

# A Microfluidic Device for Electrofusion of Biological Membranes

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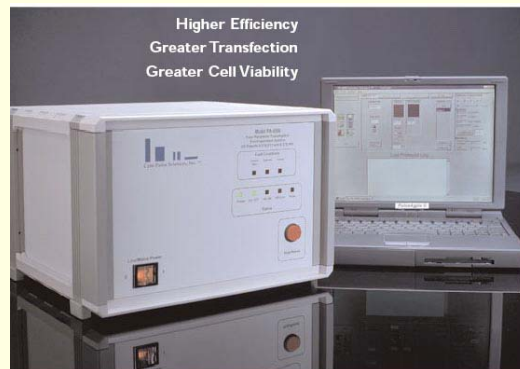
The University of Tokyo



# Framework

## Electrofusion of liposomes and cells in a MEMS device

### Electrofusion



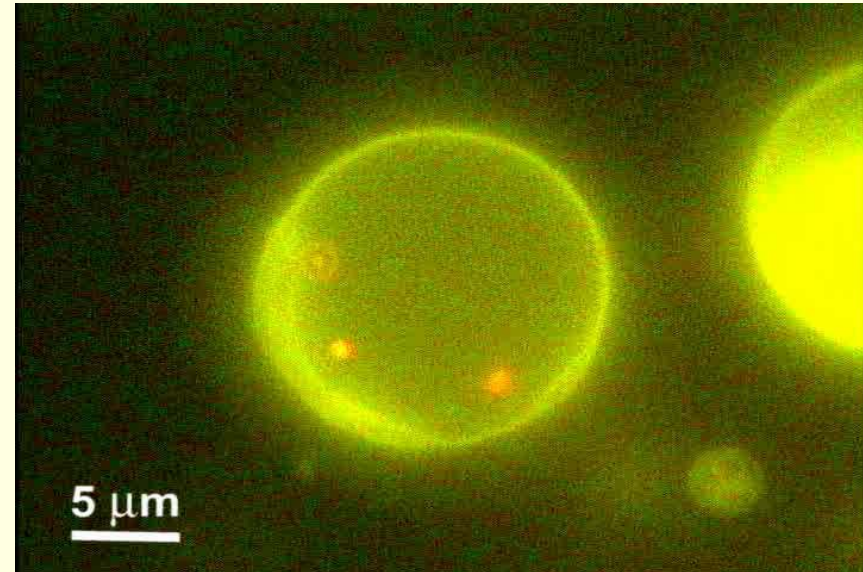
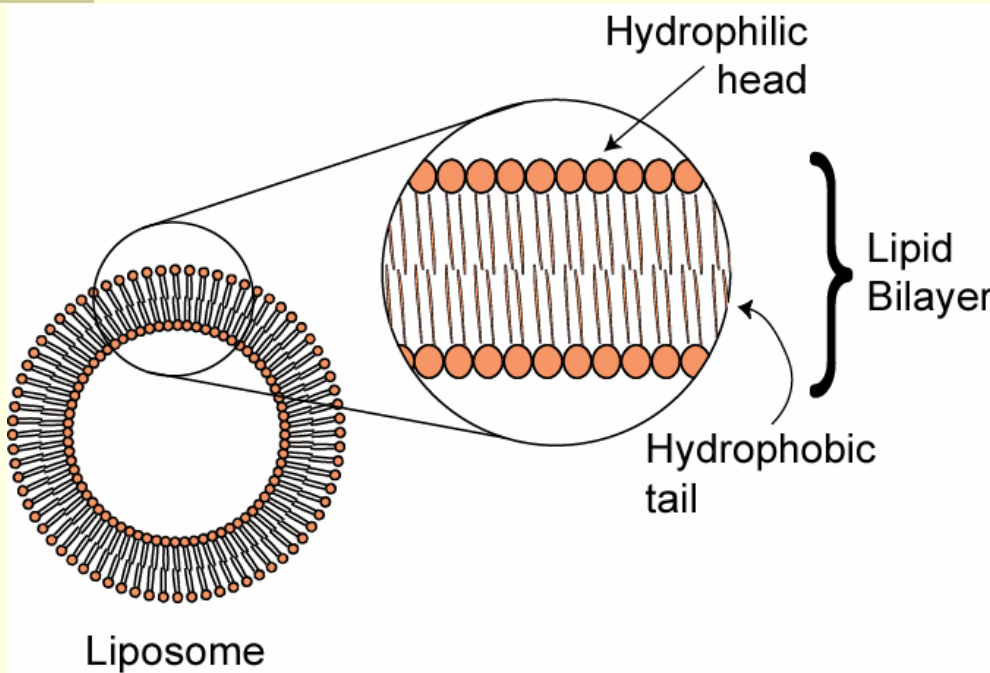
Hybrid cell  
production

