

# LID-PEG Training Session

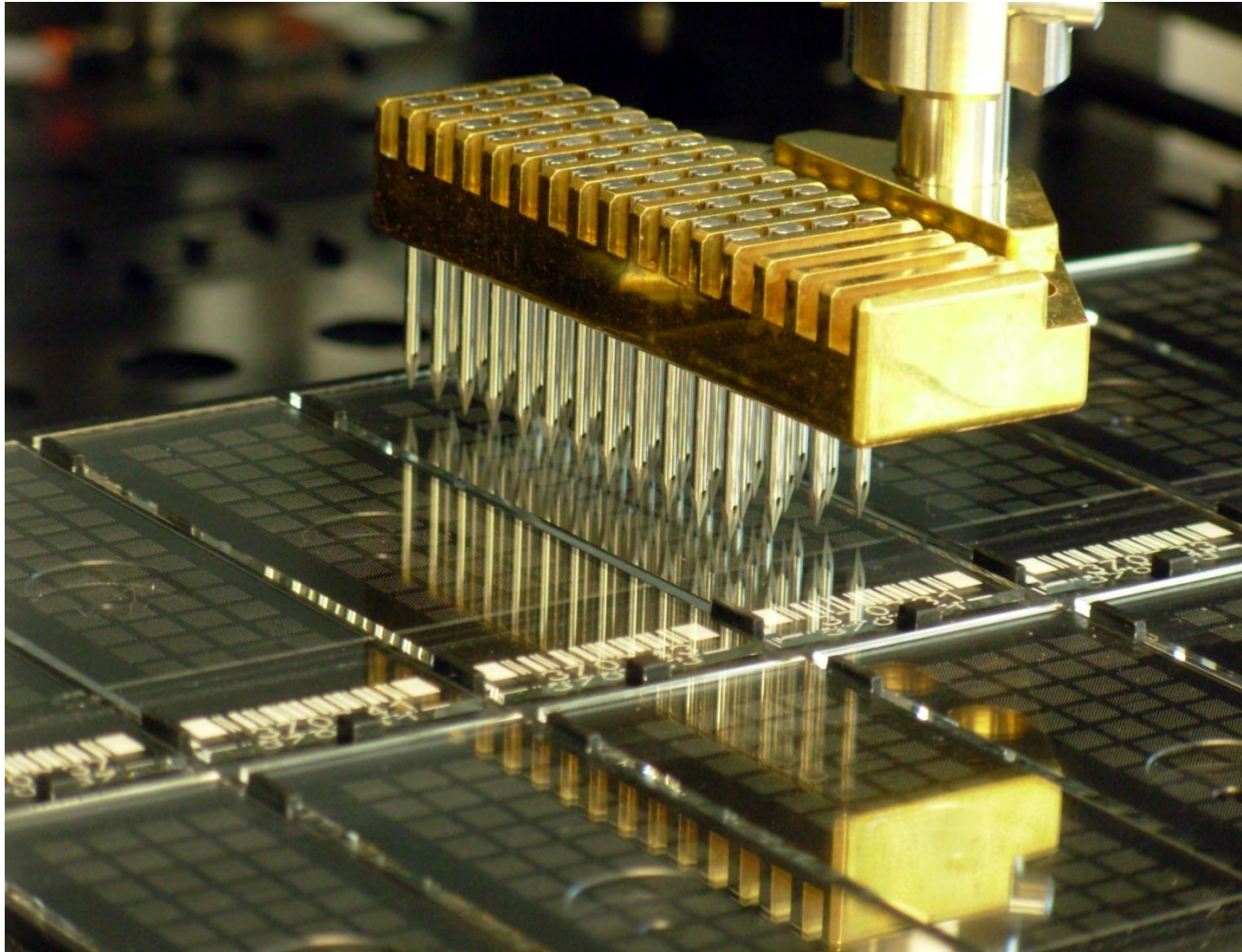
Module2:

