

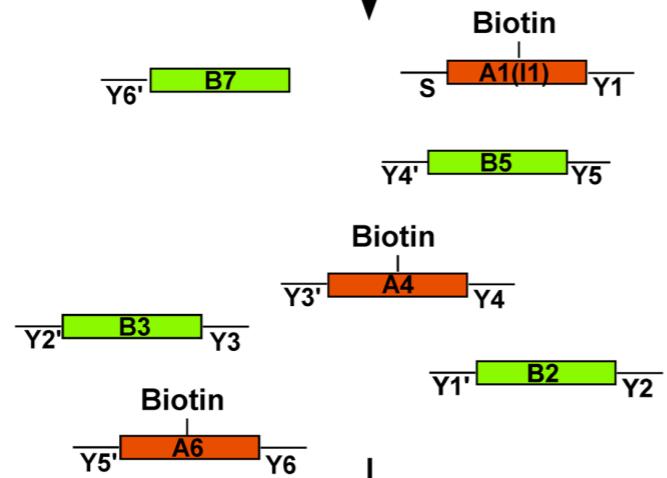
## Seed Formation

9 DNA strands per tile



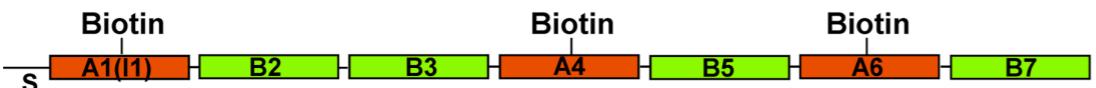
(1) Seed Tiles Annealed Separately

7 specific tiles with sticky ends



(2) Tiles Mixed Together

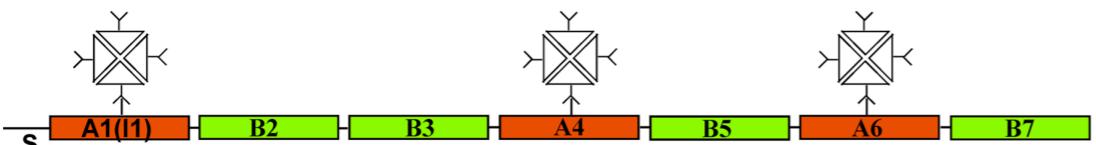
seeds



First Generation

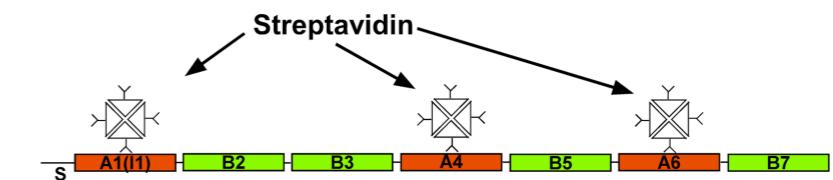
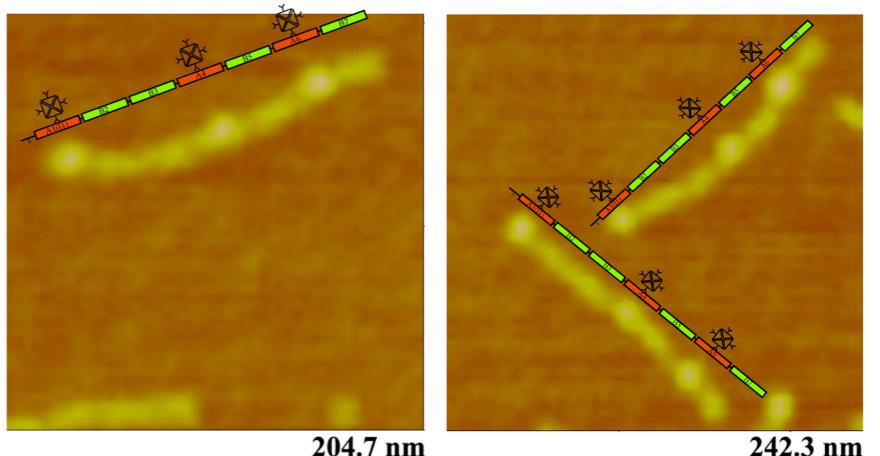
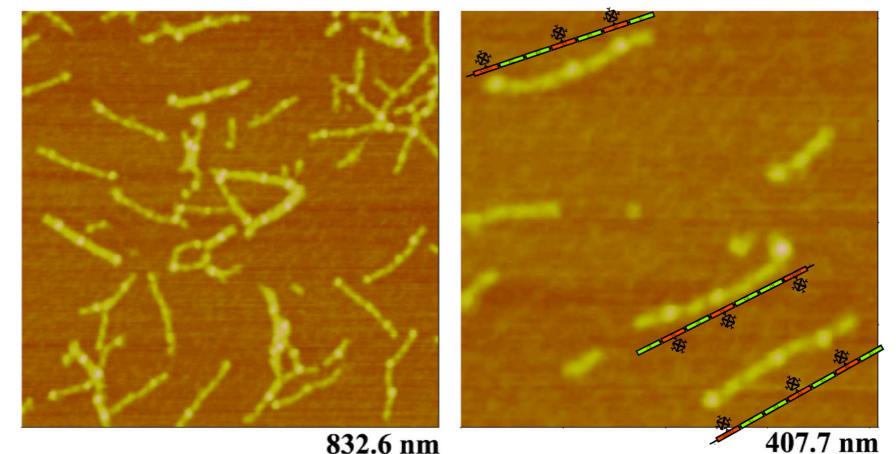


(3) Streptavidin Added

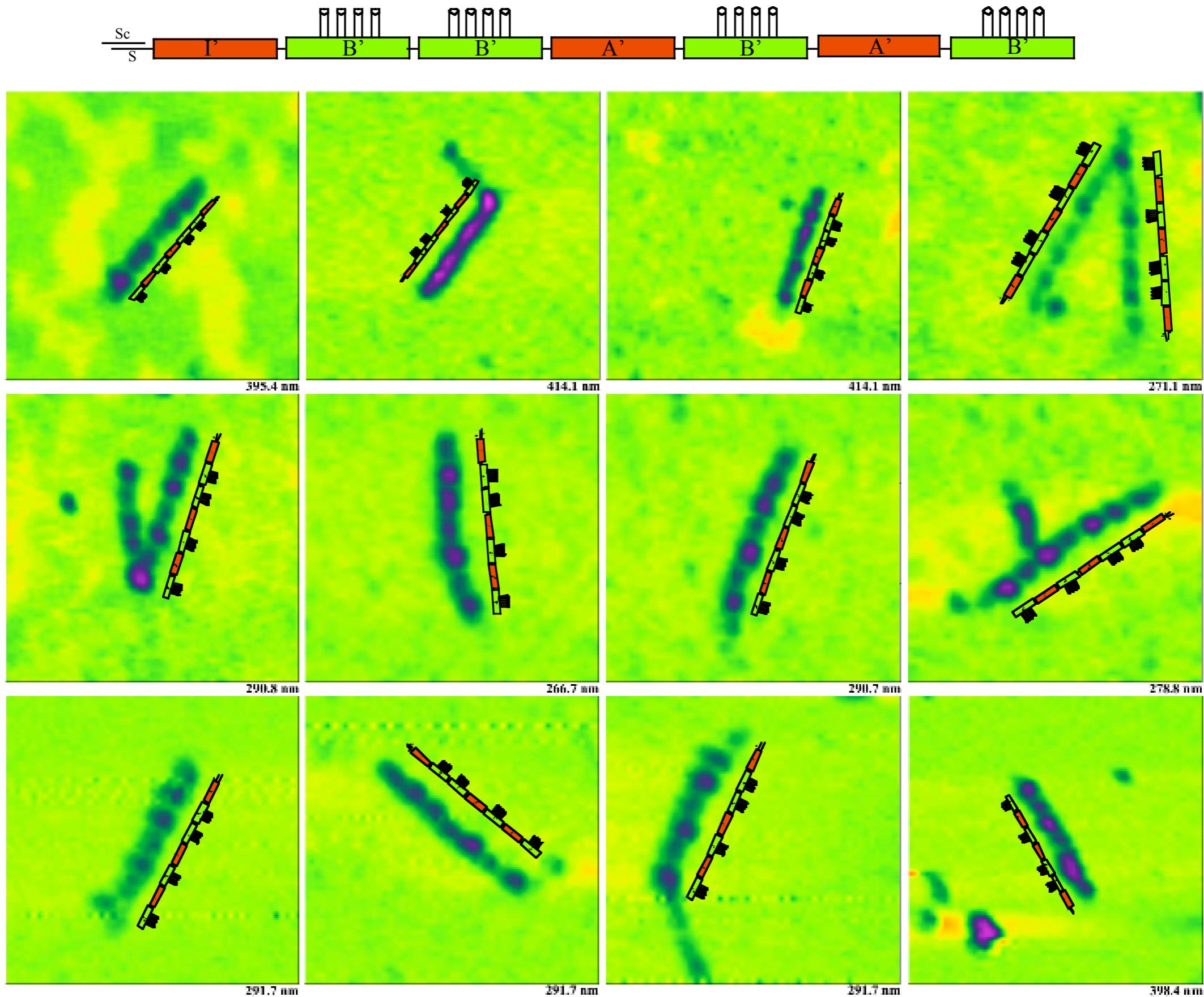


AFM Imaging

AFM images of seeds



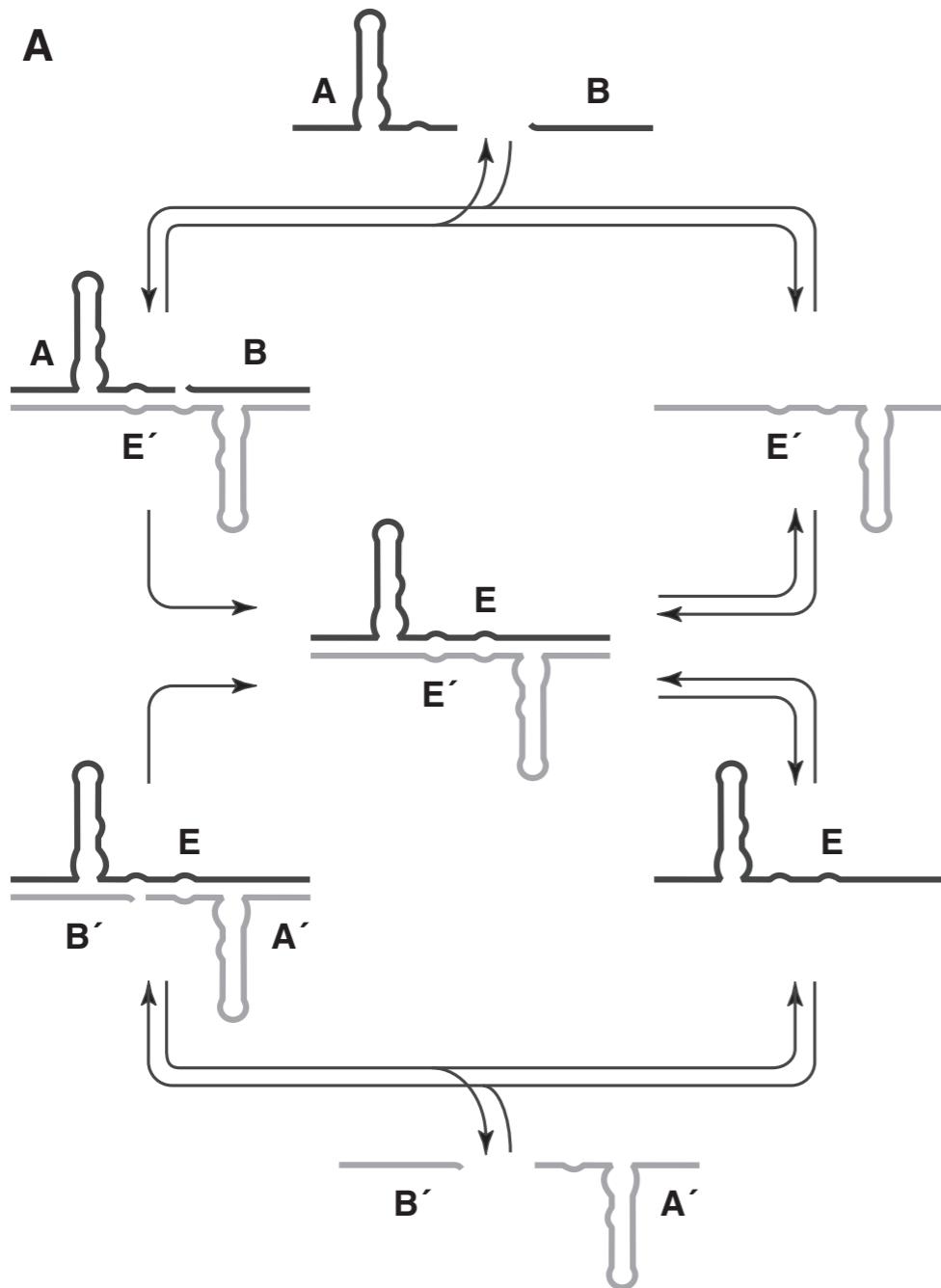
# Daughters Labeled with Hairpins ~ 60% yield



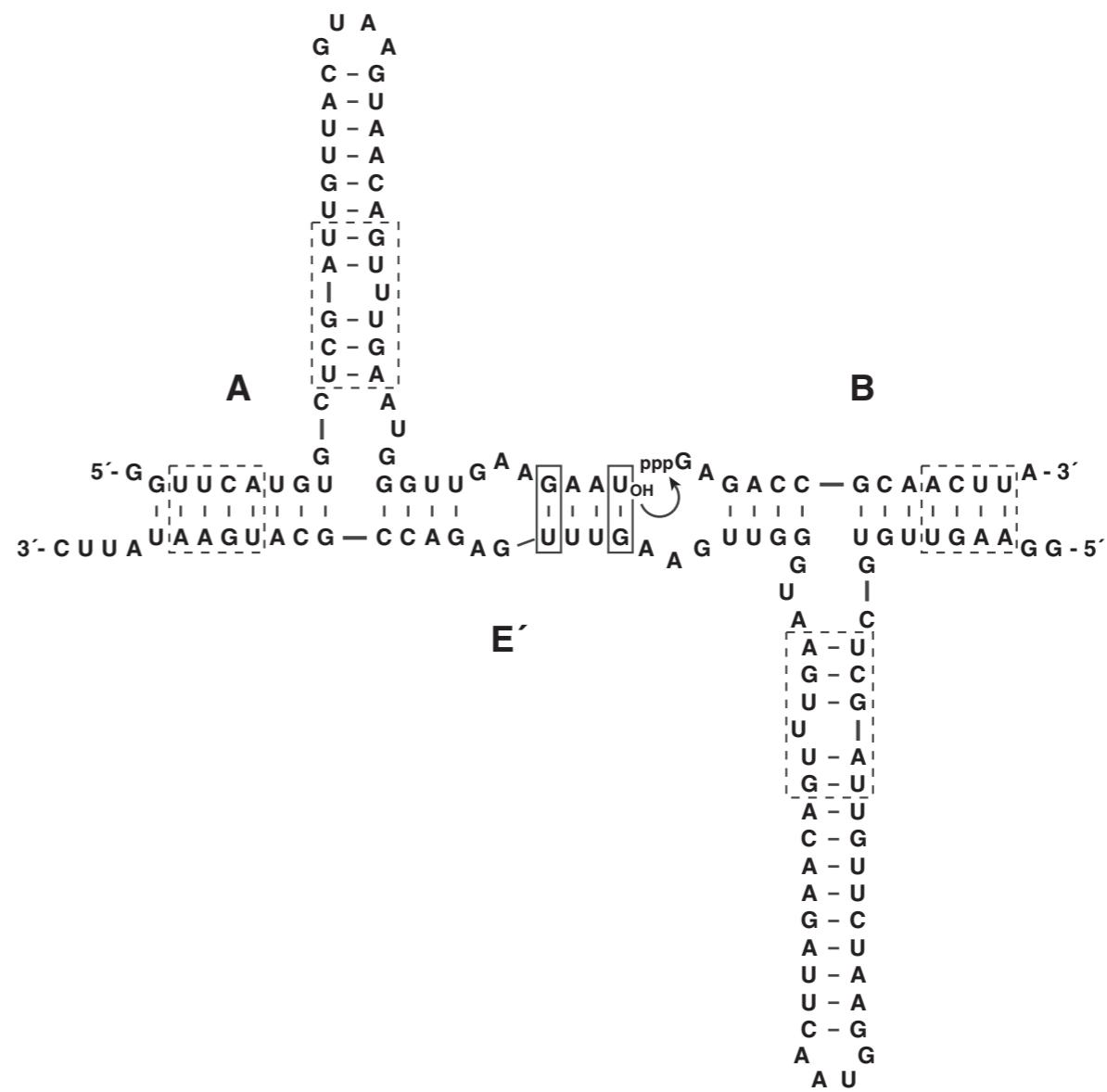
# Self-Sustained Replication of an RNA Enzyme