### MEMS

- Lab-on-a-chip integration
- Power reduction
- Future industrial fabrication

# What Is a Liposome ?

## Synthetic lipid-bilayer container

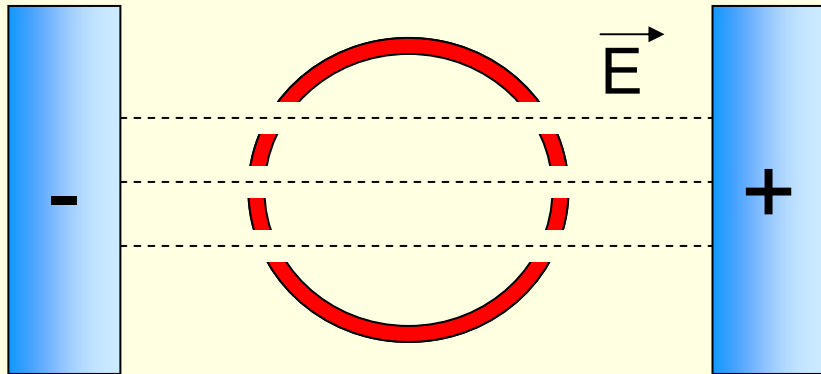


- **Cell-like** membrane structure
- Ability to **enclose** materials

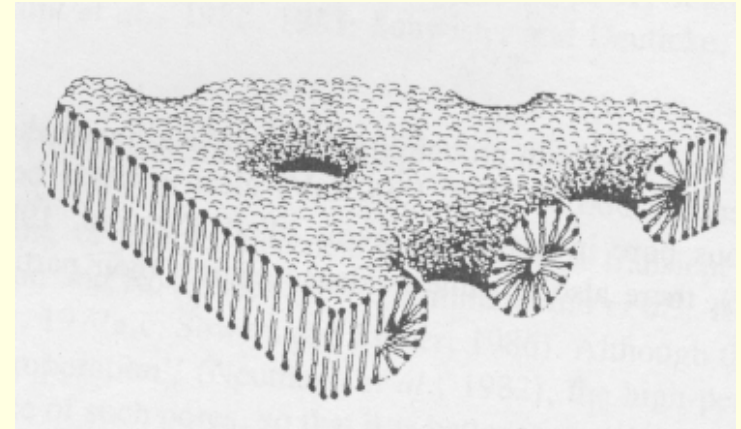


Drug, biological  
and artificial material  
delivery into cell

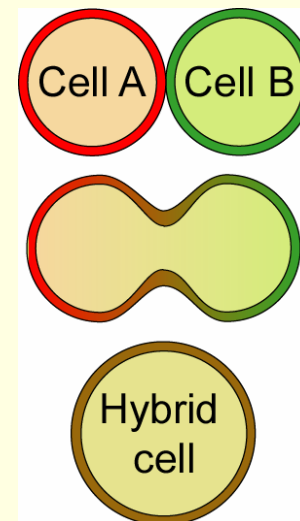
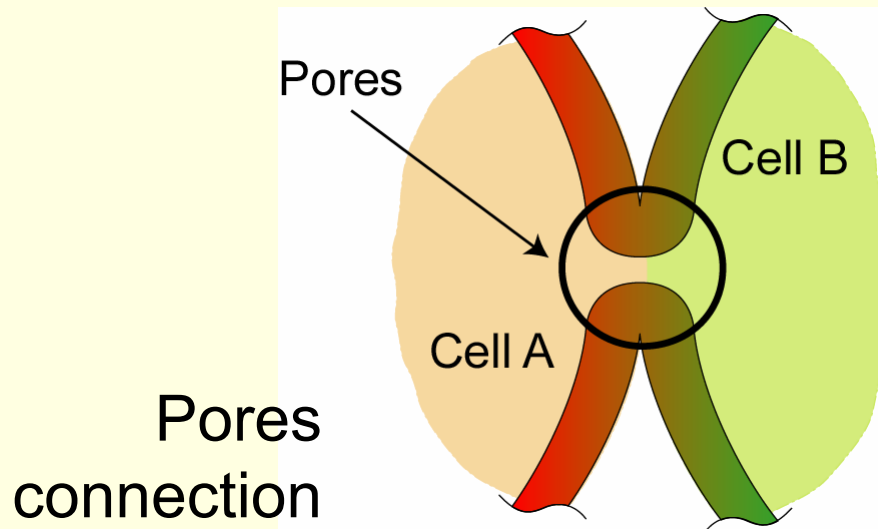