Glycan Array Production  
& Screening

Presented by- Ryan McBride

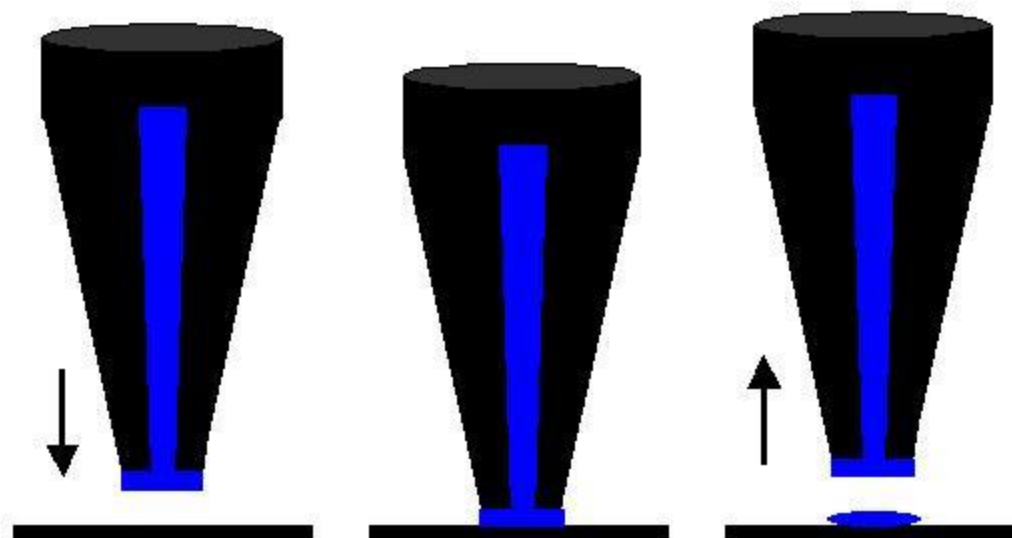
# Outline

- Contact Microarray Printing Basics
- Microarray Design/Printing
- Post Processing
- Sample Analysis
- Image Analysis/Data Acquisition
- Data Analysis/Output



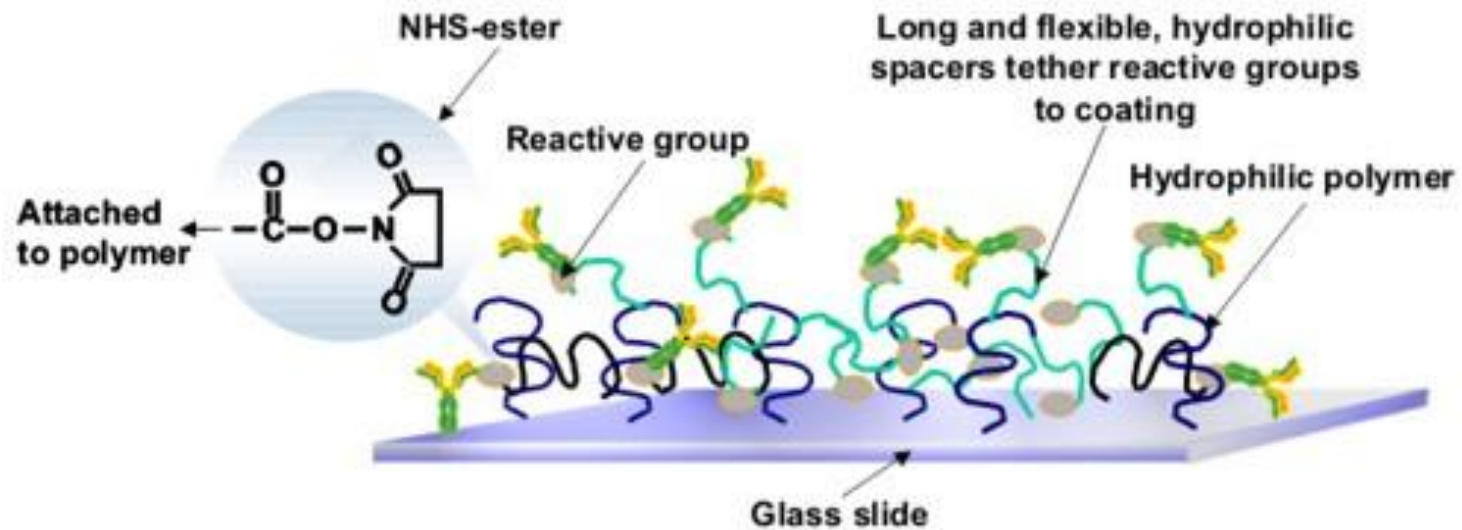
# Contact Microarray Printing

- Functions by passive capillary fluid uptake and delivery
- Stealth Microarray Spotting Pins
  - ~1nL of material deposited/spot
  - 25 – 100 spots per refill (source visit)



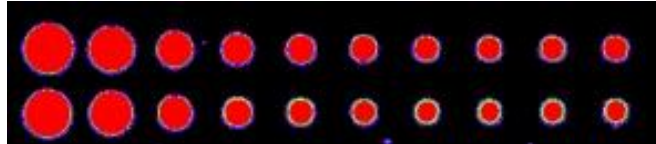
# Slide H

- Made by Schott/Nexterion (Germany)
- Polymer chemistry composed of a scaffold containing NHS-ester groups to bind amine-functionalized/containing compounds and azide groups for the polymerization of the scaffold