Tracey A. Lincoln and Gerald F. Joyce\*

SCIENCE, 323, 1229, 2009

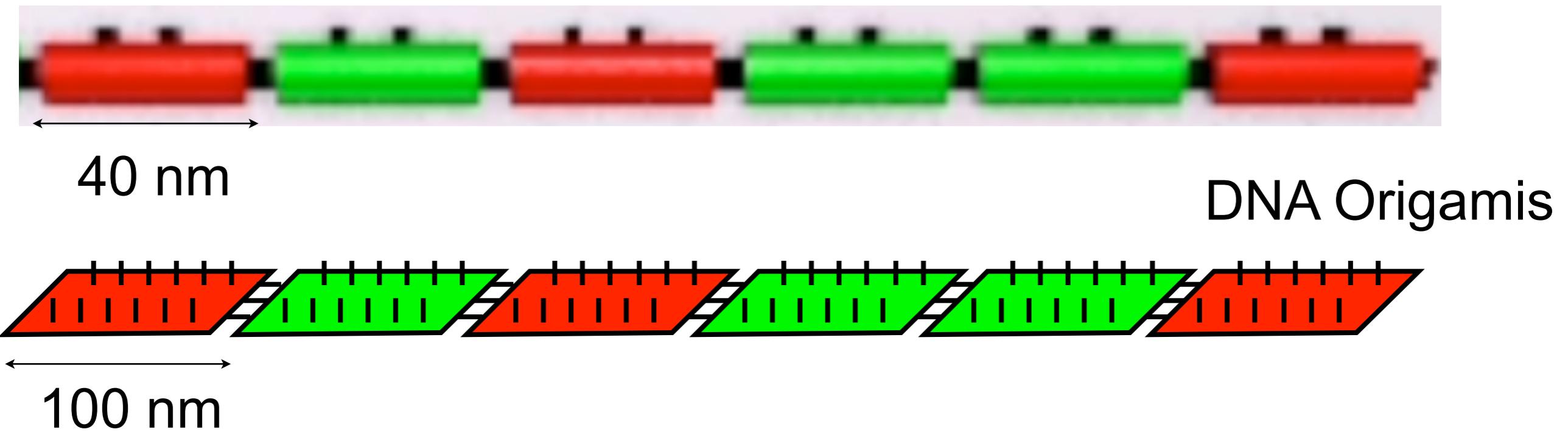
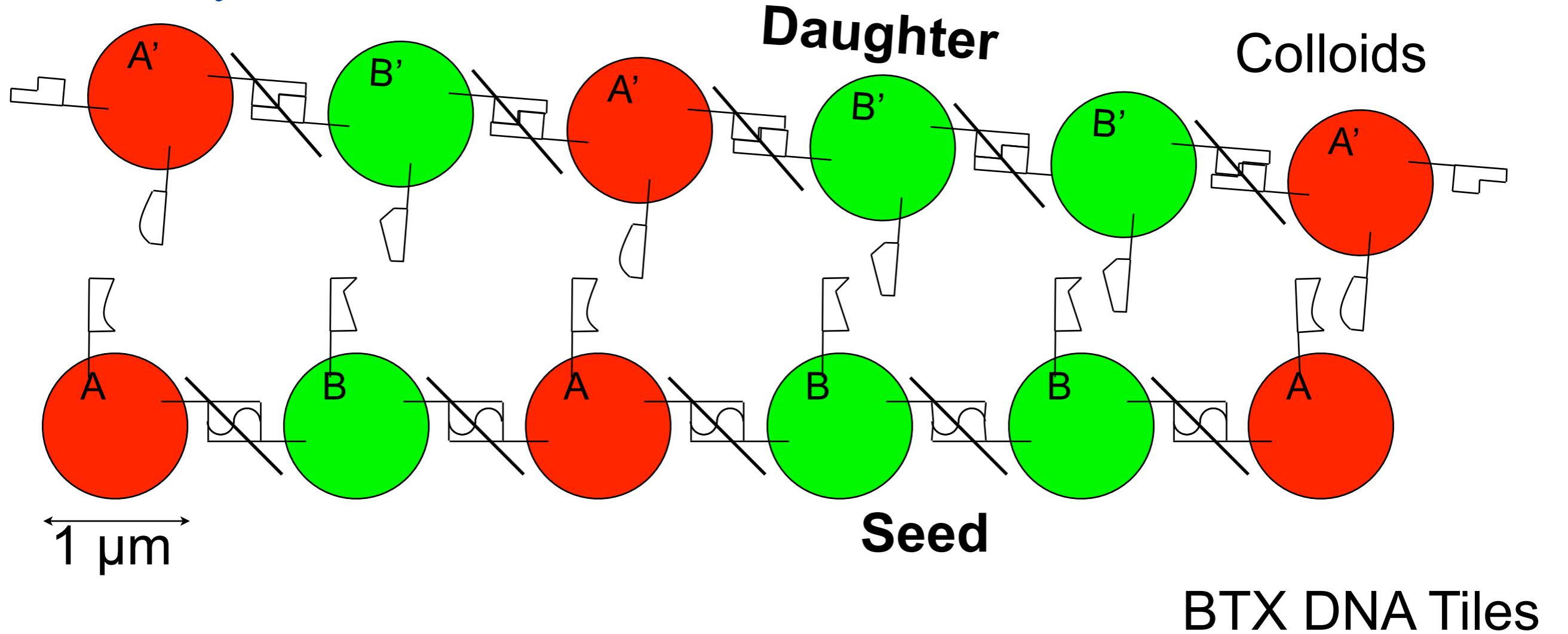


**B**



Why don't we try two tiles? And do it with Origamis

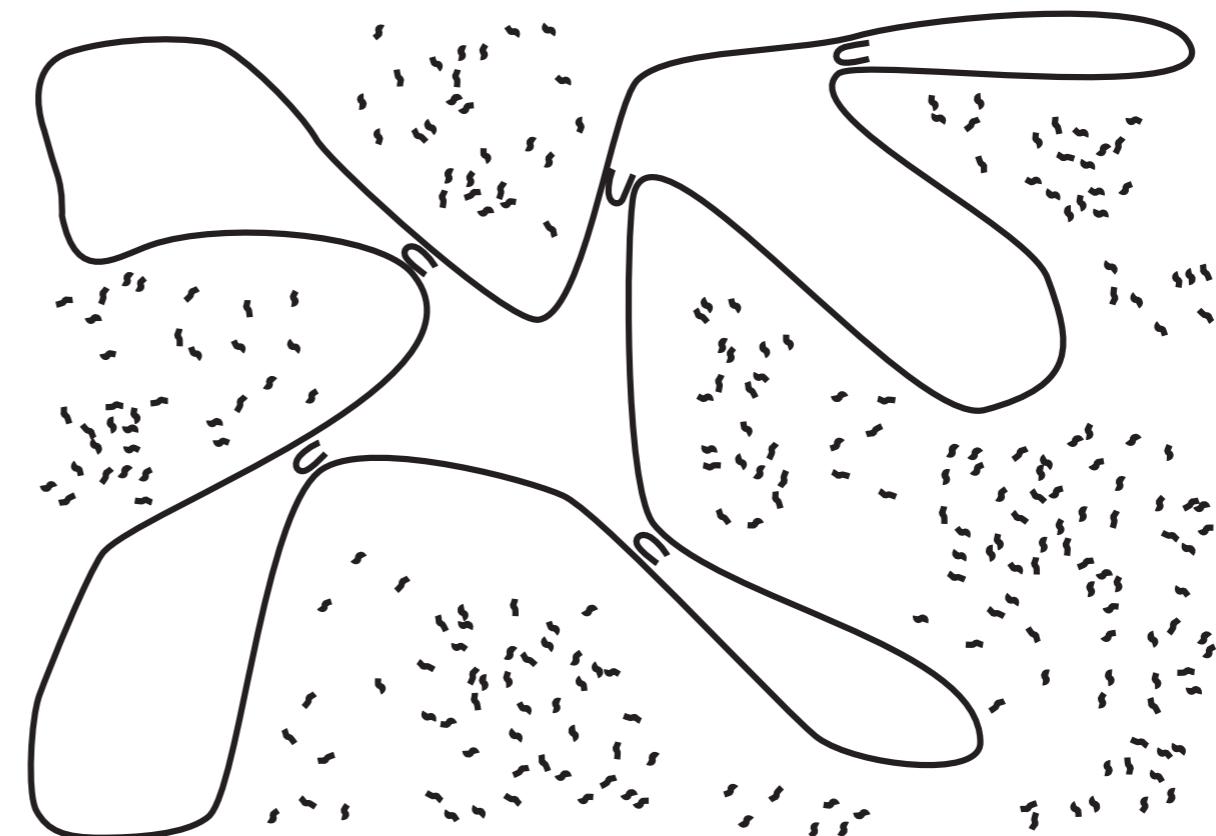
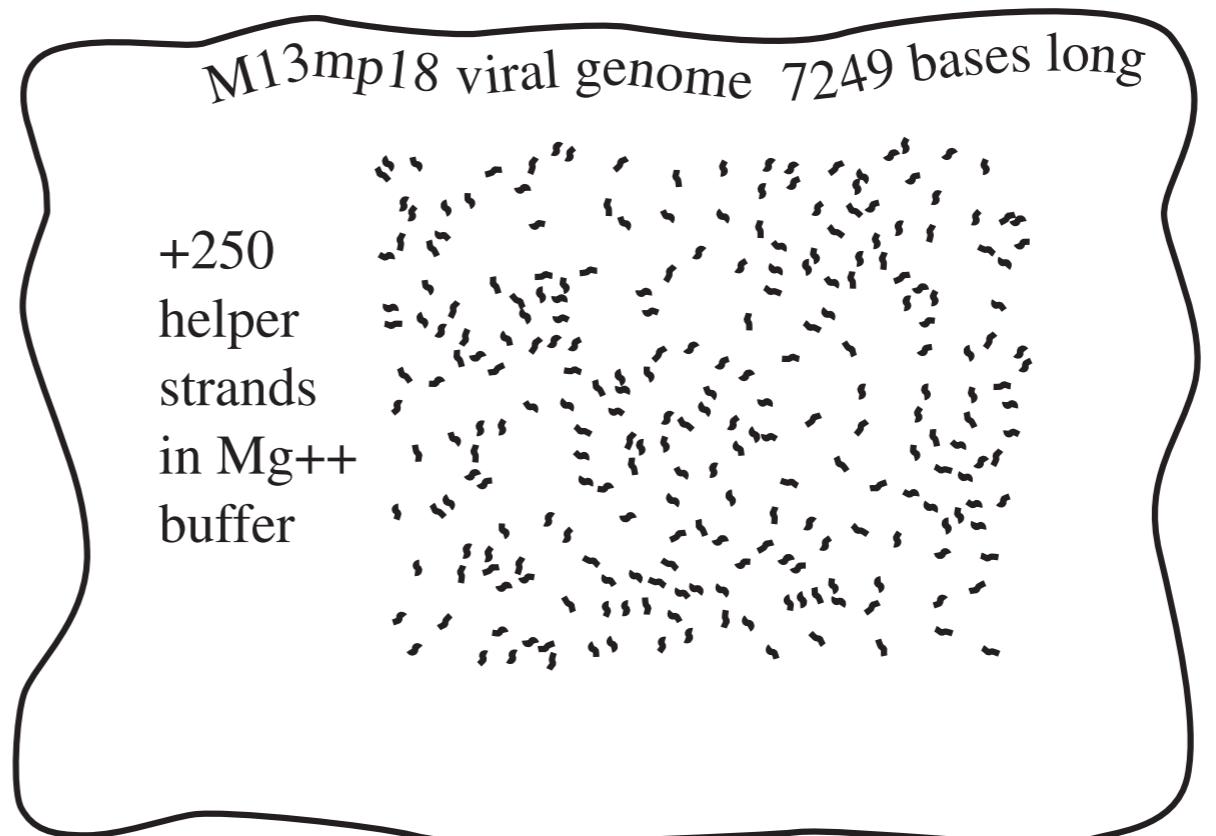
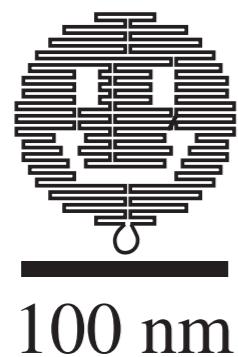
# Different systems, same idea