# Principle of Electroporation



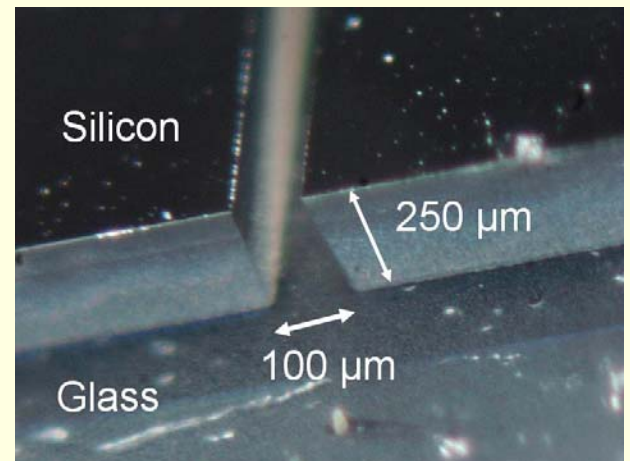
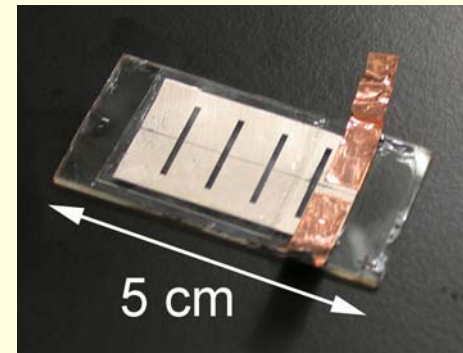
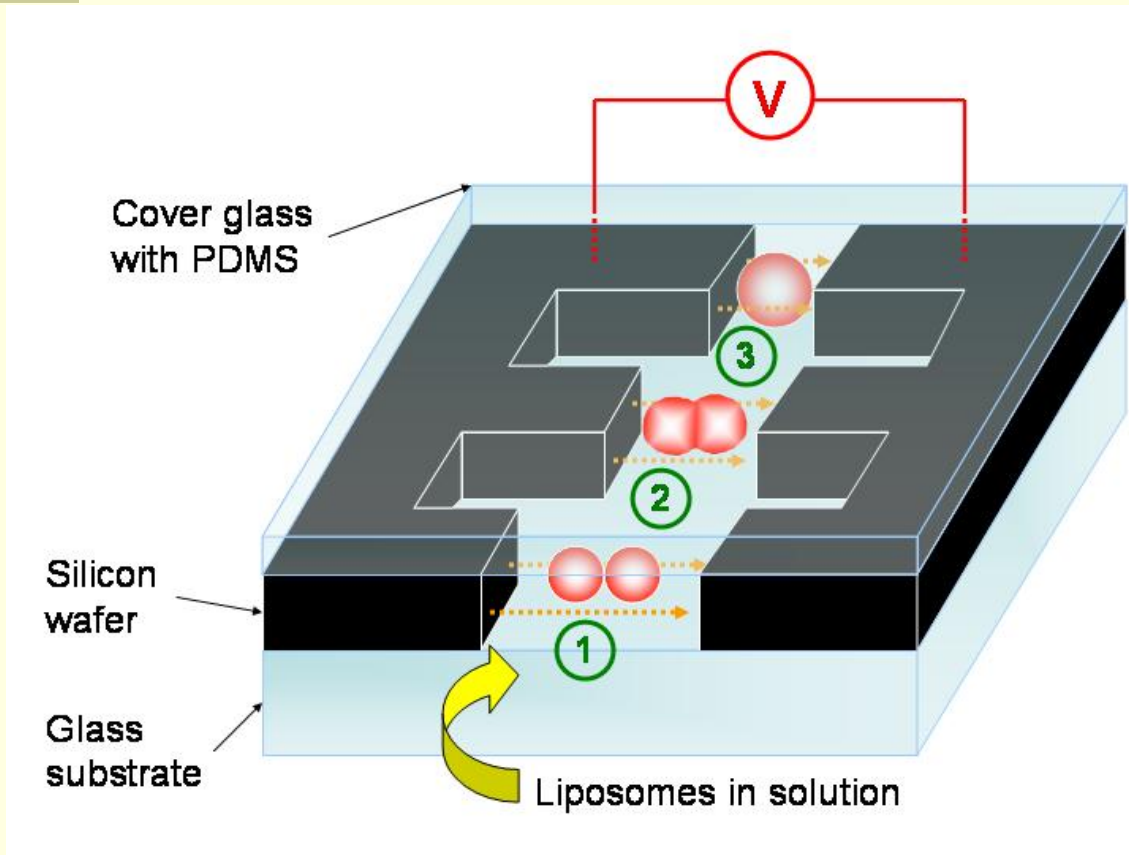
Electroporation:  $E \geq 1 \text{ kV.cm}^{-1}$