# Spotting Issues

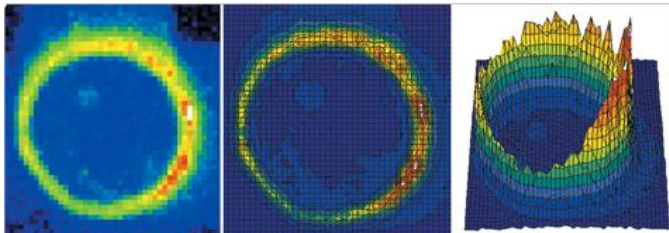
Spot Rush → Spot Taper



## Solution

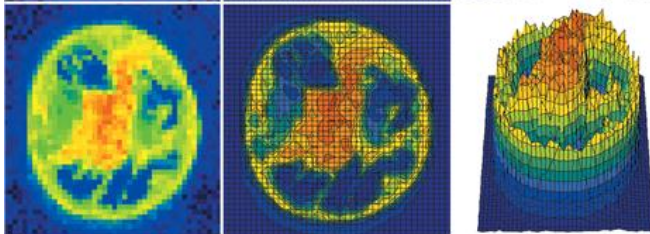
- “dab” pins onto a “pre-”spotting slide (PolyL-Lys)
- like using a quill pen

Coffee stain Spots (Donuts)



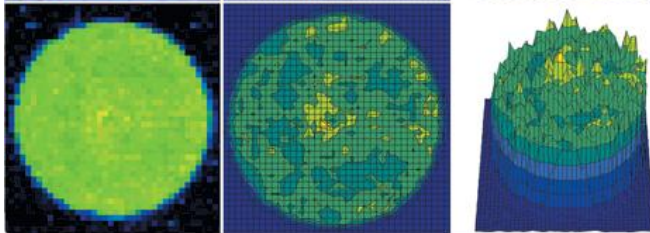
- increase humidification time post print or during printing
- change buffer composition

Spiked spots



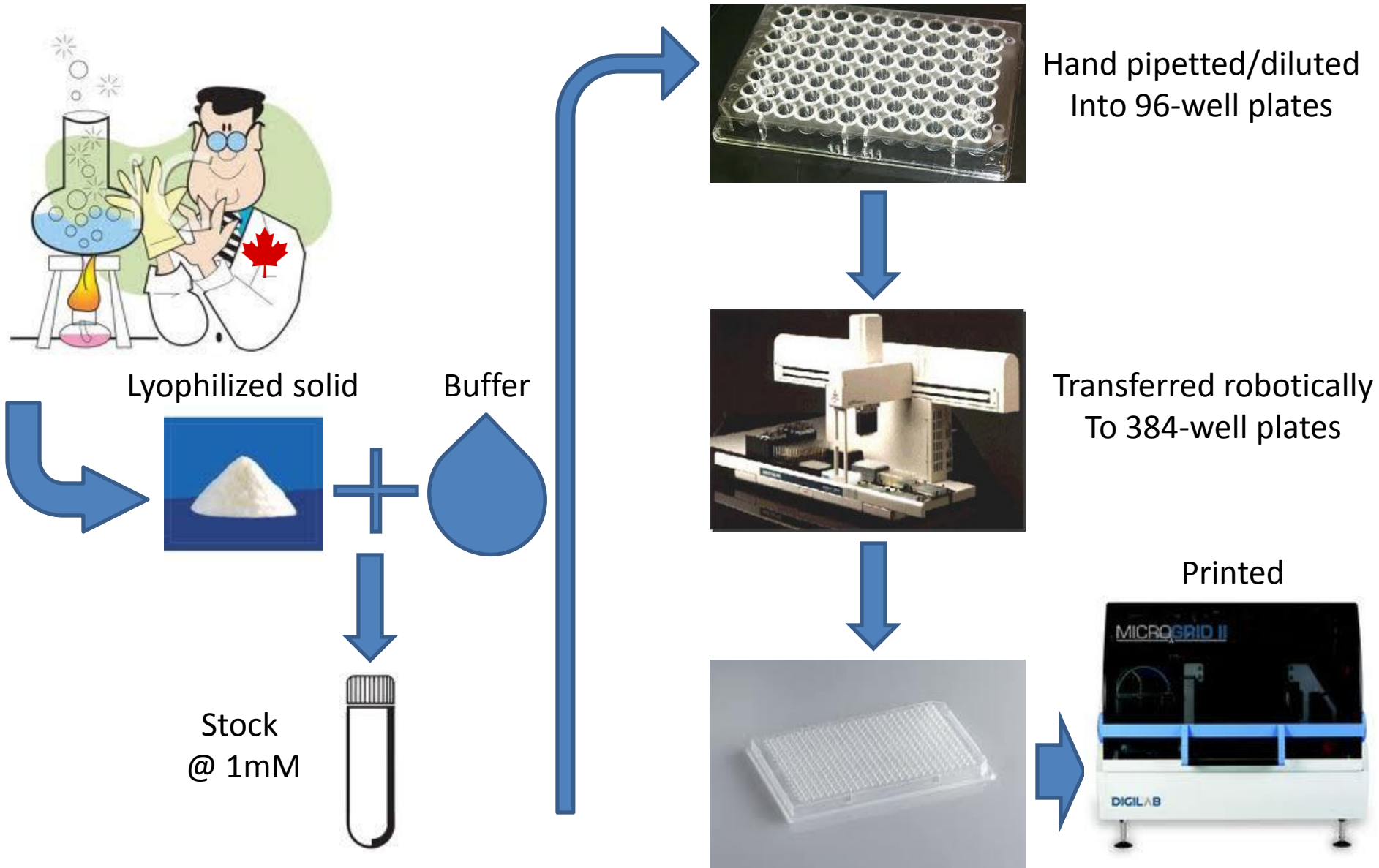
- change buffer composition
- addition of detergents or organics