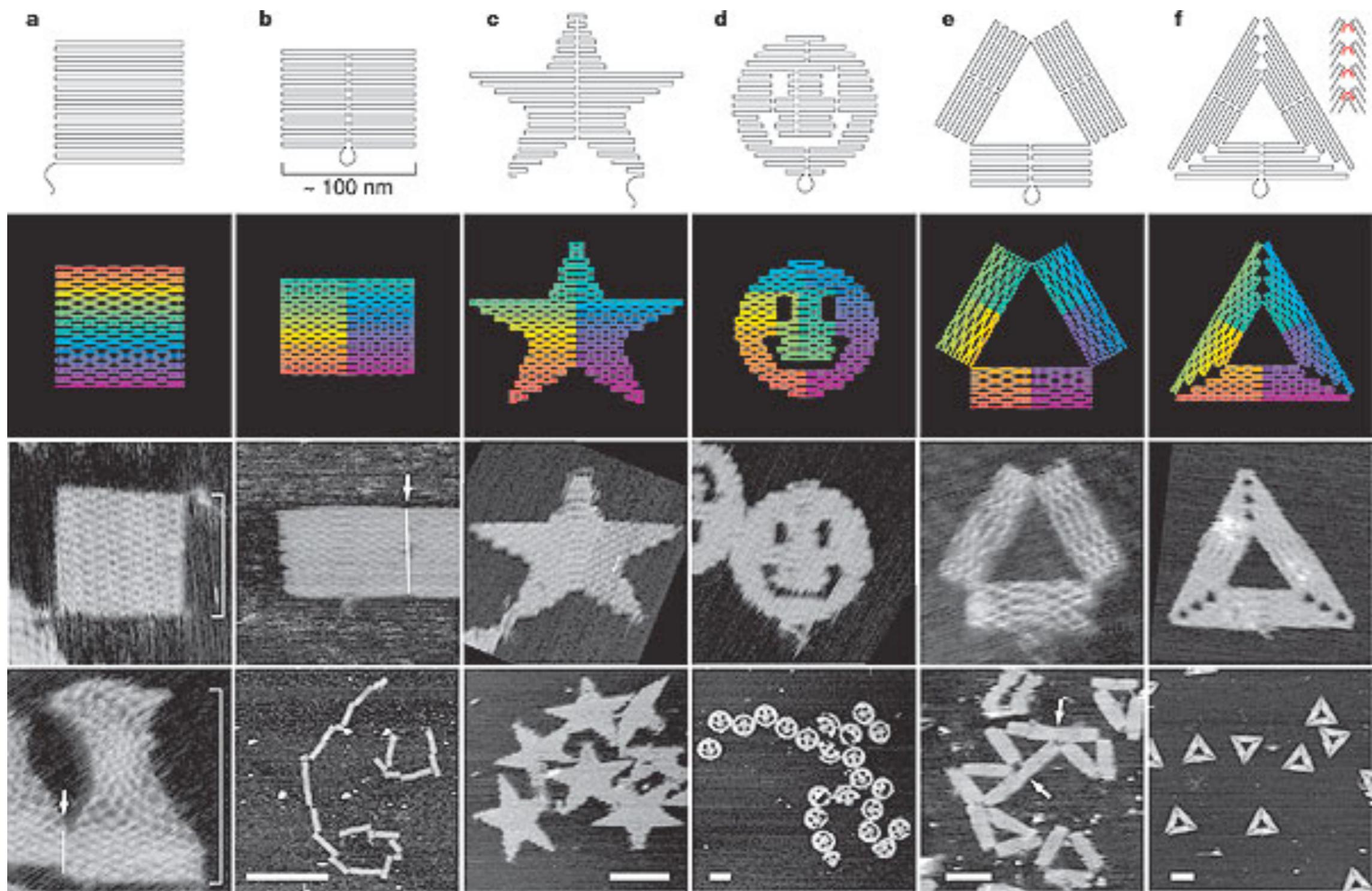
# Design of DNA origami

Paul W.K. Rothemund

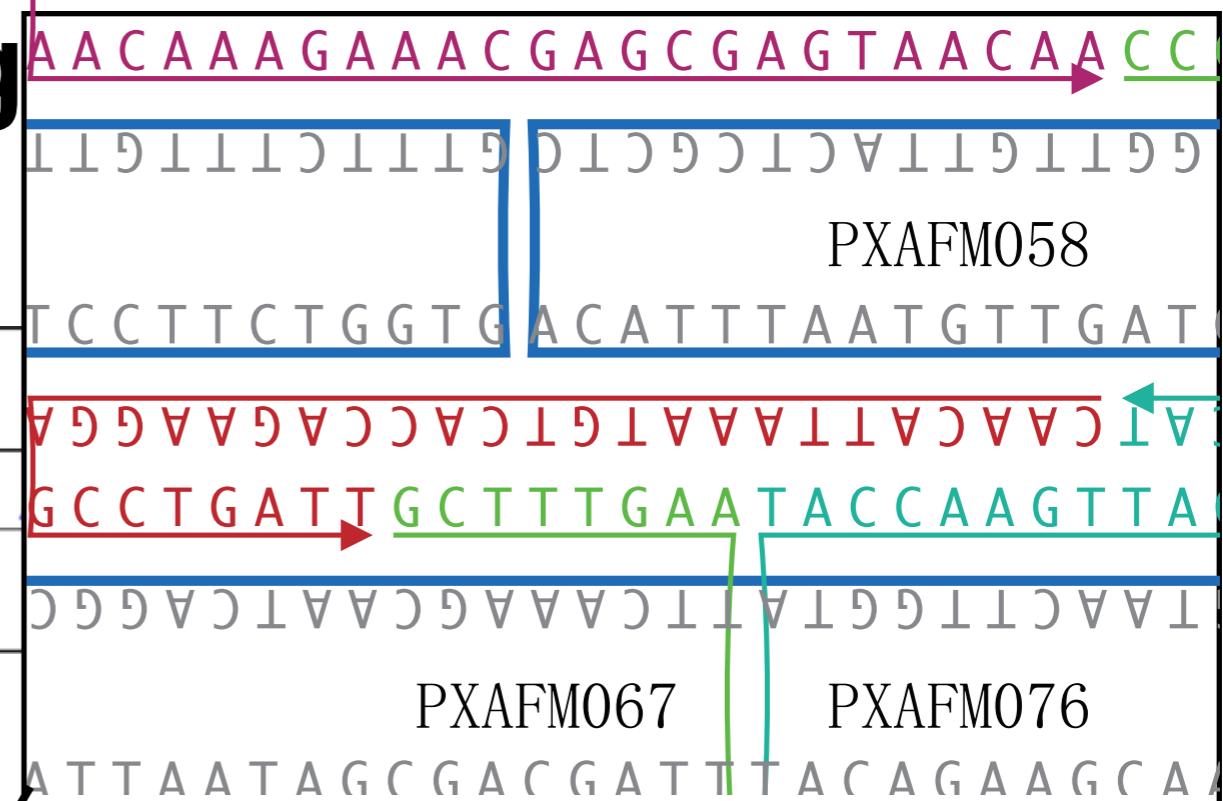
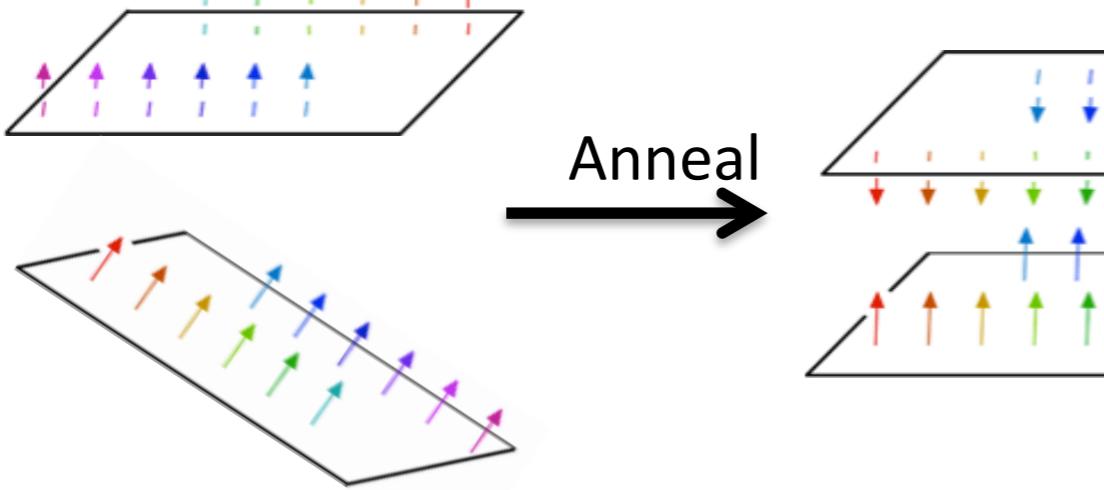
Computer Science and Computation and Neural Systems  
California Institute of Technology, Pasadena, CA 91125  
pwkr@dna.caltech.edu

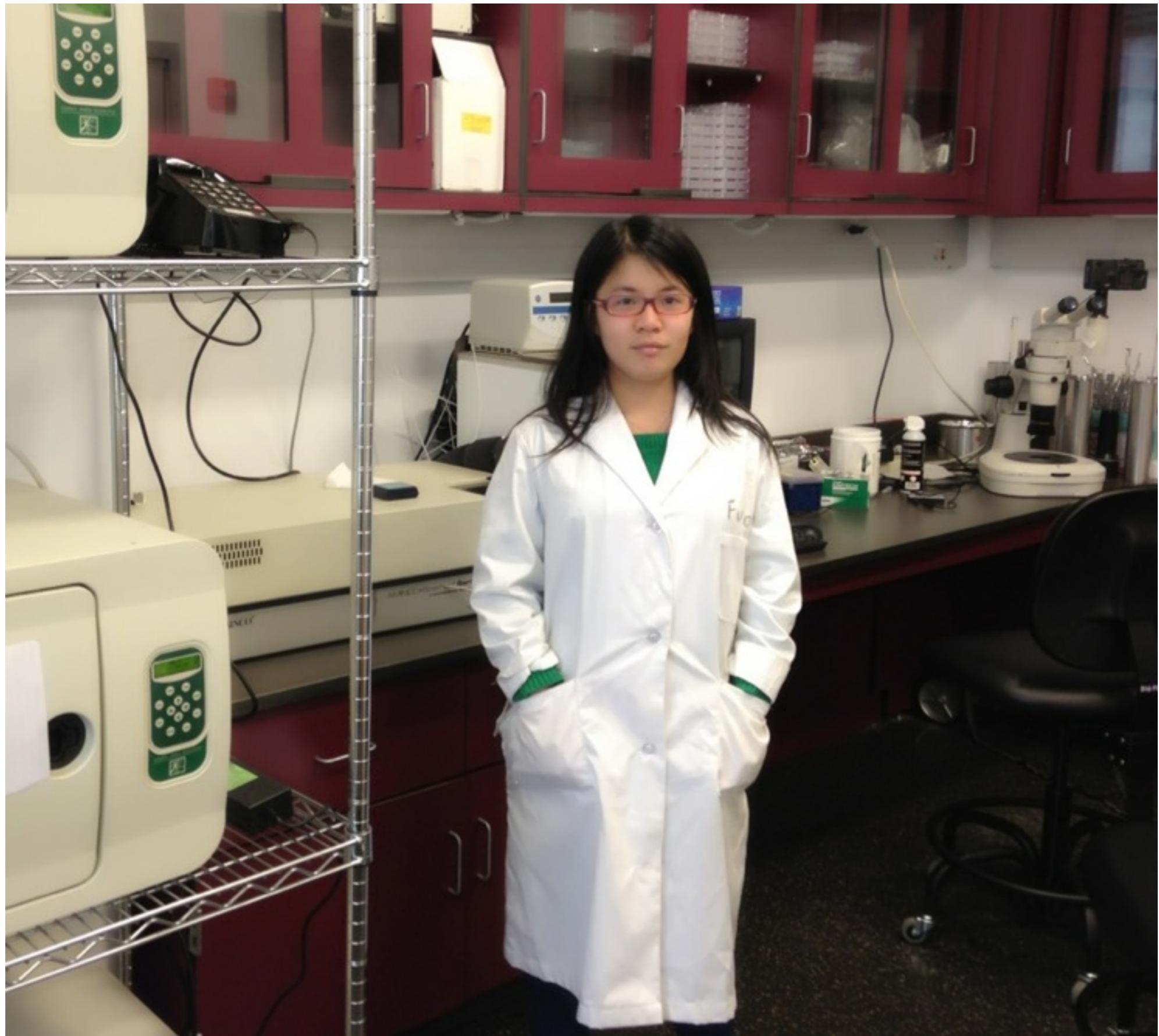


# Rothemund's DNA Origamis



# DNA Origami Design

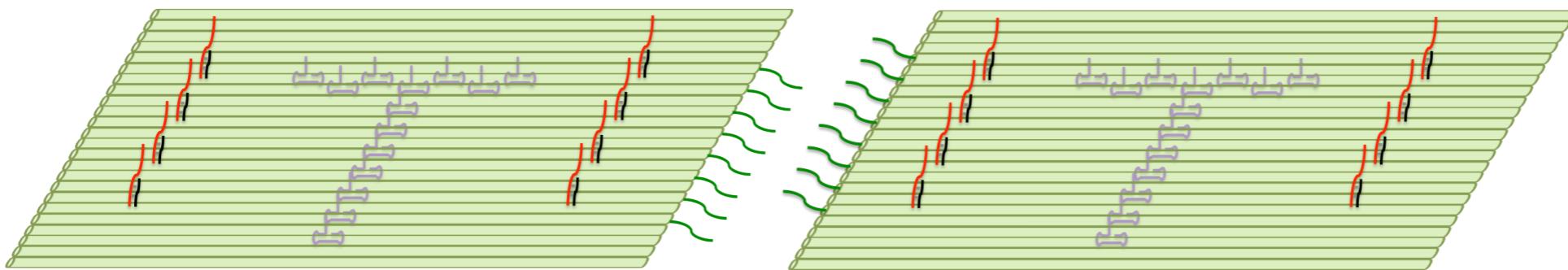




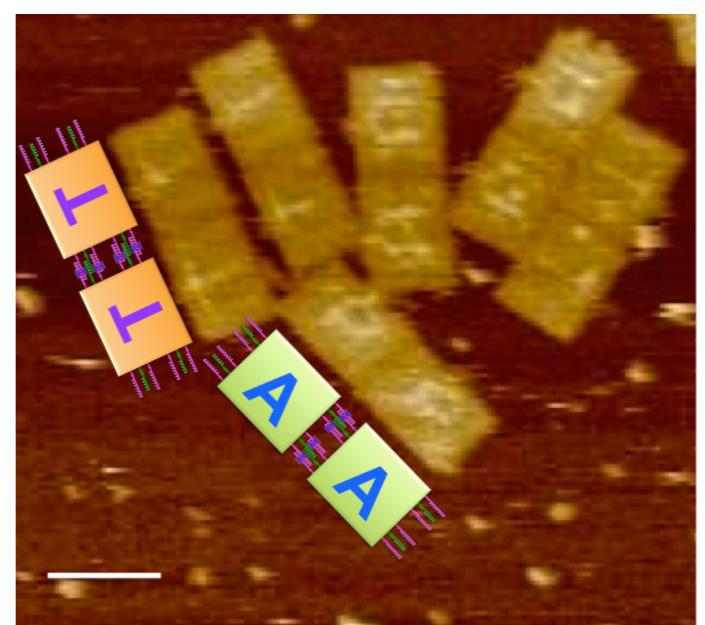
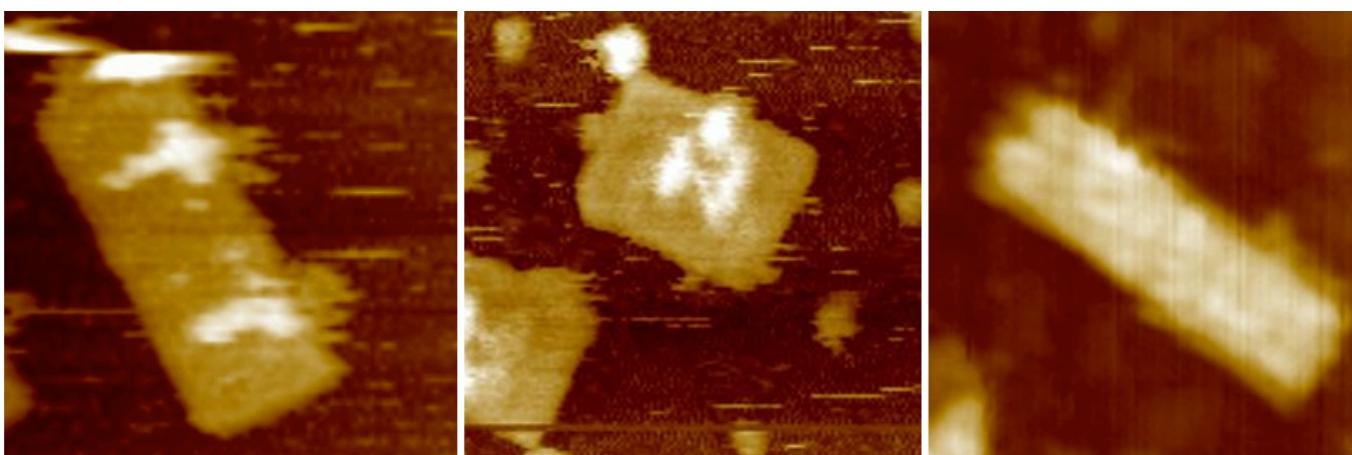
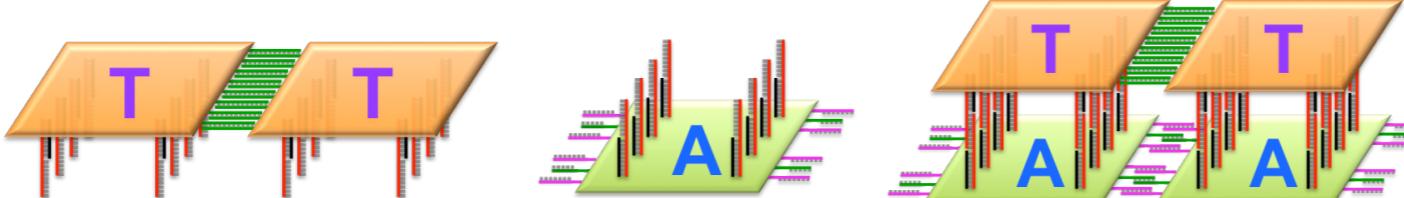
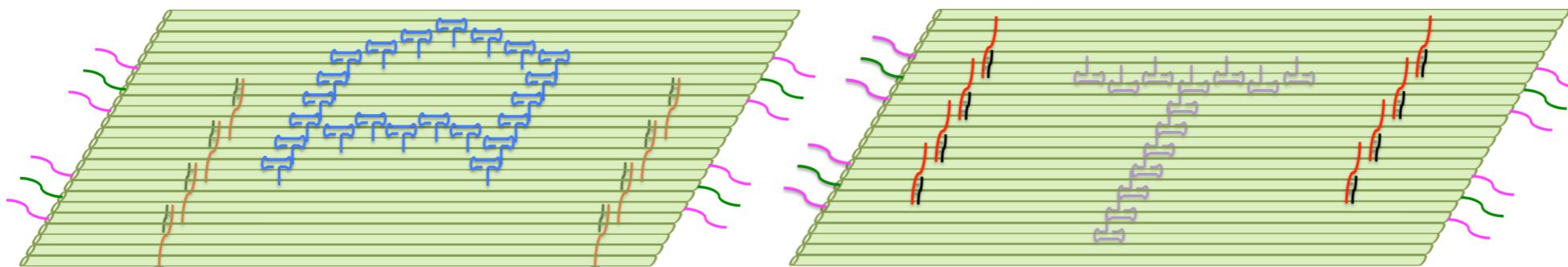
Xiaojin He      HKUST/NYU

# Basic tile set for self-replication

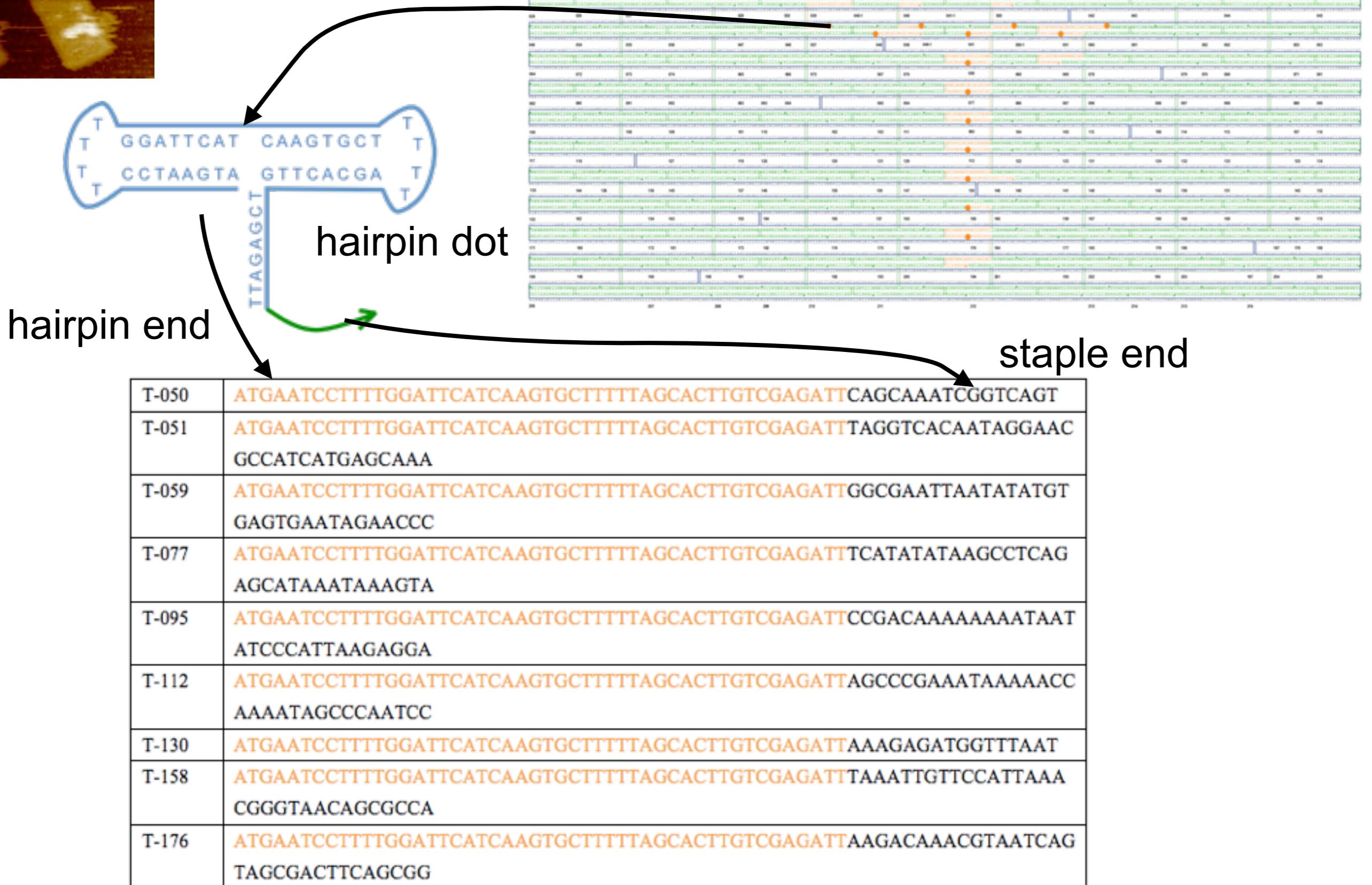
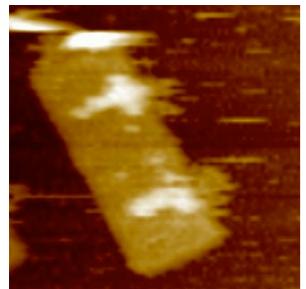
## SEED TILES