Pore size: 10-100 nm



# Microdevice Design



- Reduction of power consumption
- Easy to fabricate

# Electrofusion Protocol

## Fusion

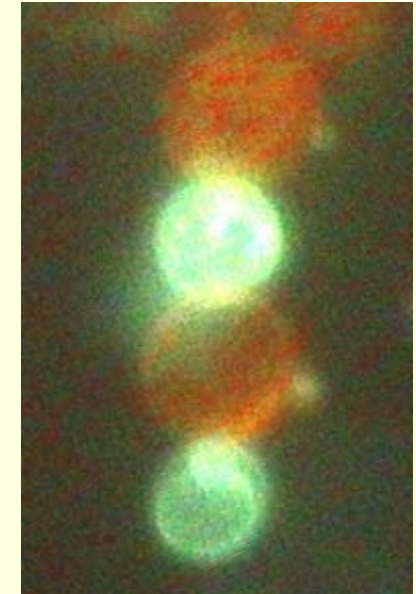
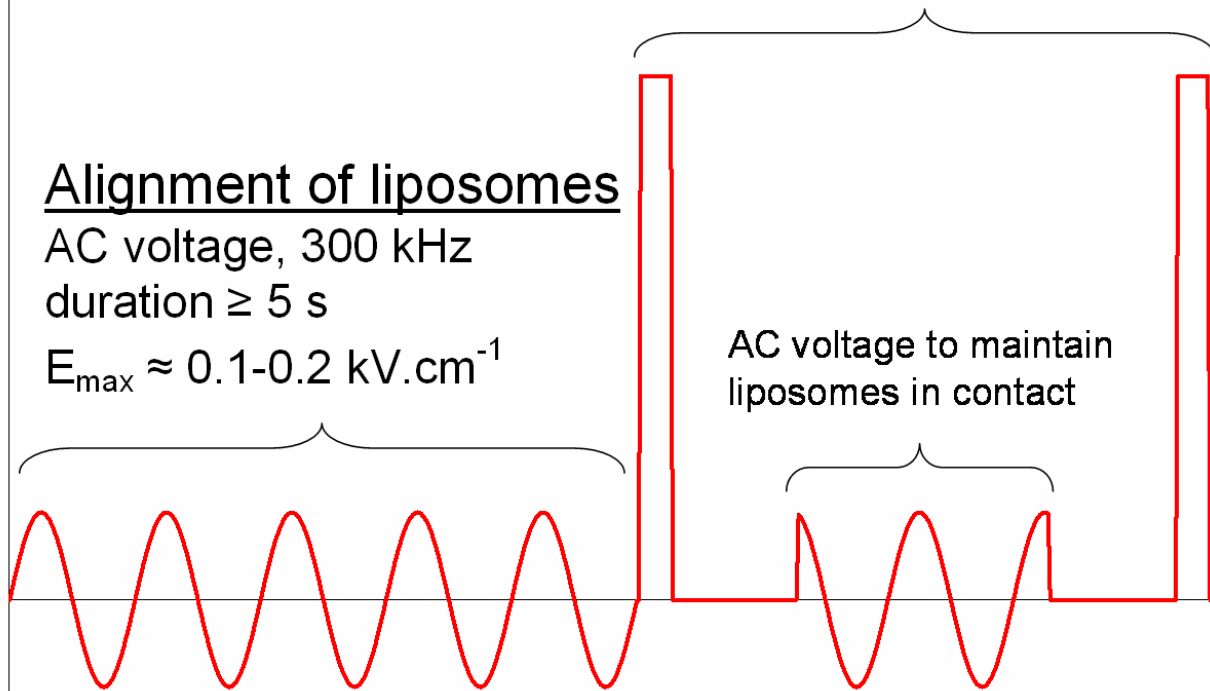
5-6 DC pulses of 10  $\mu$ s every 0.2 s,  $E_{\max} \approx 1-15 \text{ kV.cm}^{-1}$

## Alignment of liposomes

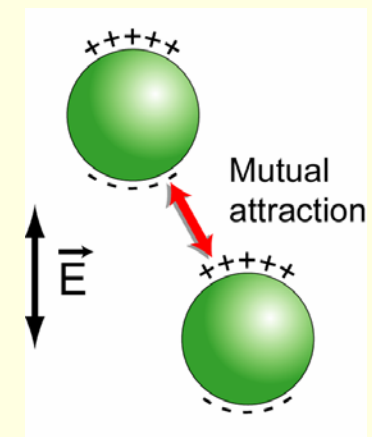
AC voltage, 300 kHz  
duration  $\geq 5$  s