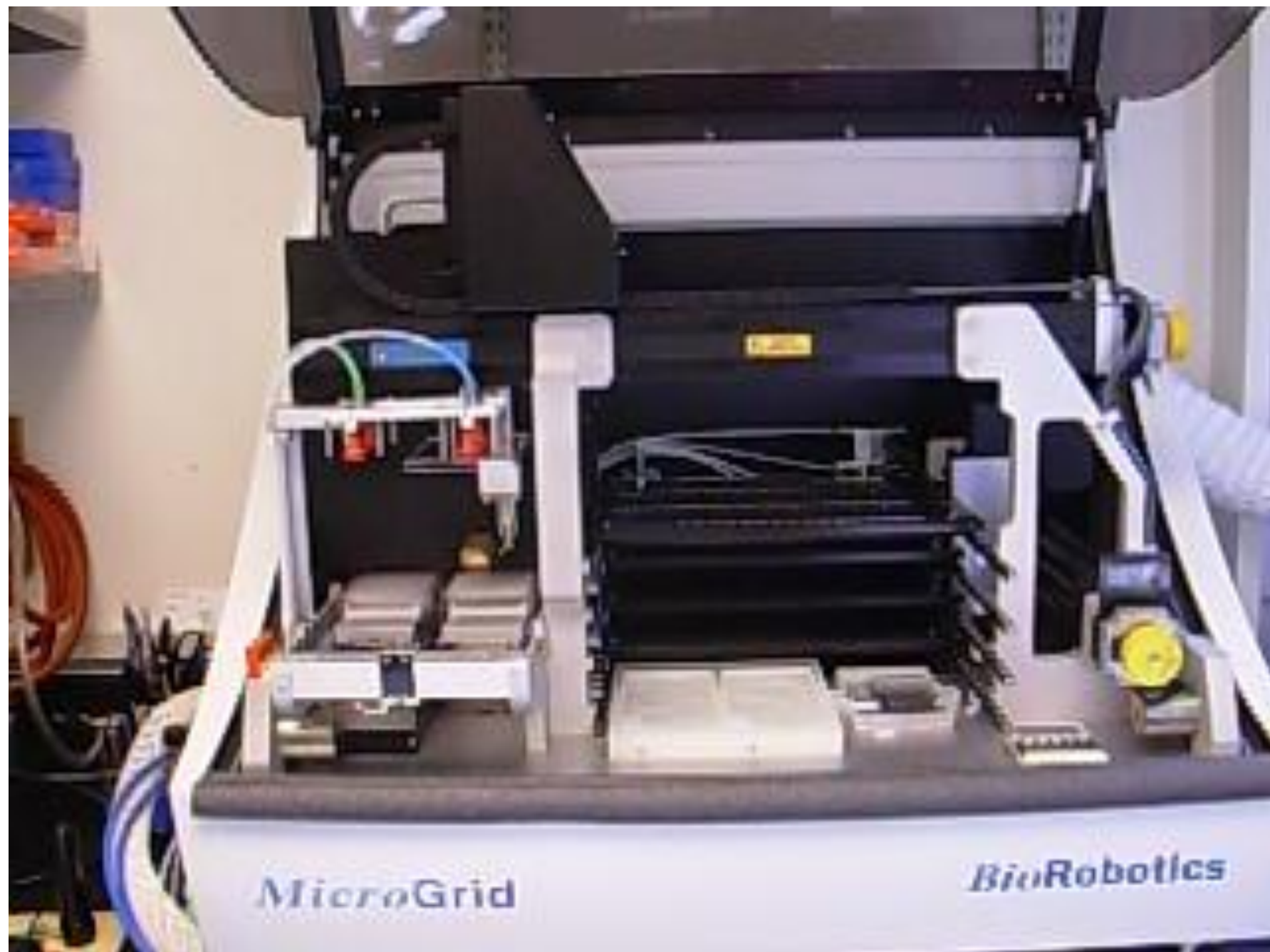
Uniform spot



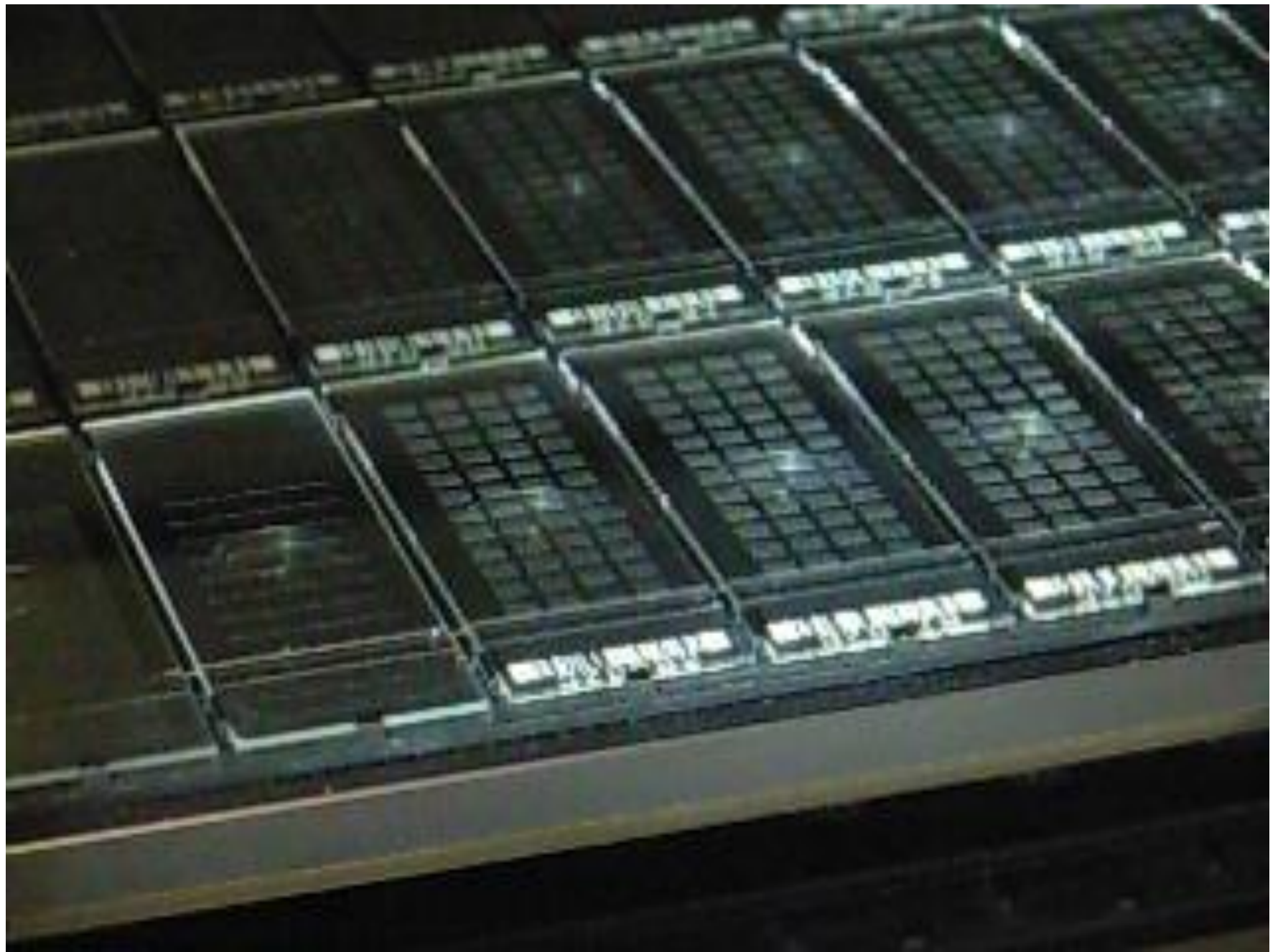
Invitrogen