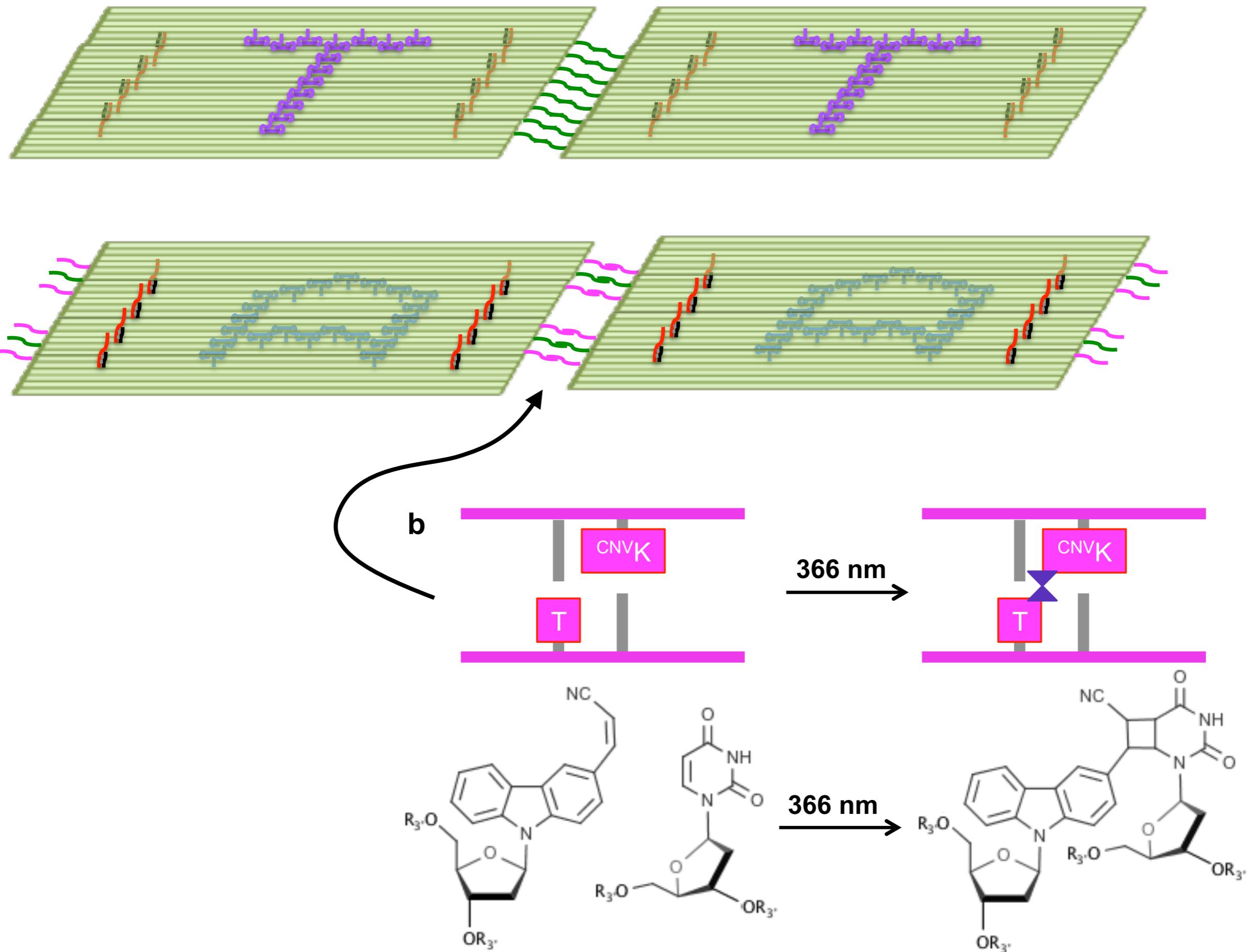


## LATER-GENERATION TILES

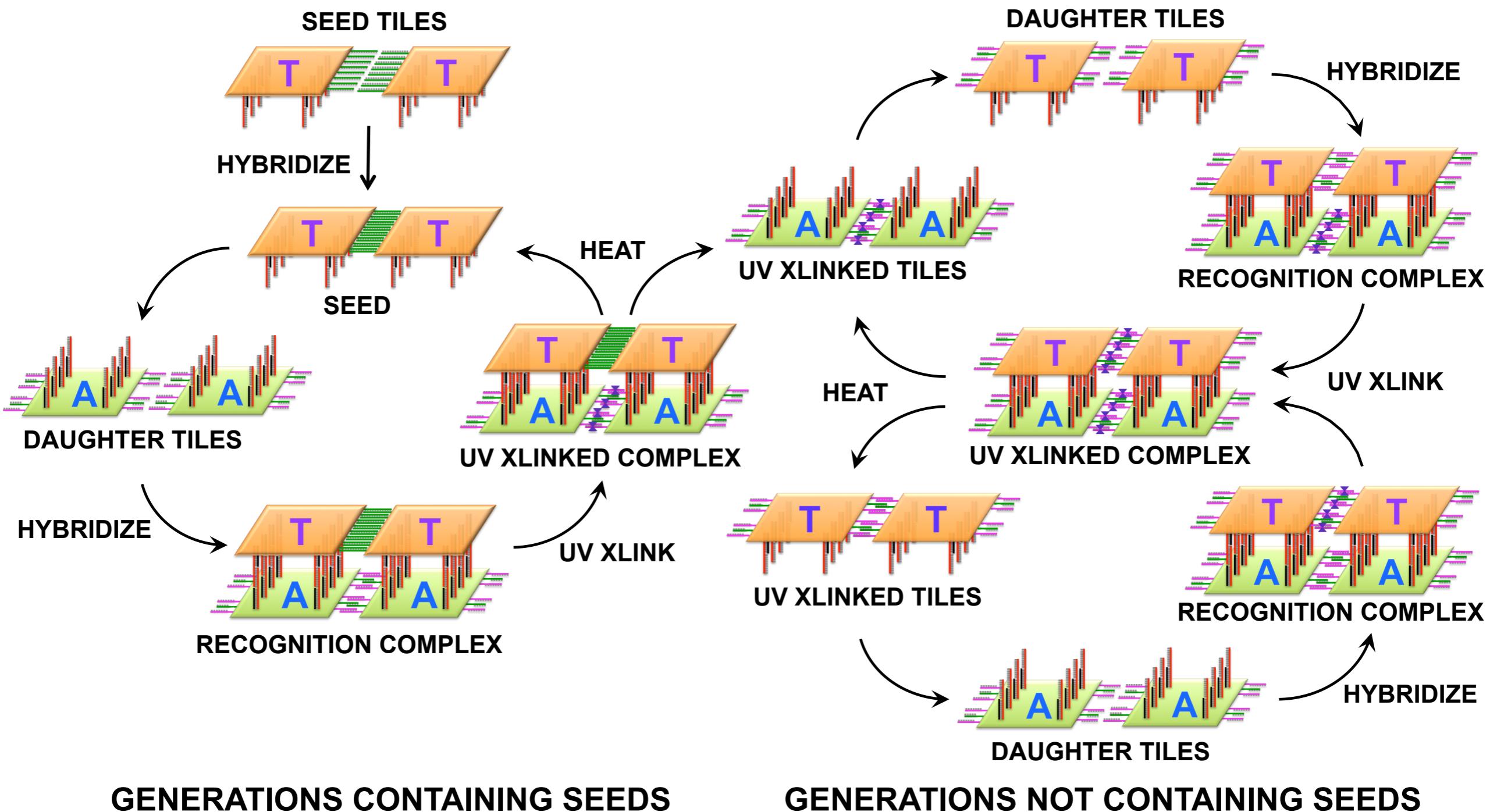


# Labelling with letter “T”

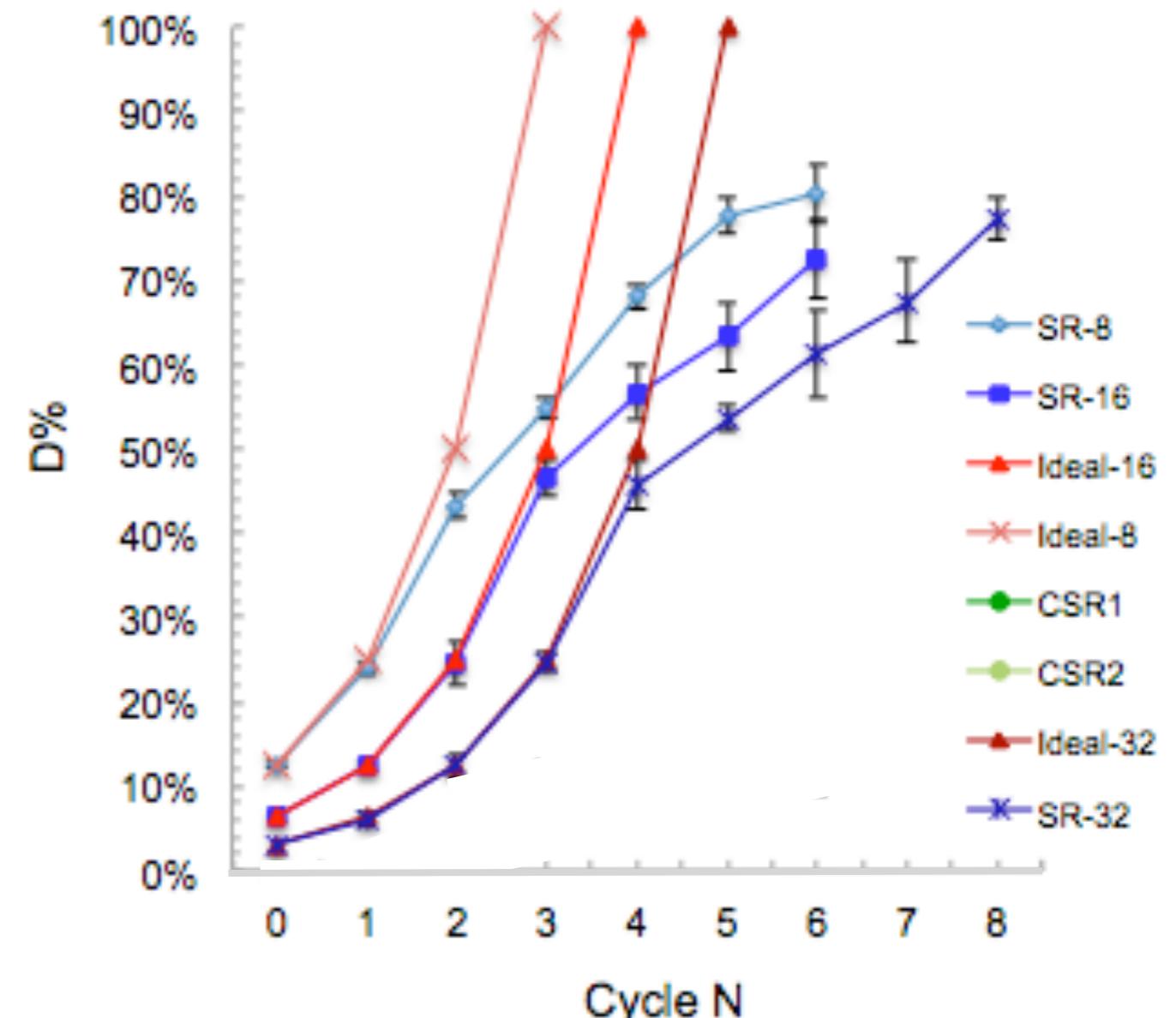
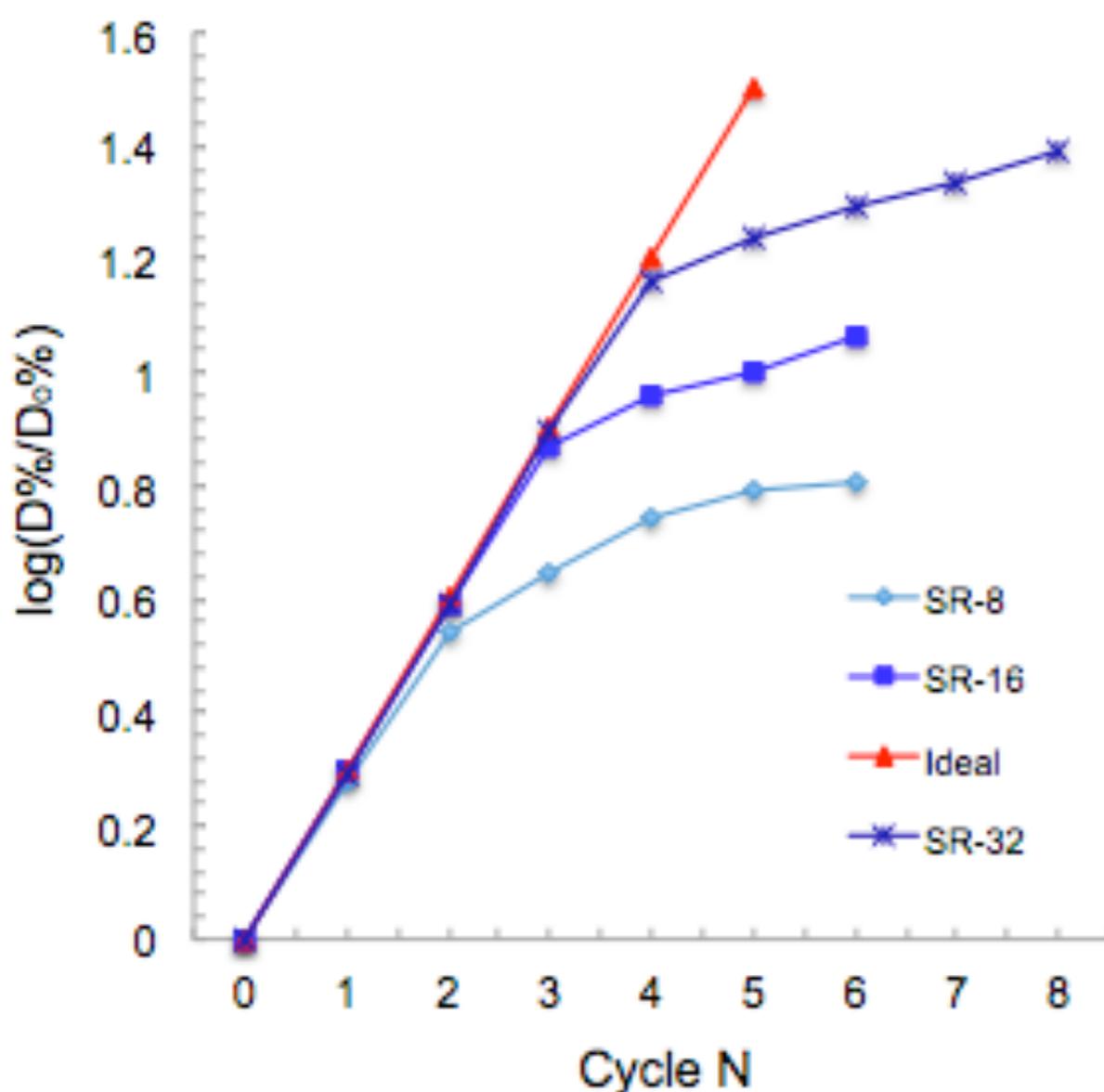




