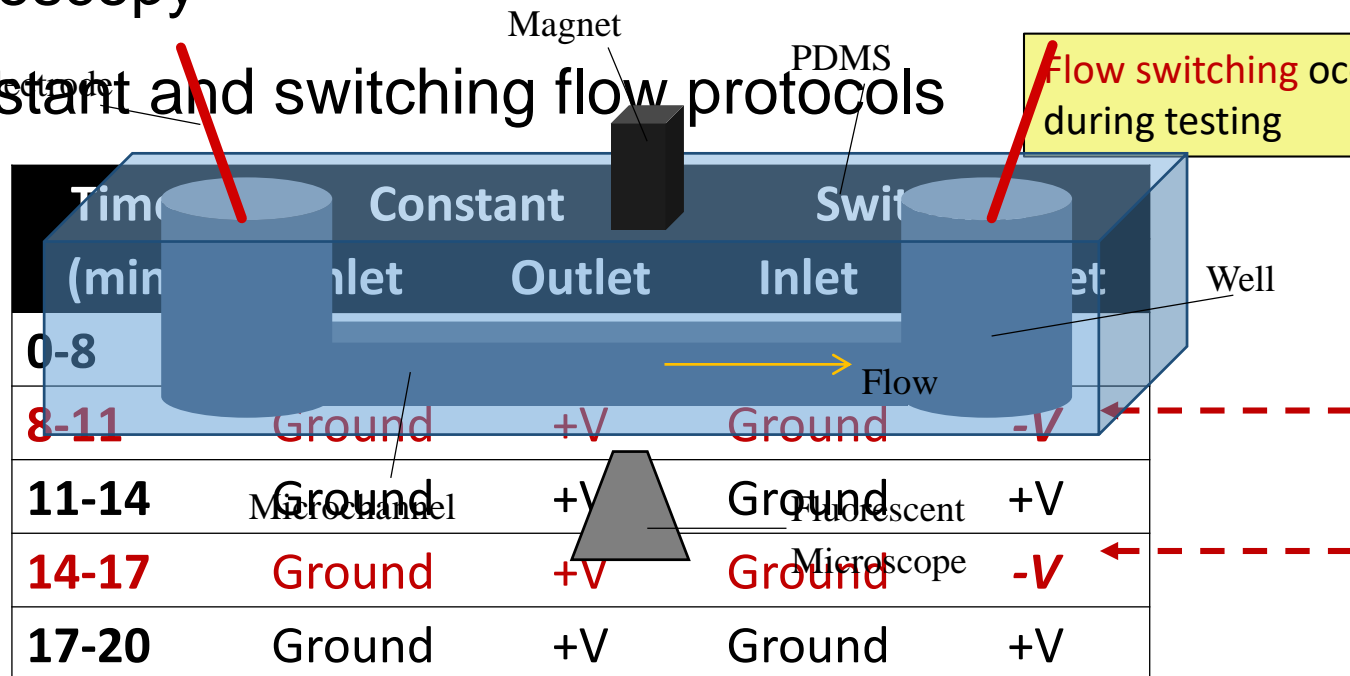
- Constant flow vs. **switching flow**



• Testing Procedure

- PDMS microchannel bound to glass slide
- Analysis performed using inverted fluorescent microscopy