# Workflow

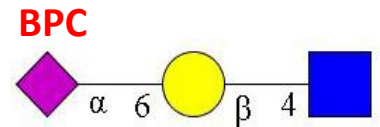
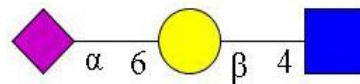
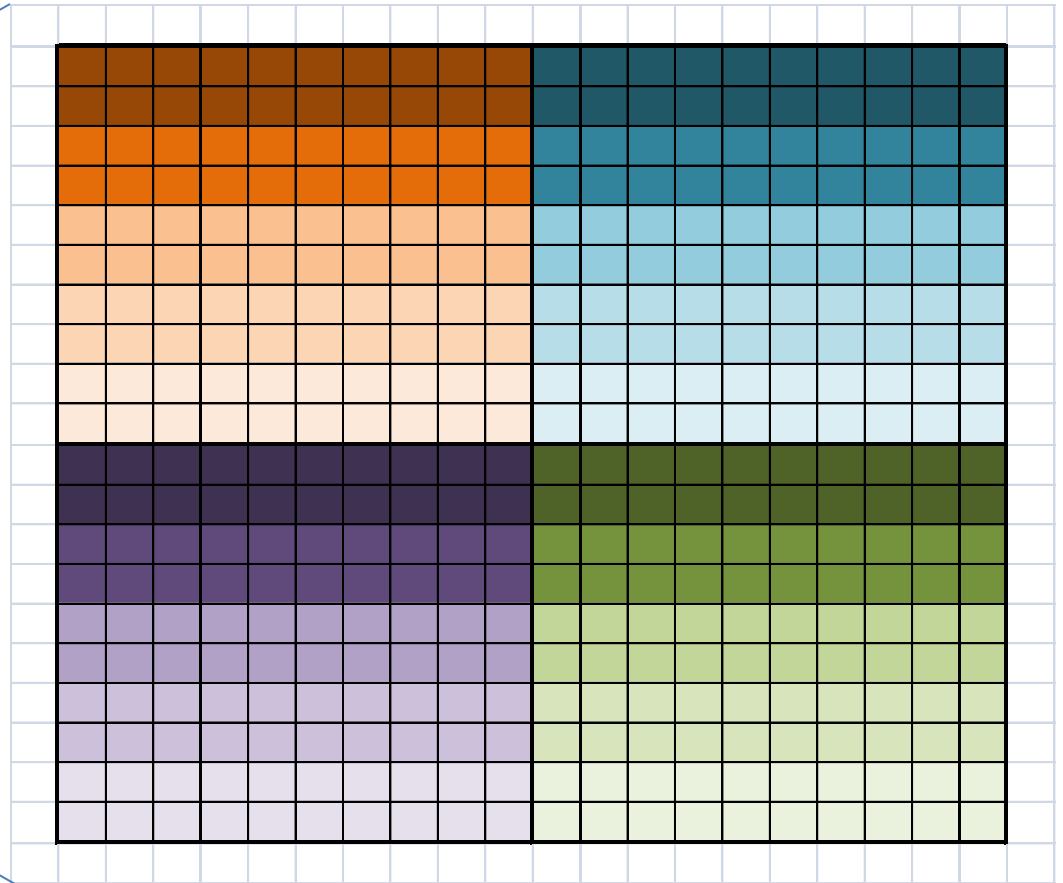