# Replication cycles - cool/UV/heat -repeat

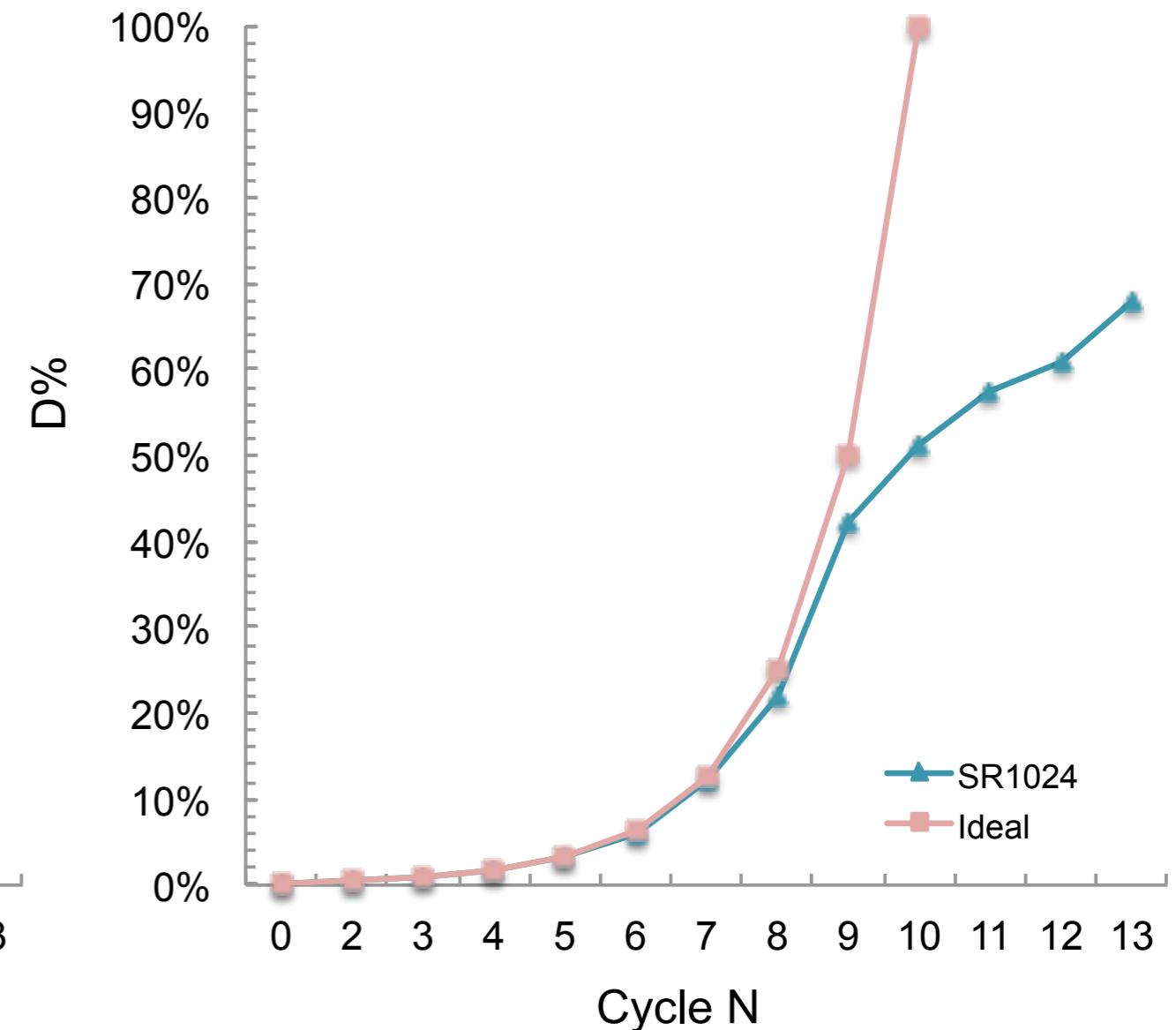
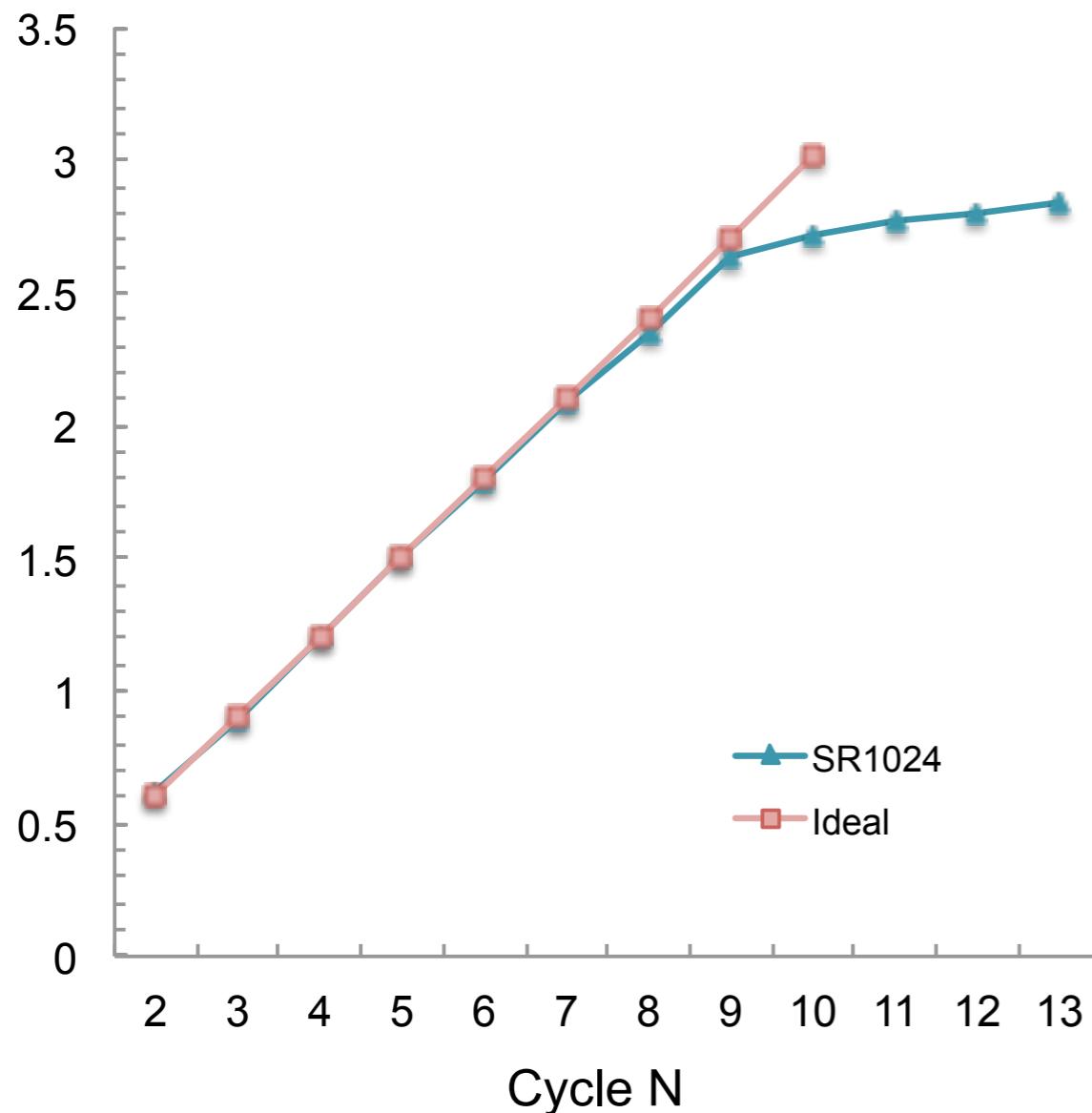


# Number of Dimers Doubles each cycle!



Here's 500X multiplication of seed

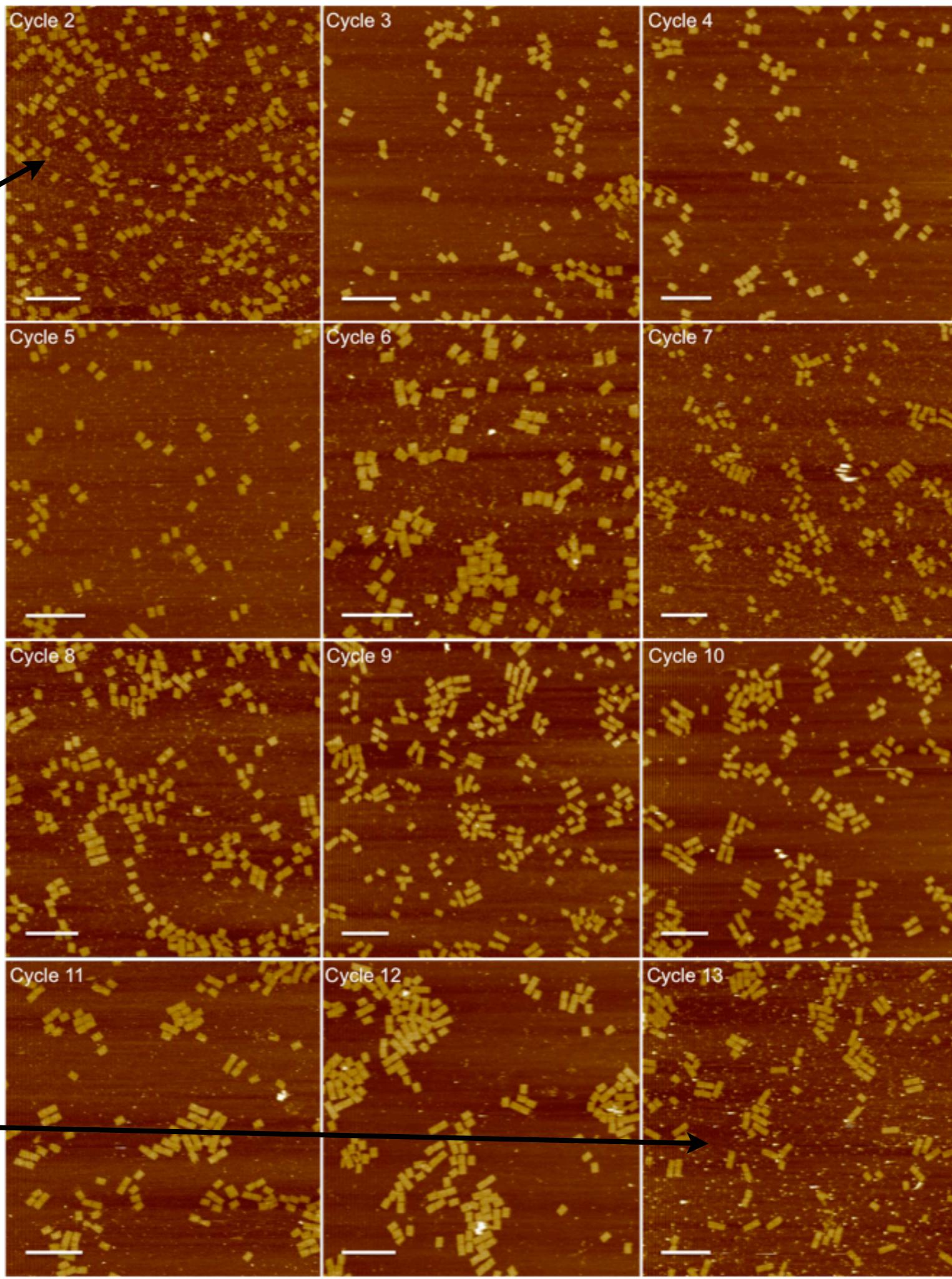
## Self-Replication Plot (1:1024)



# AFM Images of 1024 replication

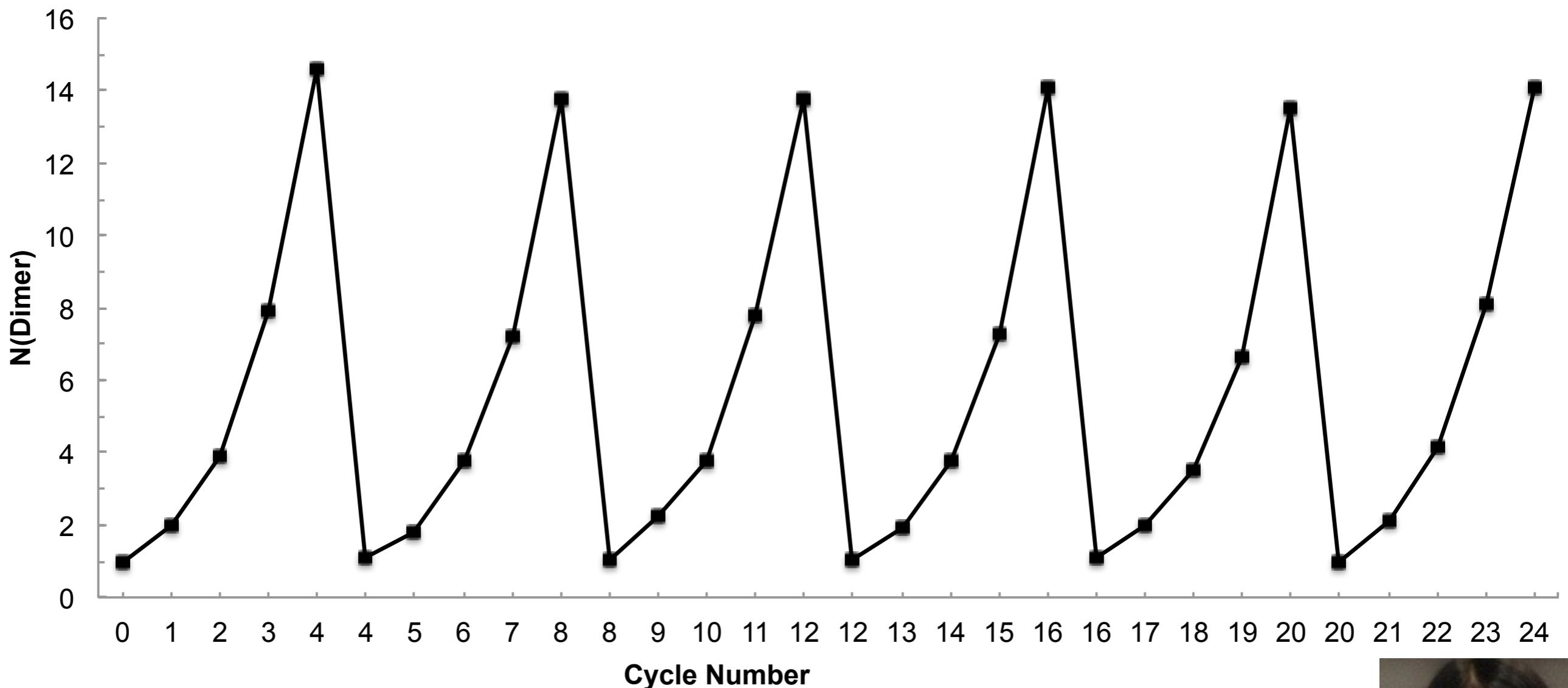
mostly single tiles

mostly dimer tiles



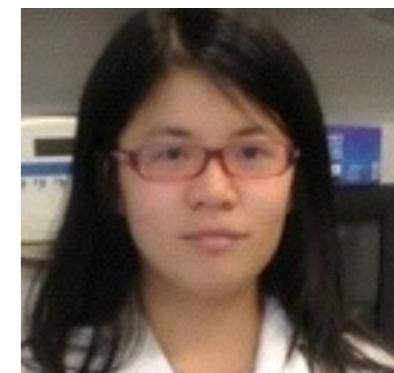
# Replication by Serial Dilution

- ◆ Use the self-replicated sample (ratio: 1:32) after four cycles
- ◆ Allow approximately 14-fold amplification before transferring ~6% of the mixture to a new reaction tube that contained a fresh supply of monomers.