$E_{\max} \approx 0.1-0.2 \text{ kV.cm}^{-1}$

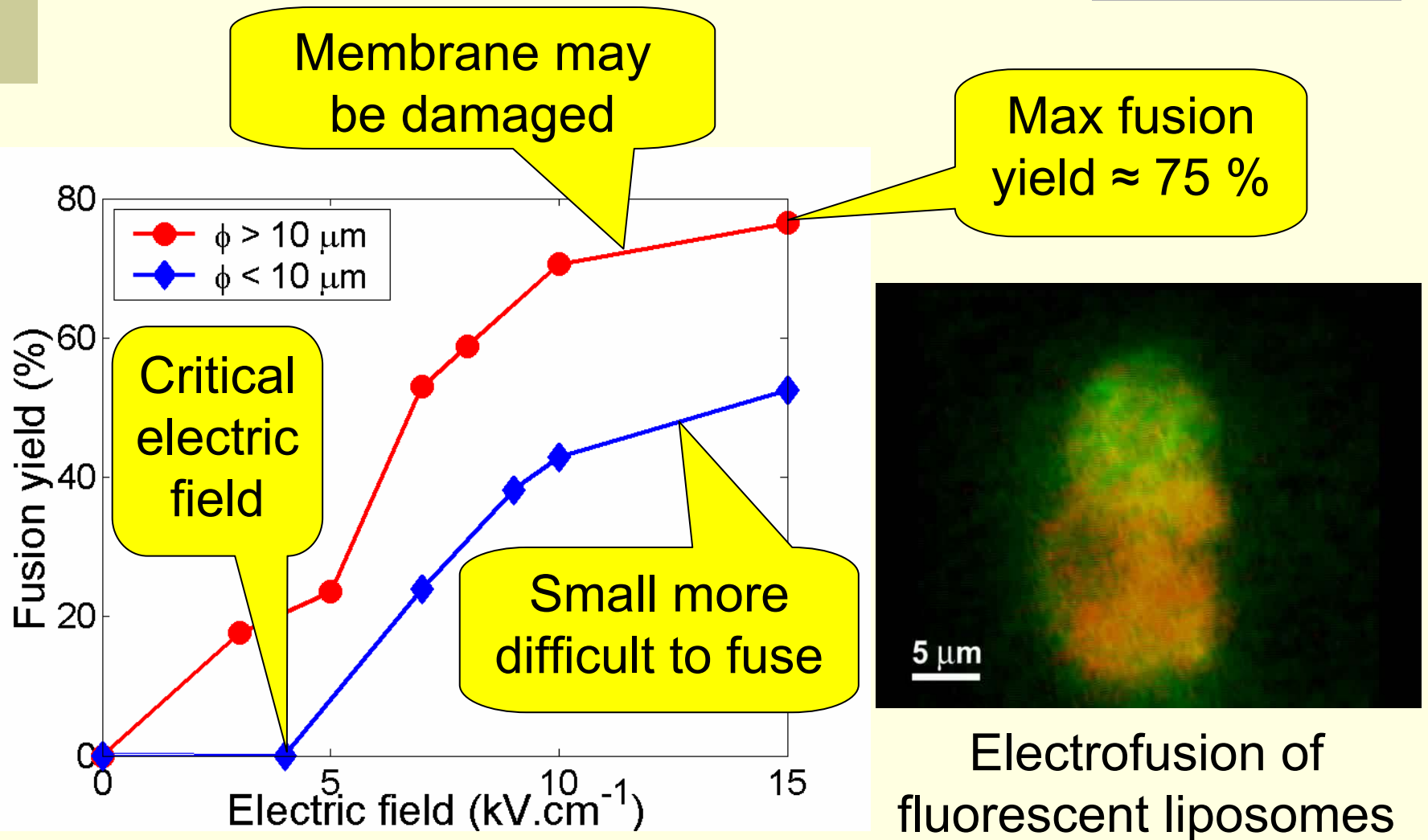
AC voltage to maintain liposomes in contact