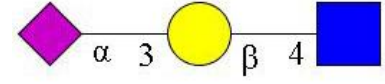
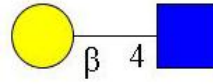
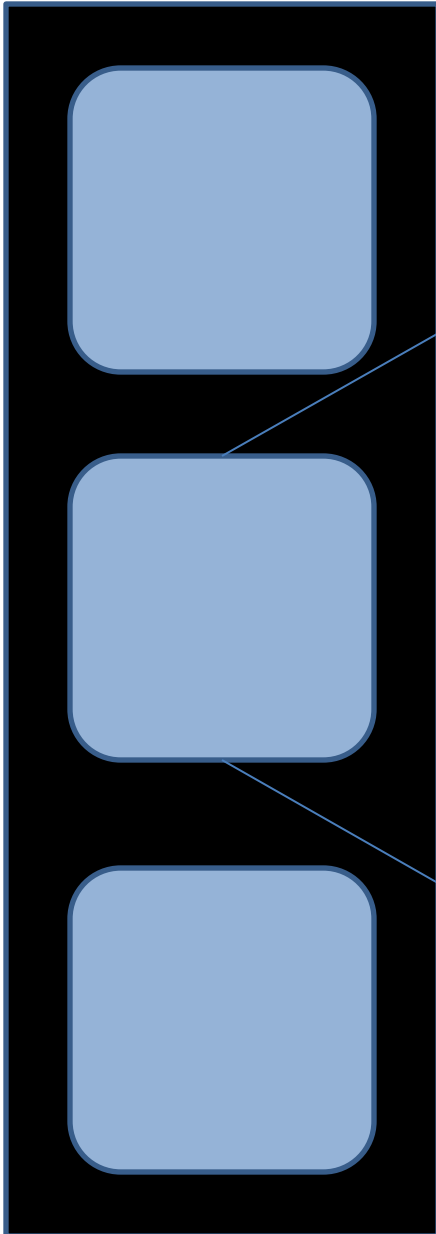