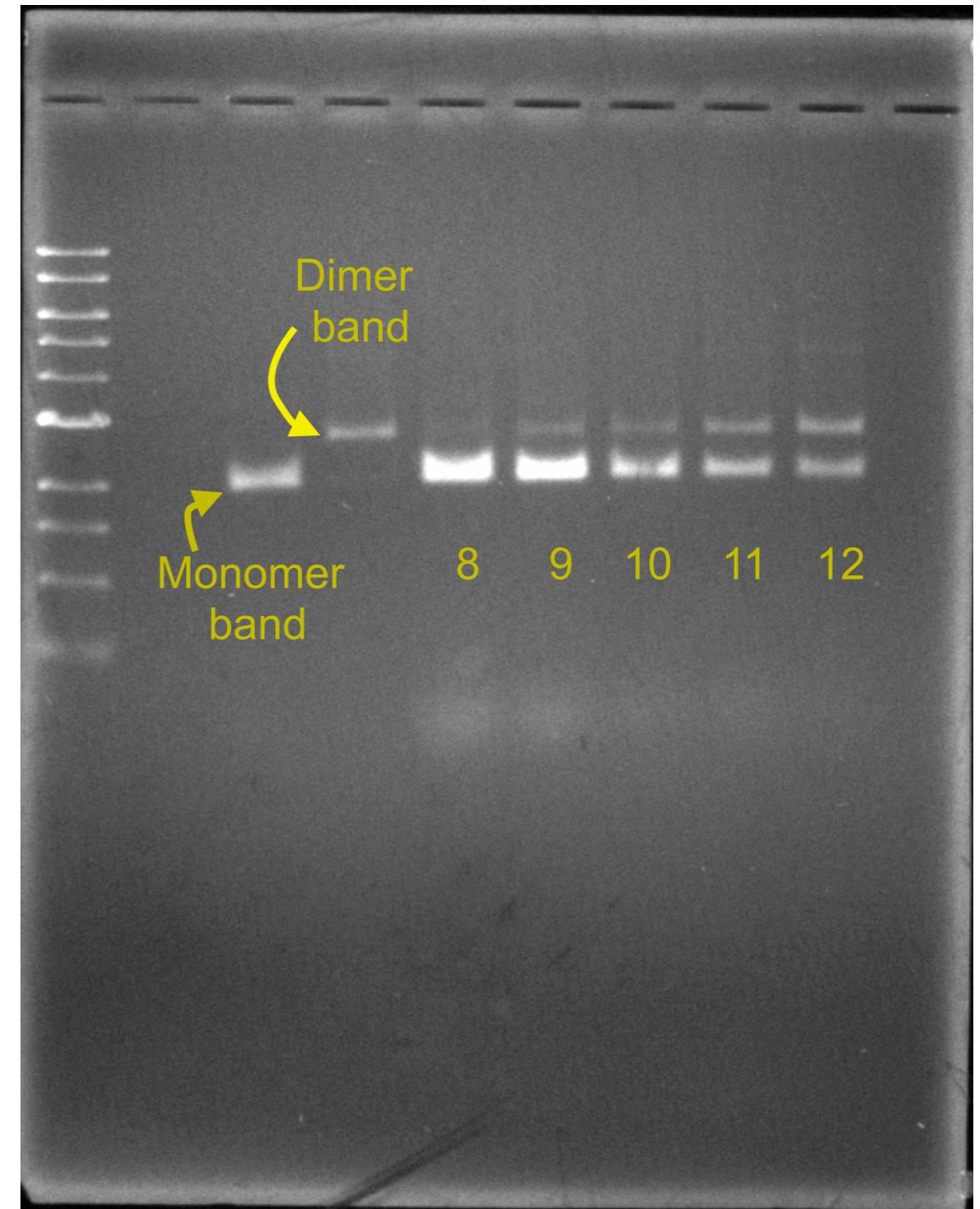
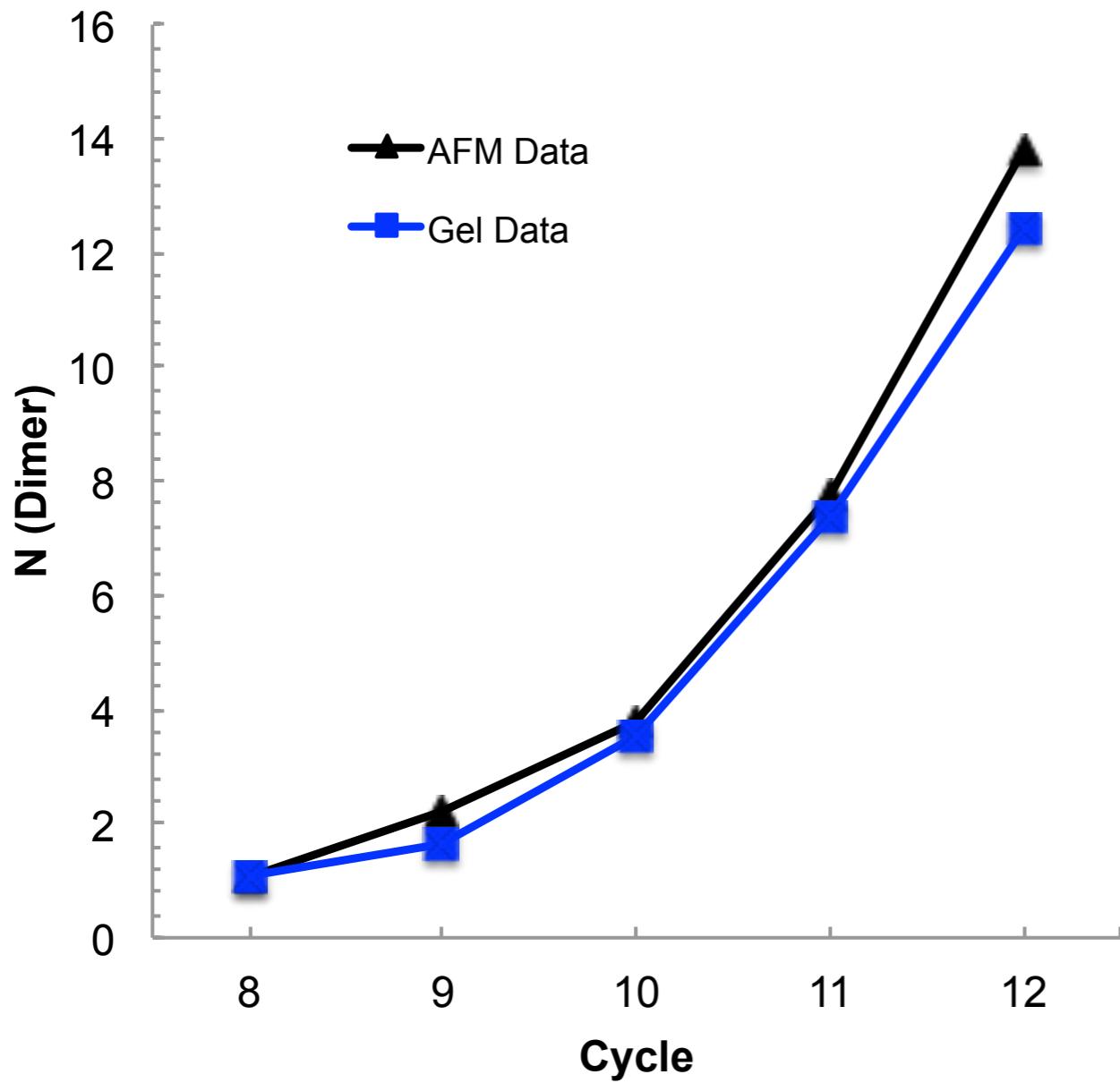


Total Amplification 7.5 million

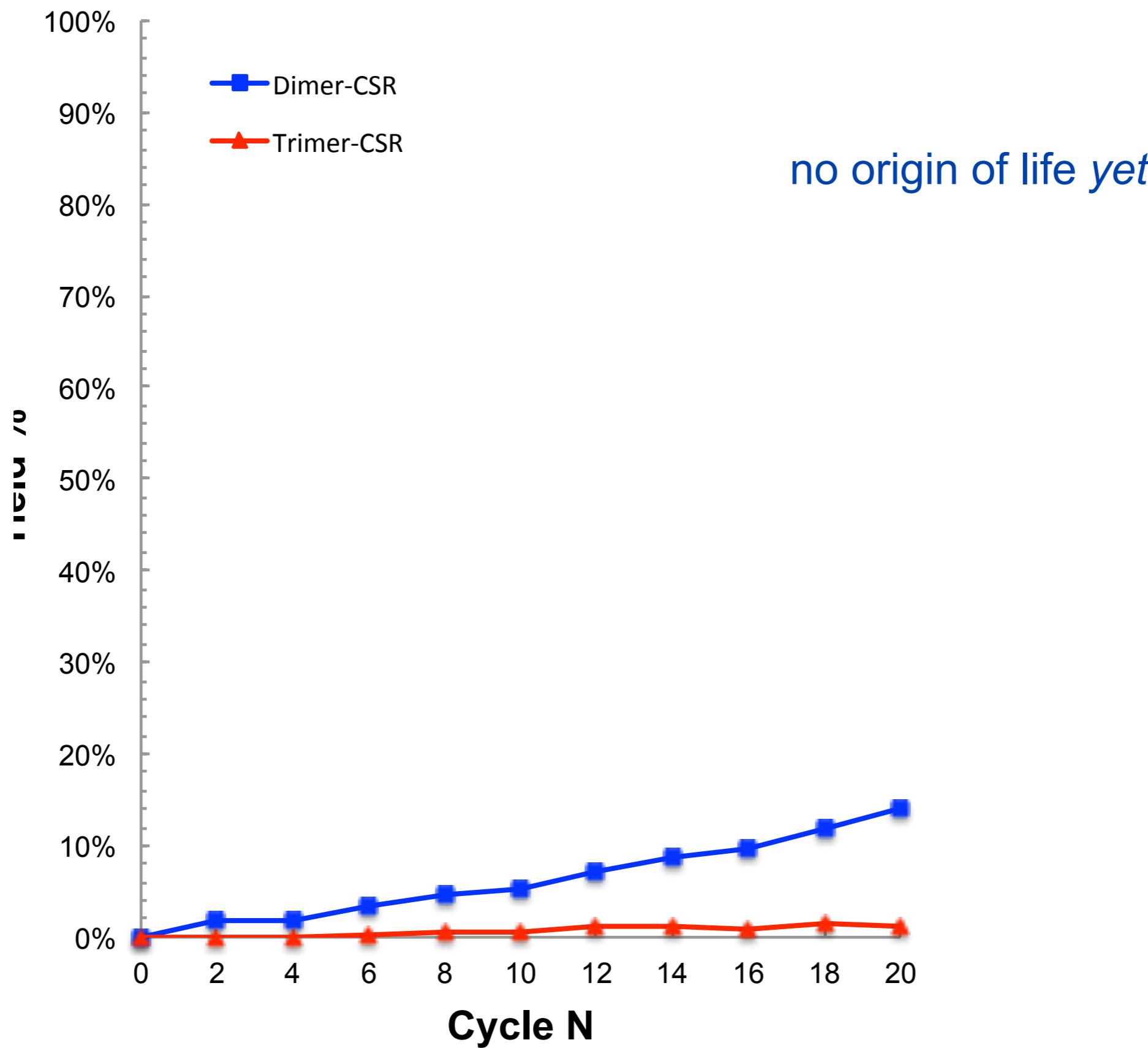
Xiaojin  
He



# Gel analysis shows same growth as counting Origami



# Temperature and light cycles NO SEED



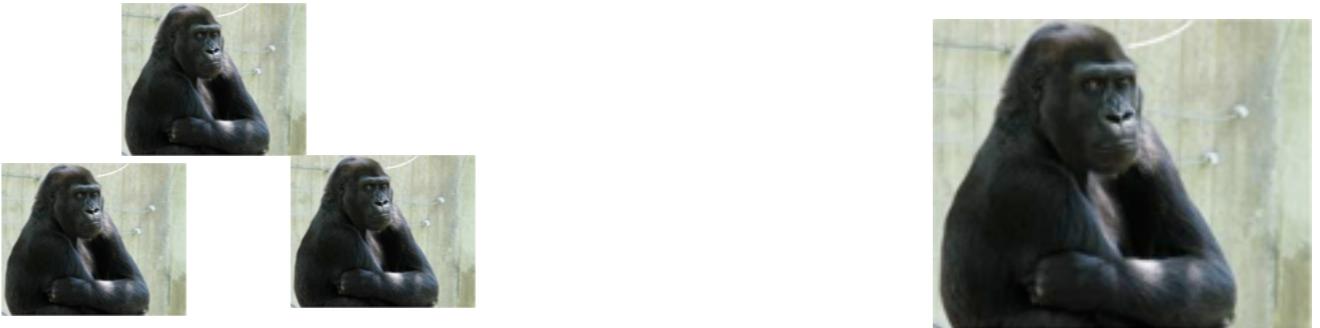
## [Evolution - Wikipedia, the free encyclopedia](#)

[en.wikipedia.org/wiki/Evolution](https://en.wikipedia.org/w/index.php?title=Evolution&oldid=98300000) ▾

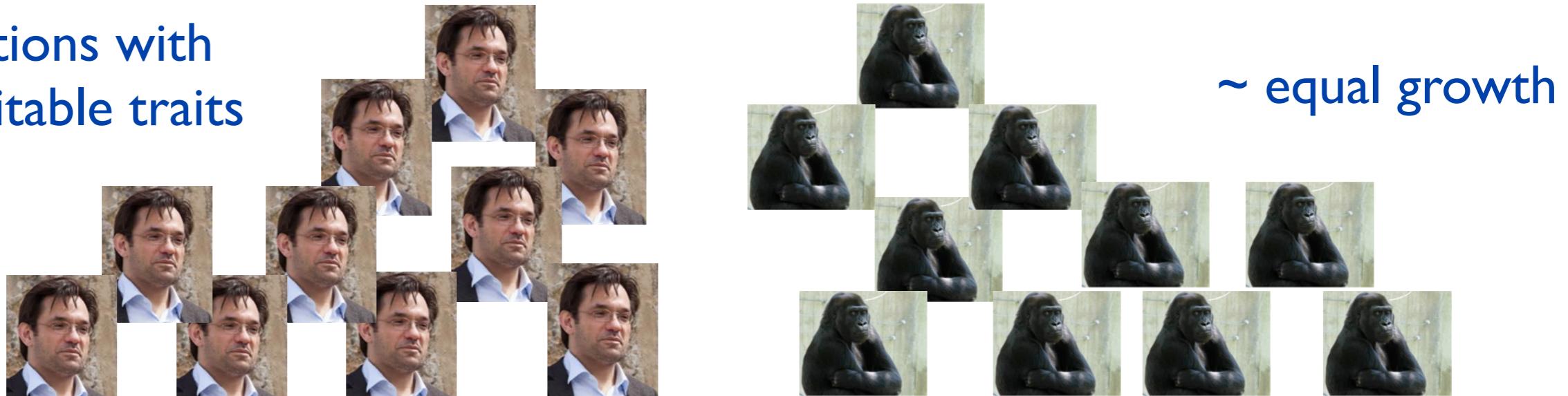
**Evolution** is the change in the inherited characteristics of biological populations over successive generations. **Evolutionary** processes give rise to diversity at ...

# Schematic Evolution

Original Species



Mutations with  
inheritable traits



~ equal growth

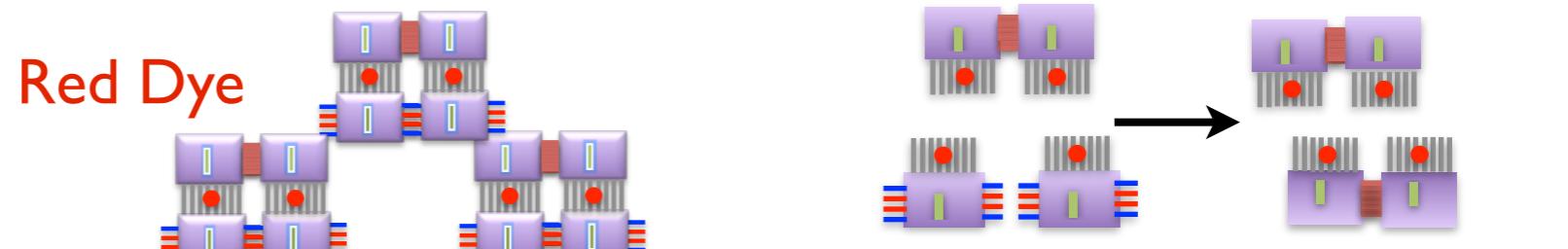
Environment Changes - Fire - need theory of Plasmas - advantage to one species



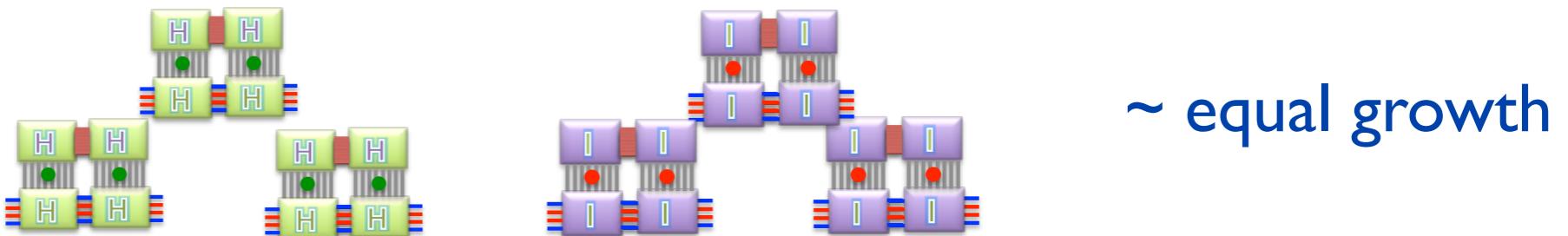
Environment Changes - advantage to one species  
growth rate higher - species takes over