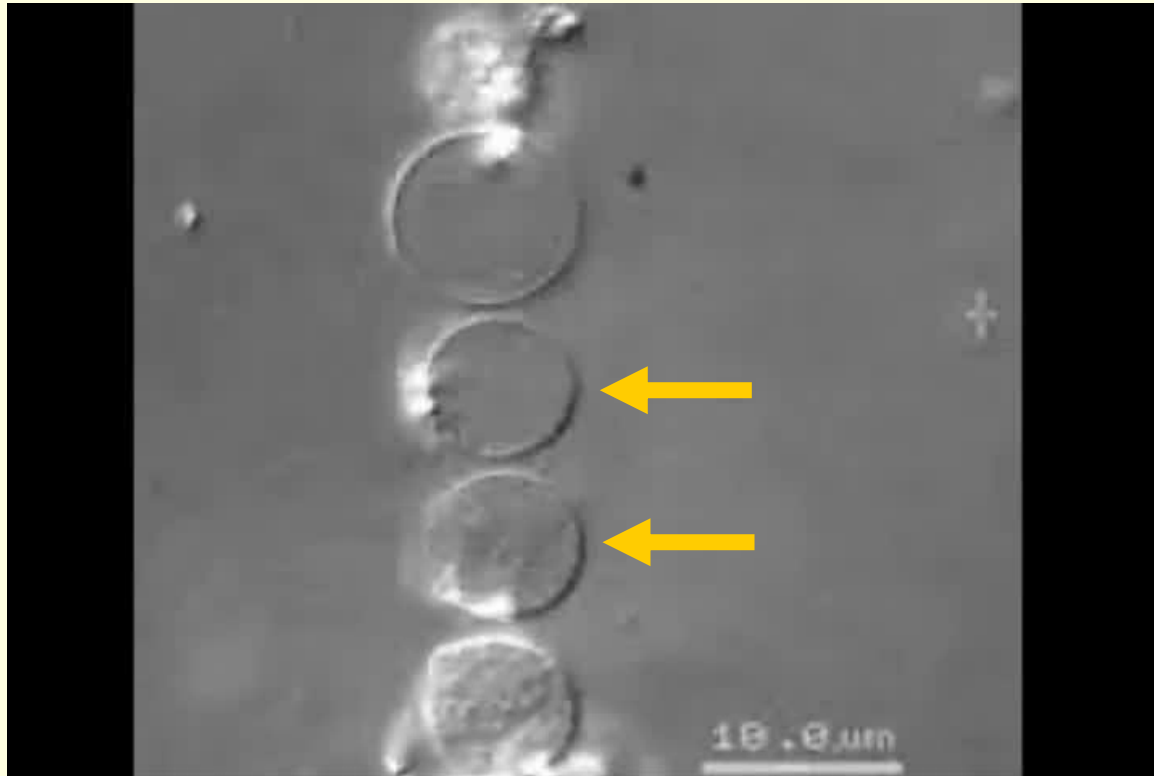
Alignment of polarized vesicles



# Liposome Fusion



# Electrofusion of Bacteria



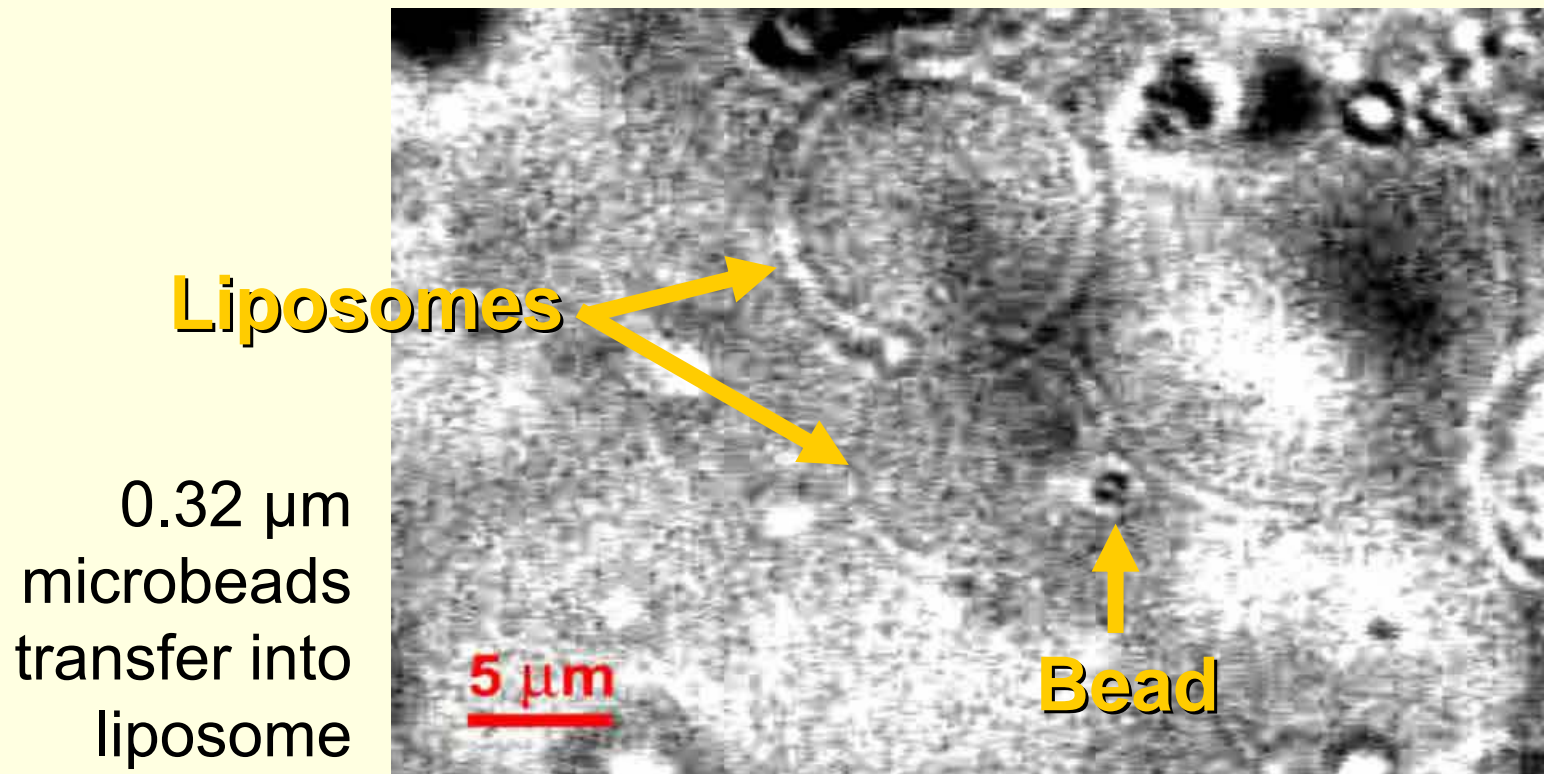
Membranes must be soft and fluid enough

*E. Coli* provacuoles fusion with bulk electrodes



# Towards Transfer of Microstructure

## Microstructure delivery by vesicle-vesicle electrofusion



# Conclusions

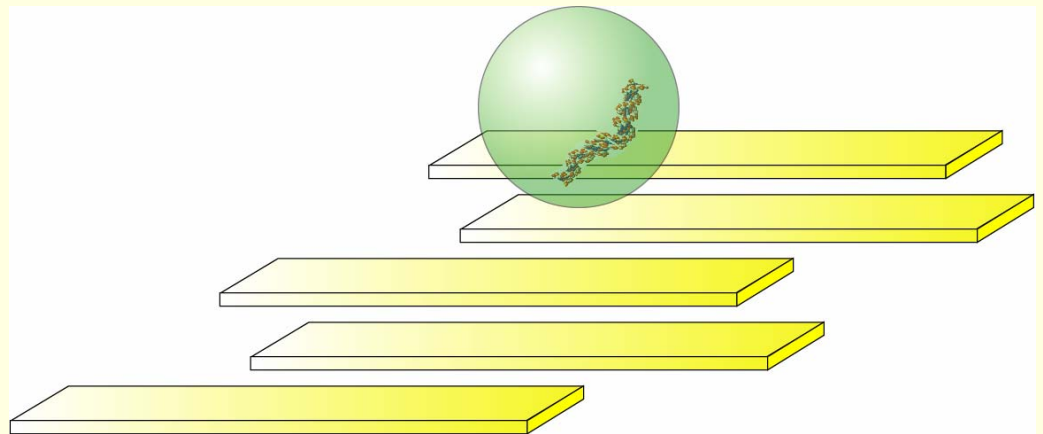
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- A microfluidic device for electrofusion of liposomes at low power level has been designed and tested. Fusion yield can reach 75 %,
- Possibility to fuse cells but the membrane must be soft and fluid to obtain good fusion yield,
- Possibility to deliver microstructures into liposomes.

# Prospects



- Cell labeling for biomedical applications
- Drug and gene delivery
- Single molecule and femtoliter drug manipulation



# Acknowledgements

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