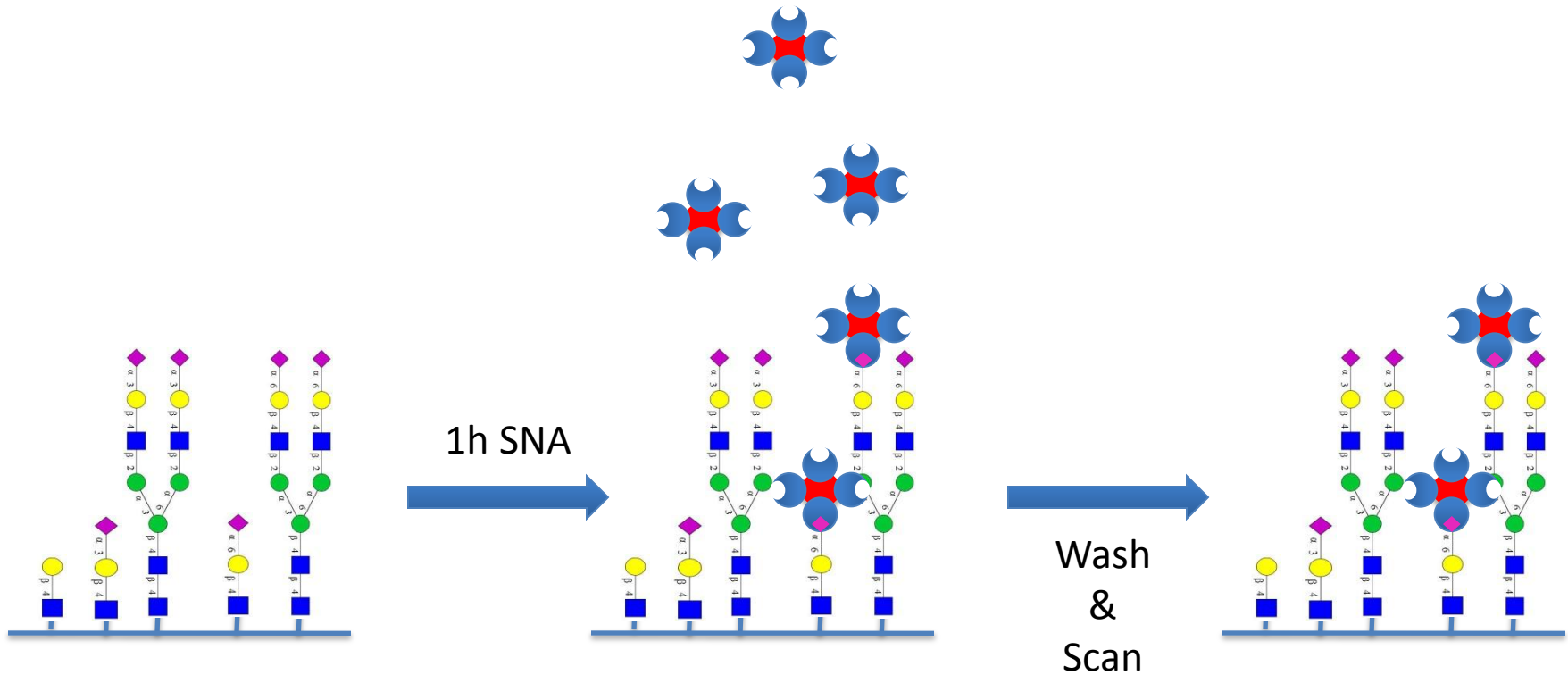






# Array Design



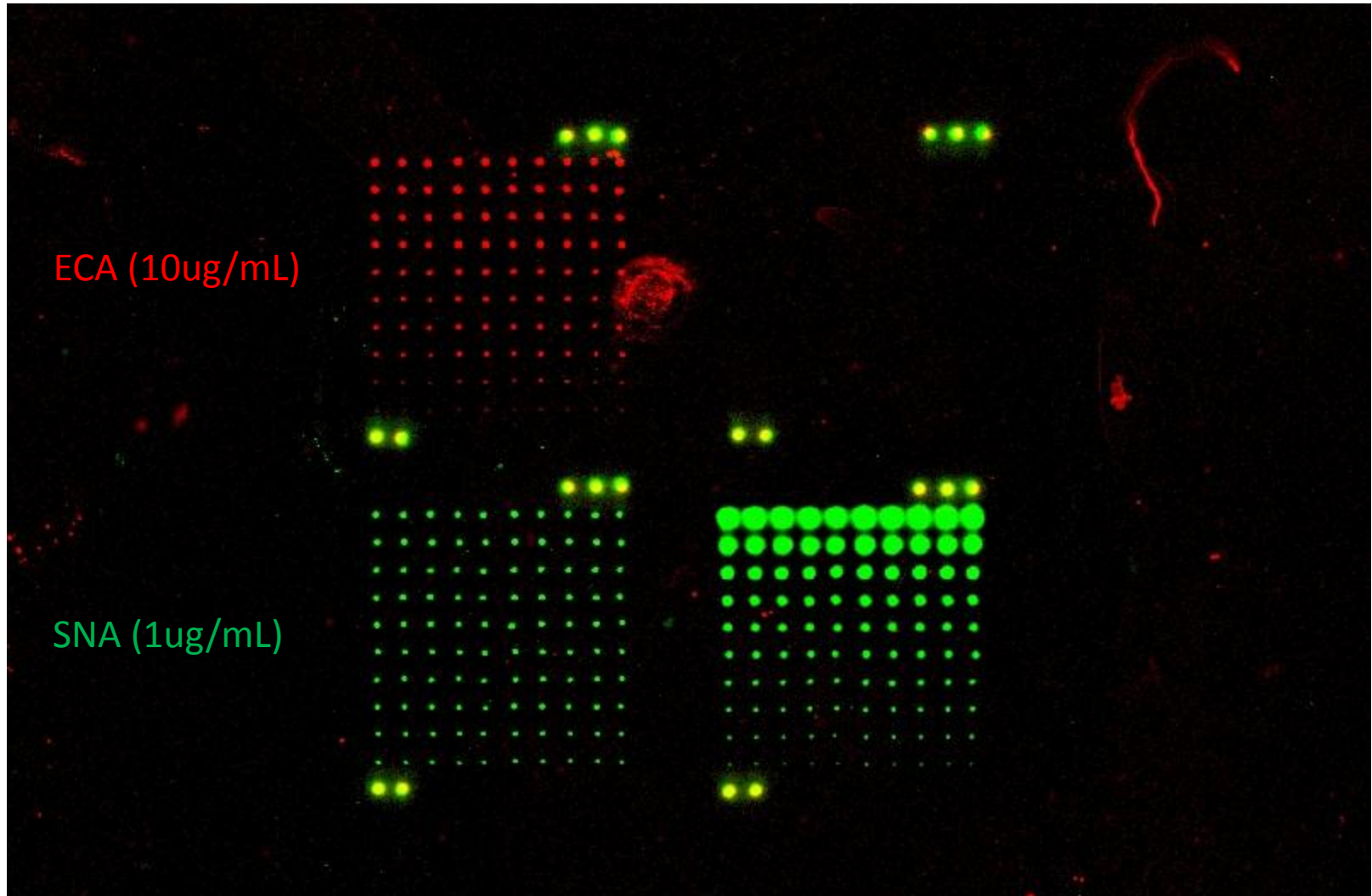
# Sample Incubation



 SNA - Biotin

 Streptavidin –  
AlexaFluor488

# Image Acquisition



# Data Acquisition

MR	MC	R	C	Sample
1	2	11	8	M024 100uM
1	2	12	8	M053 100uM
1	2	13	8	M024 100uM
1	2	14	8	0
1	2	1	9	0
1	2	2	9	M036 100uM
1	2	3	9	M042 100uM
1	2	4	9	M036 100uM
1	2	5	9	M042 100uM
1	2	6	9	M036 100uM
1	2	7	9	M042 100uM
1	2	8	9	M012 100uM
1	2	9	9	M010 100uM
1	2	10	9	M012 100uM
1	2	11	9	M010 100uM
1	2	12	9	M012 100uM
1	2	13	9	M010 100uM
1	2	14	9	0
1	2	1	10	0
1	2	2	10	0
1	2	3	10	M020 100uM