# Red - Green Origami Evolution

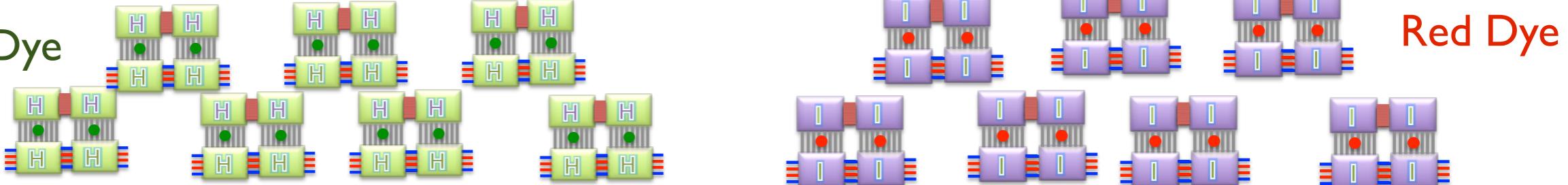
Original Species



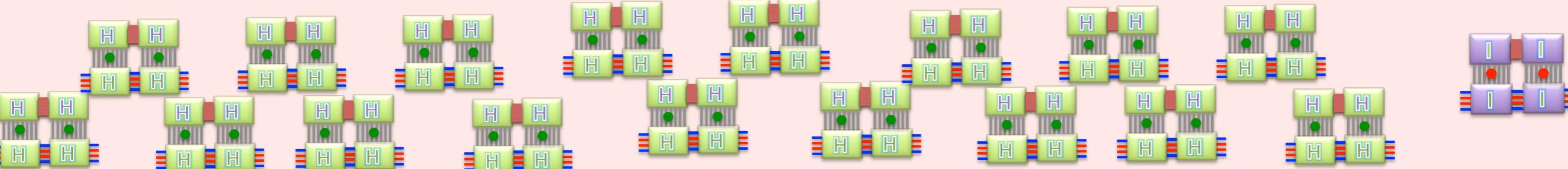
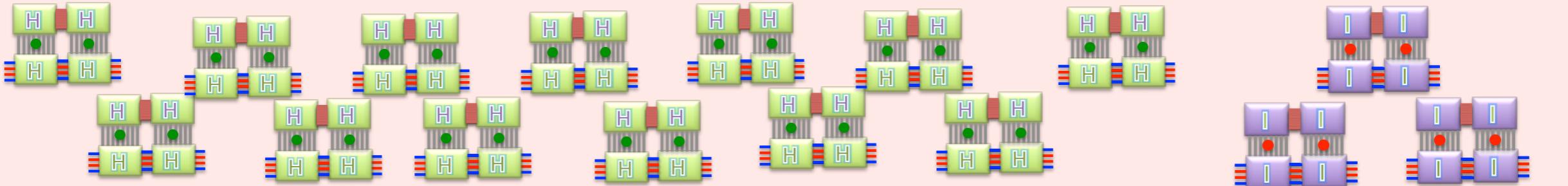
Mutations with  
inheritable traits



Green Dye



Environment Changes - Red Light - advantage to one species



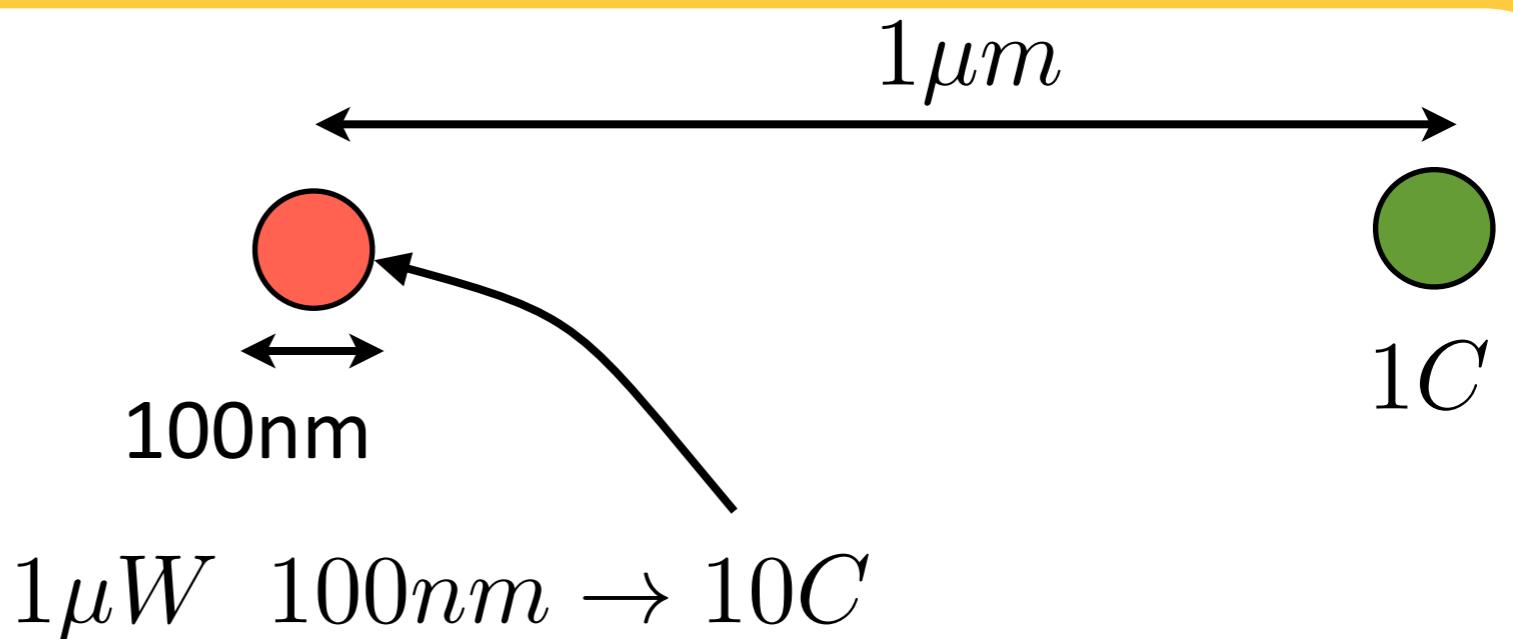
Selection - higher growth rate - Green takes over

# Laser Heating of IR Dyes

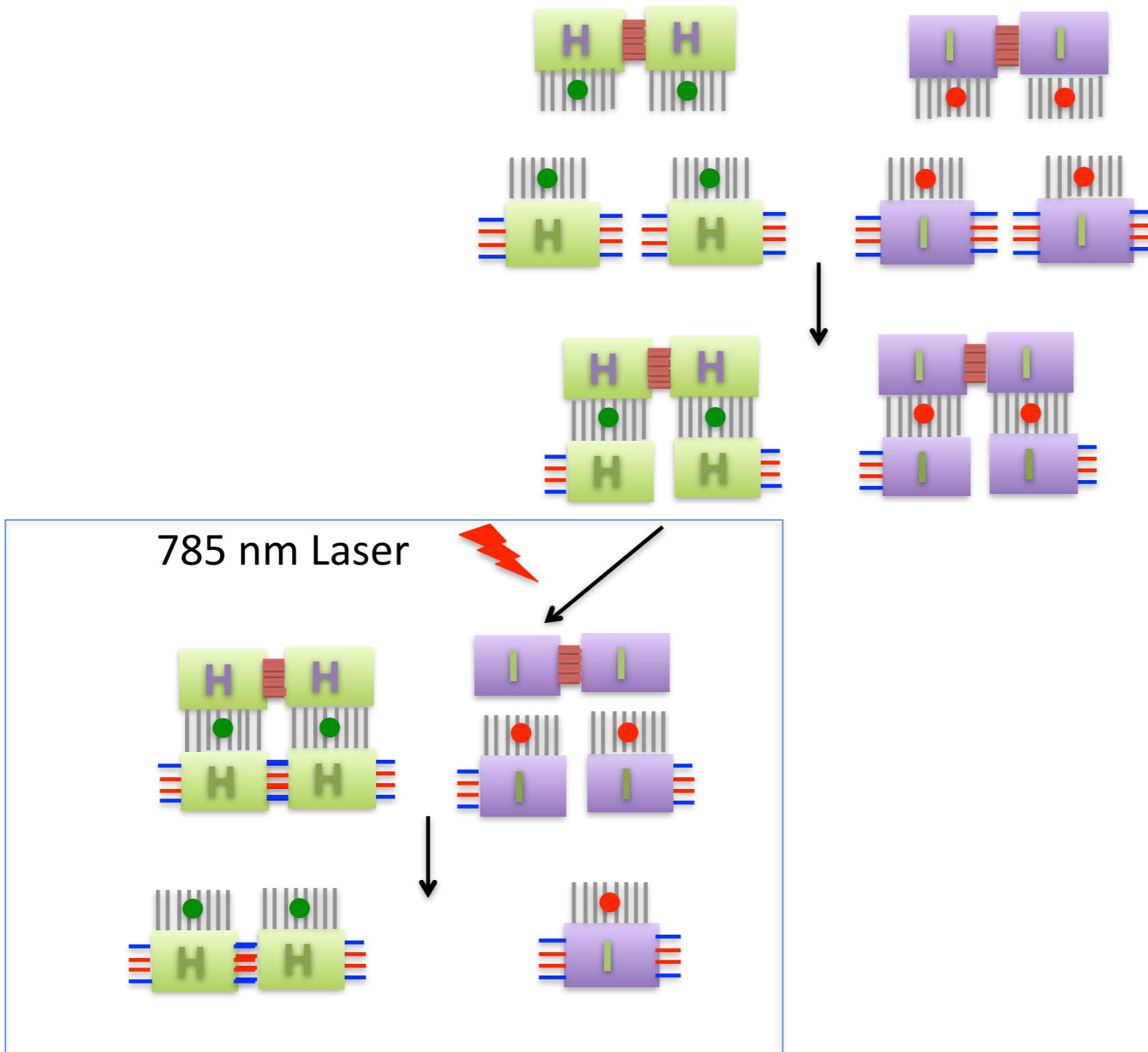
Local heating

$$j_Q \sim \frac{\dot{Q}}{4\pi r^2}$$

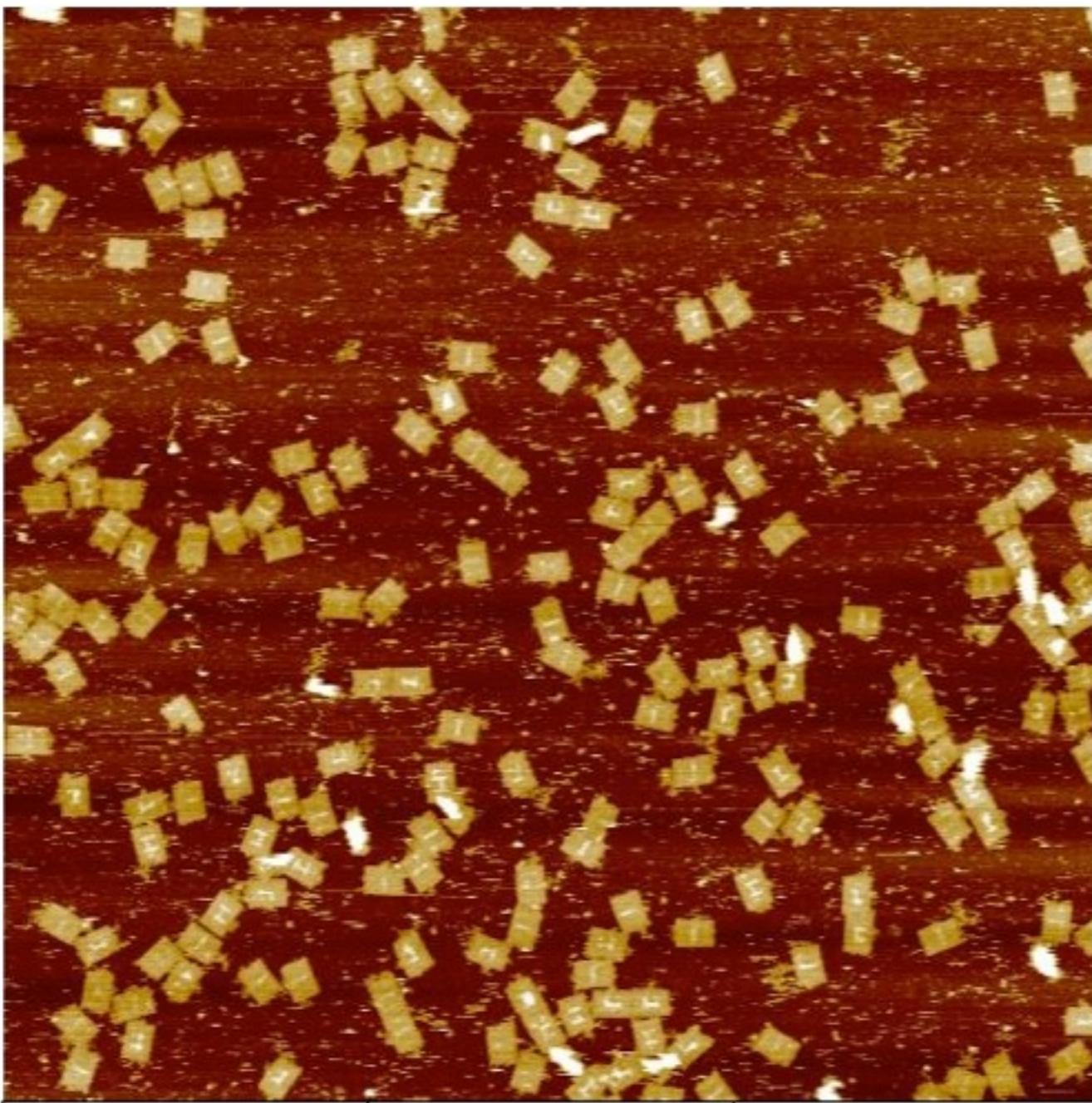
$$T = T_0 + \delta T / r$$



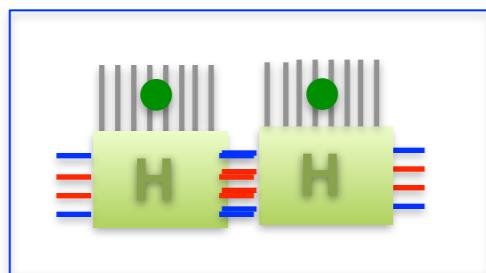
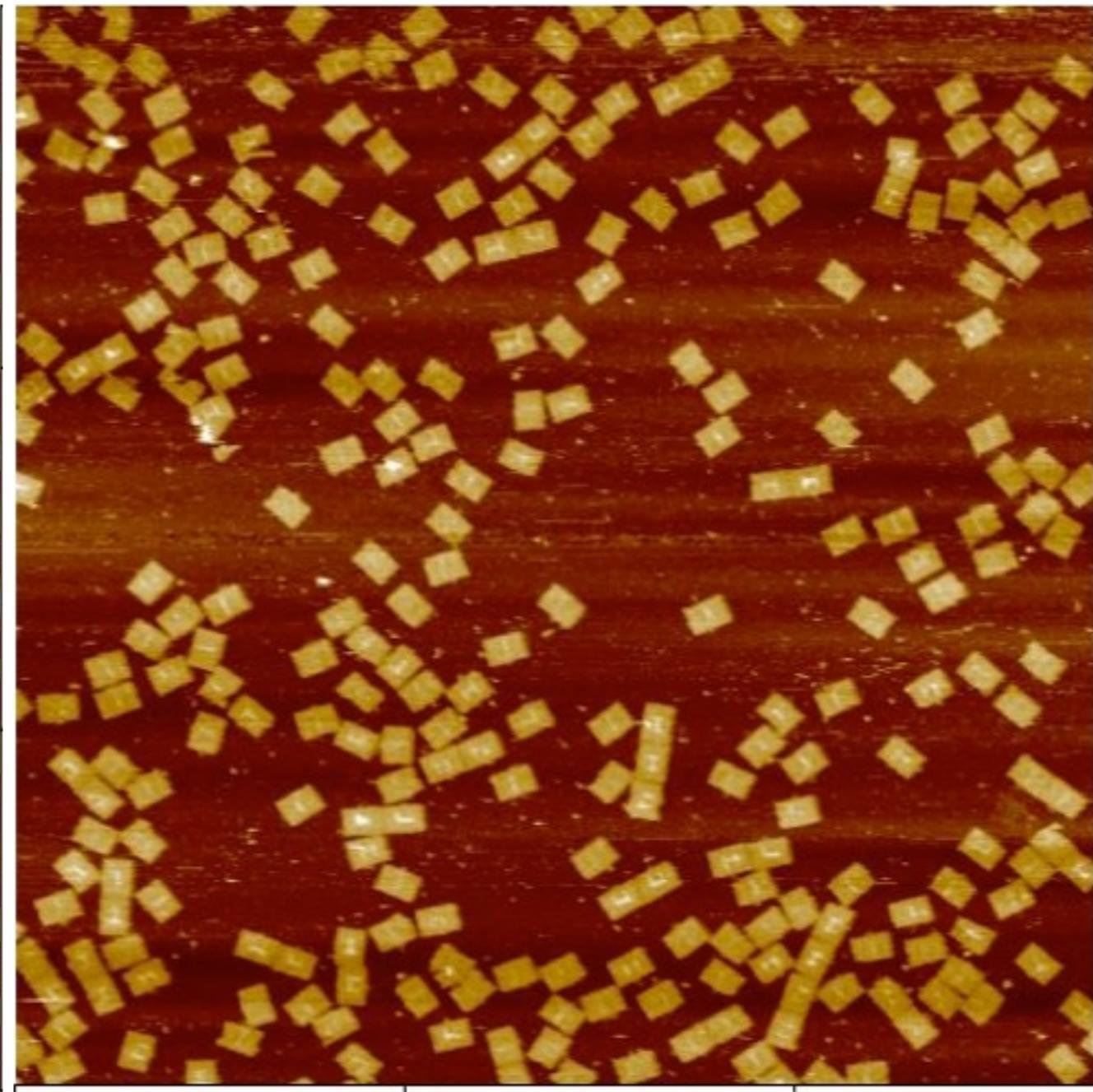
Seed HH	FG H-2 IR700	SG H-1 IR700	Seed II	FG I-4 IR800	SG I-3 IR800	
C 0	1	8	6	1	8	6