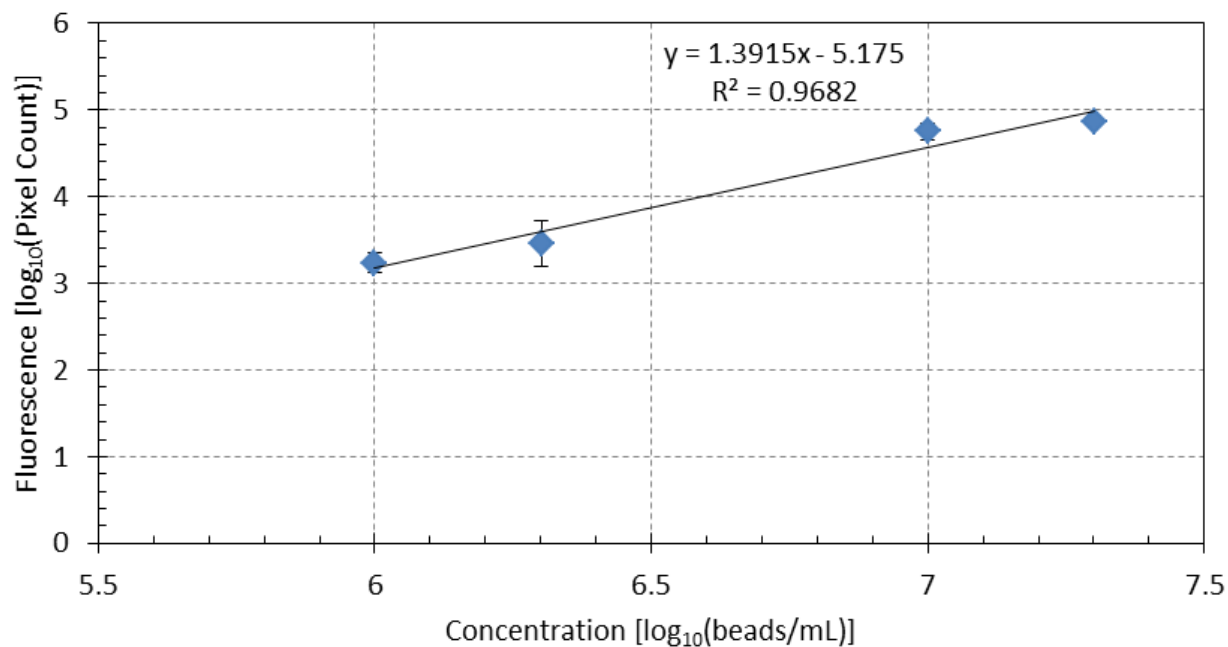
- Constant and switching flow protocols



• Calibration Curve

1×10^6 beads/mL

2×10^6 beads/mL



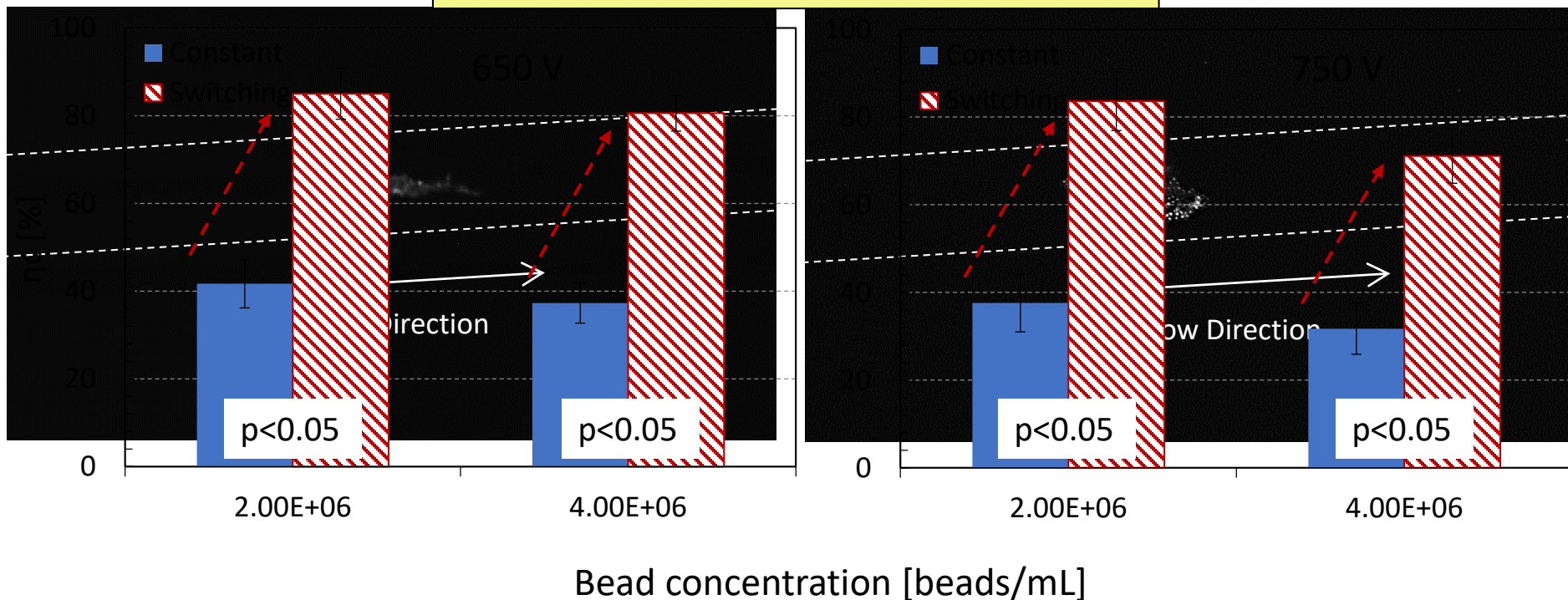
1×10^7 beads/mL

2×10^7 beads/mL

• Capture Efficiency

$$\bullet \eta_c = \frac{\text{pixel count captured}}{\text{pixel count captured} + \text{pixel count uncaptured}}$$

Capture efficiency **increased** up to **2 times** using the flow switching protocol.



- *Specific Aim 1*

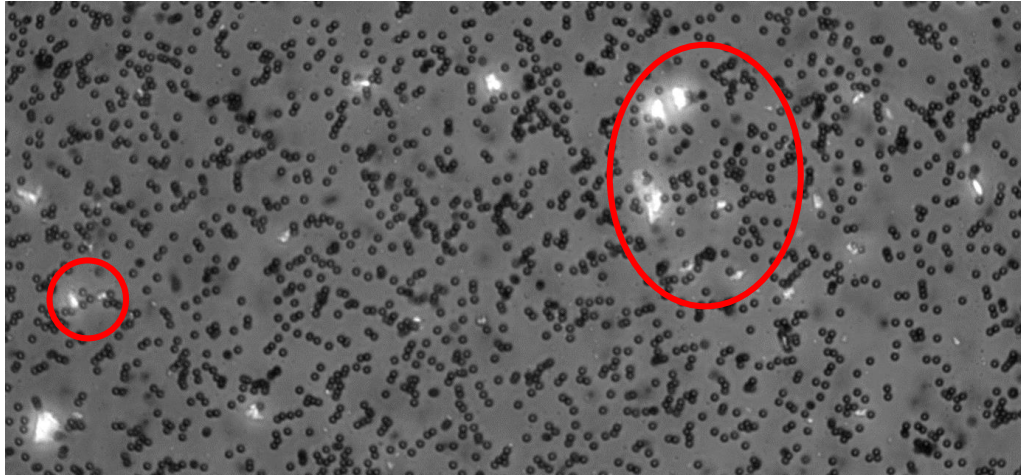
- Examine the effect of **electroosmotic flow switching** on capture efficiency in microfluidic channels

- *Specific Aim 2*

- Evaluate **bacteria binding** and create a **calibration curve** for bacteria capture in the device

- **Bacteria binding**

- Separate antibodies to prevent bead-tag complexes
- Excess beads to bind efficiently



- **Future**

- Completion of calibration curve for bacteria capture

- **Fluorescent analysis**
 - Differences in fluorescent intensity shown among varying microbead concentrations
- **Flow switching**
 - Significantly increased capture efficiency compared to constant flow
- **Bacteria binding**
 - Shows promise with binding of microbeads and fluorescent tag



Future Direction

- Completion of **bacteria study** and calibration curve
- Analysis of **different bacteria** to increase device versatility
- Development of device into a **closed system** with analysis sensors included



Acknowledgement

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Thanks & Questions