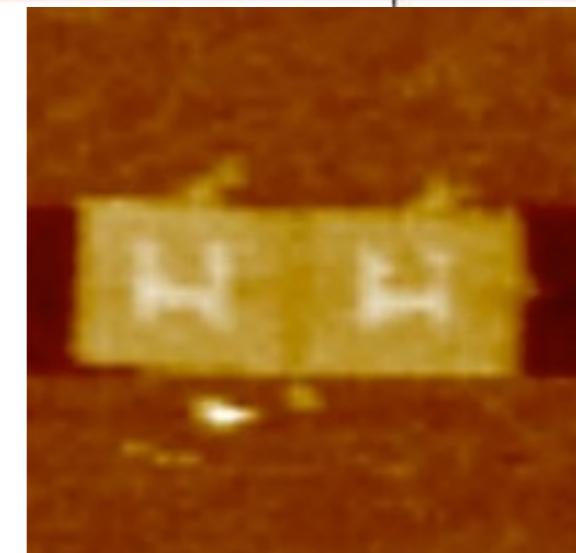
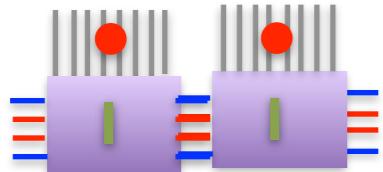
## Cycle 2

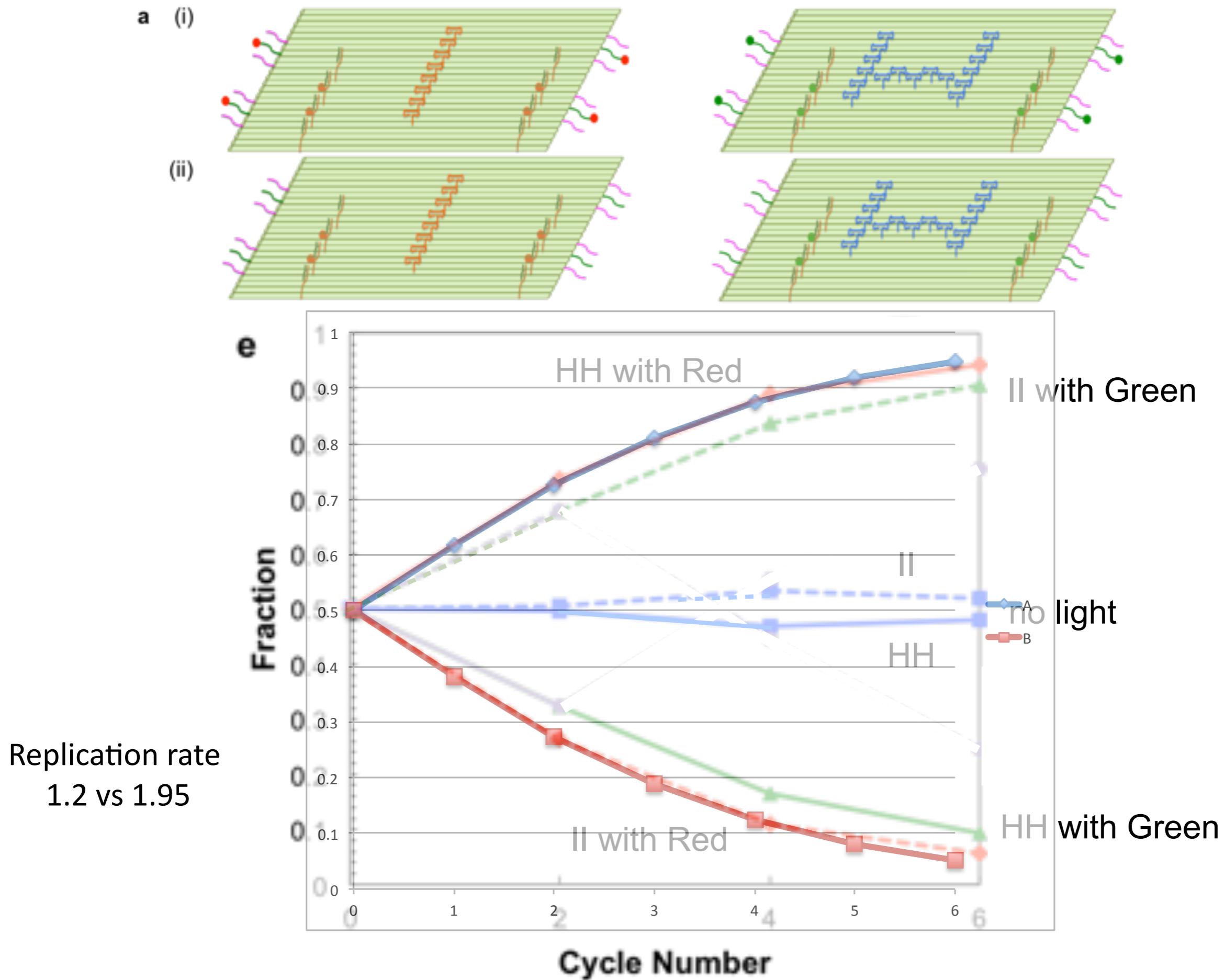


## Cycle 4



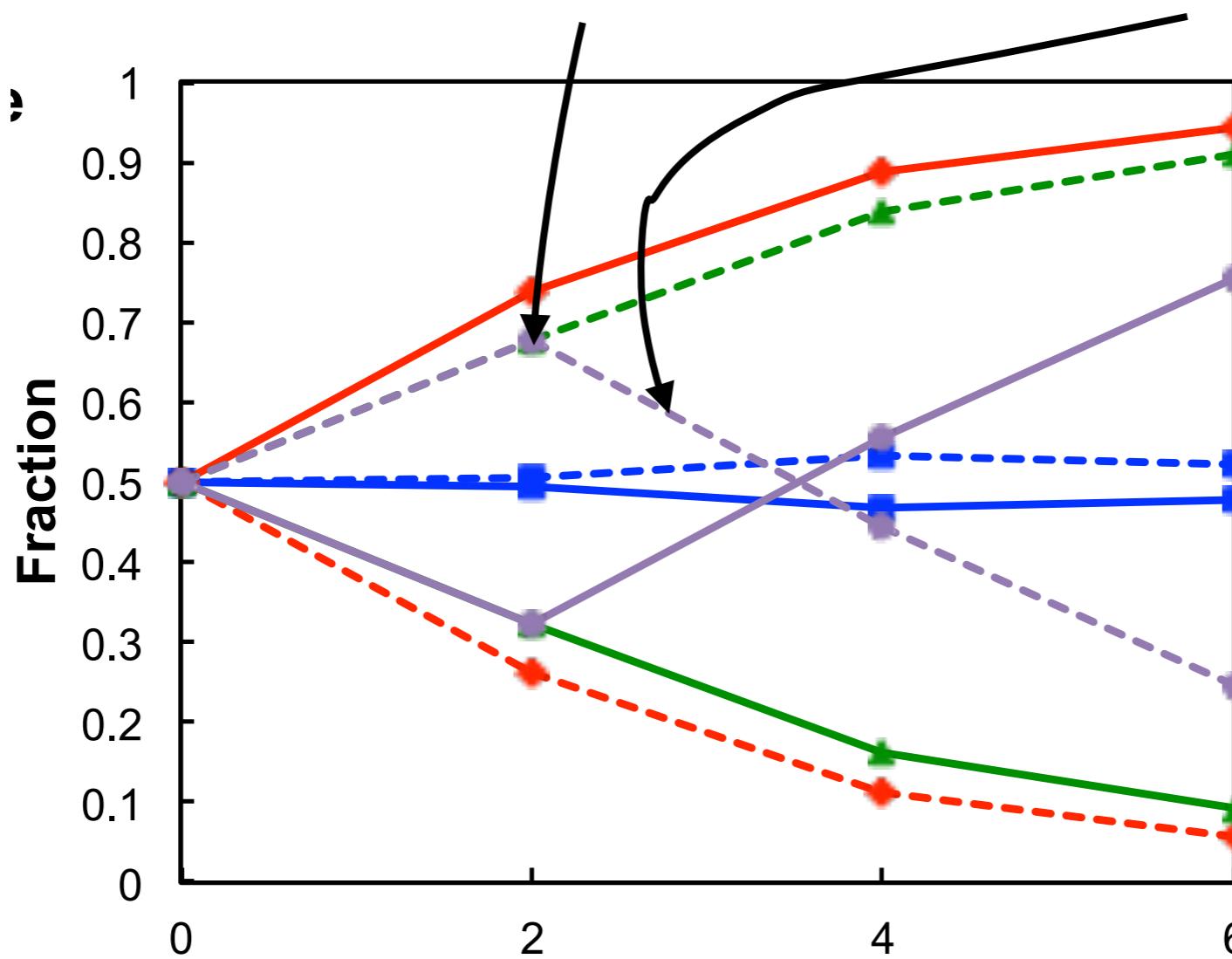
- ◆ ~ 4 C: Laser 785 nm for 20 min
- ◆ ~ 4 C: Laser 785 nm + UV for 1 h
- ◆ After each 2 cycles
- 1) add monomer H to keep  
HH: H-FG: H-SG = 1: 7: 7
- 2) Add Monomer I to keep  
 $(2^*HH + H): (2^*II + I) = 1: 1$





# Can reverse selection by switching lights

Grow in green 1 step then switch to red



# How about growing it outside?

# Roof Top - Washington Sq Park

Sun from here

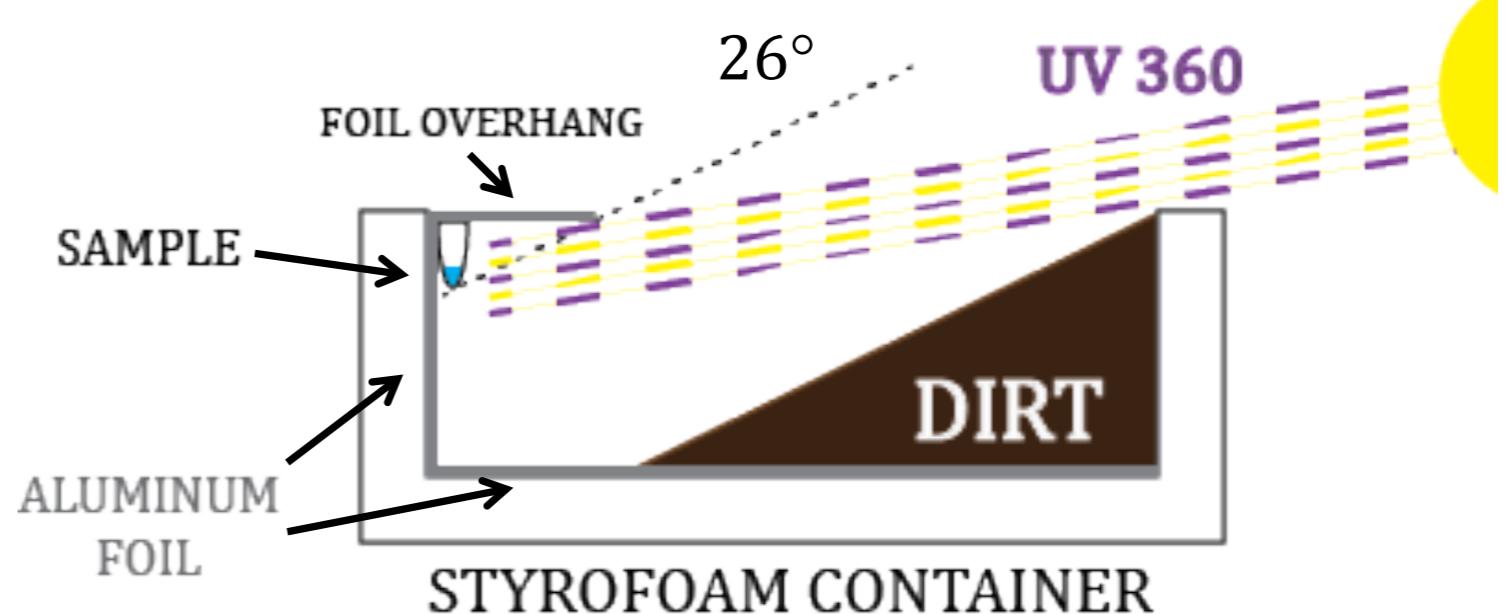


Dirt from here

# RoofTop DNA Origami Solar Replicator



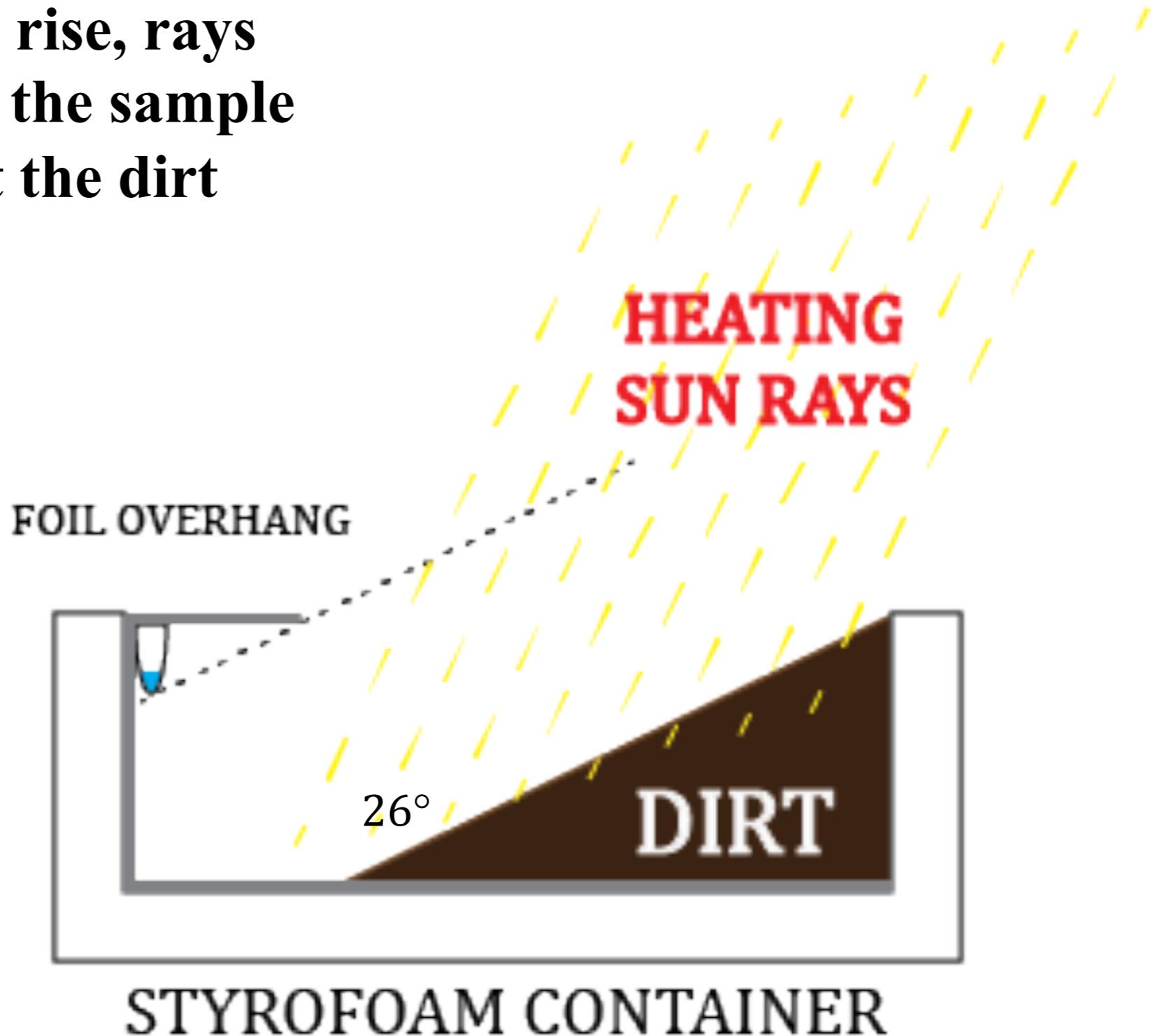
After a cold night, rays from the sunrise hit the sample for about 2 hours



As the sun continues to rise, rays are blocked from hitting the sample and now serve to heat the dirt

Why Dirt?  
Ambient temperature  
 $32\text{-}36\text{F} \rightarrow 0\text{-}2\text{C}$

Dirt Temperature  
 $32\text{-}96\text{F} \rightarrow 0\text{-}35\text{C}$



**Original Dimer:Mono Ratio – 1:30**

**After One Sunny Day:**

**Control Sample on roof (in Al Foil)**

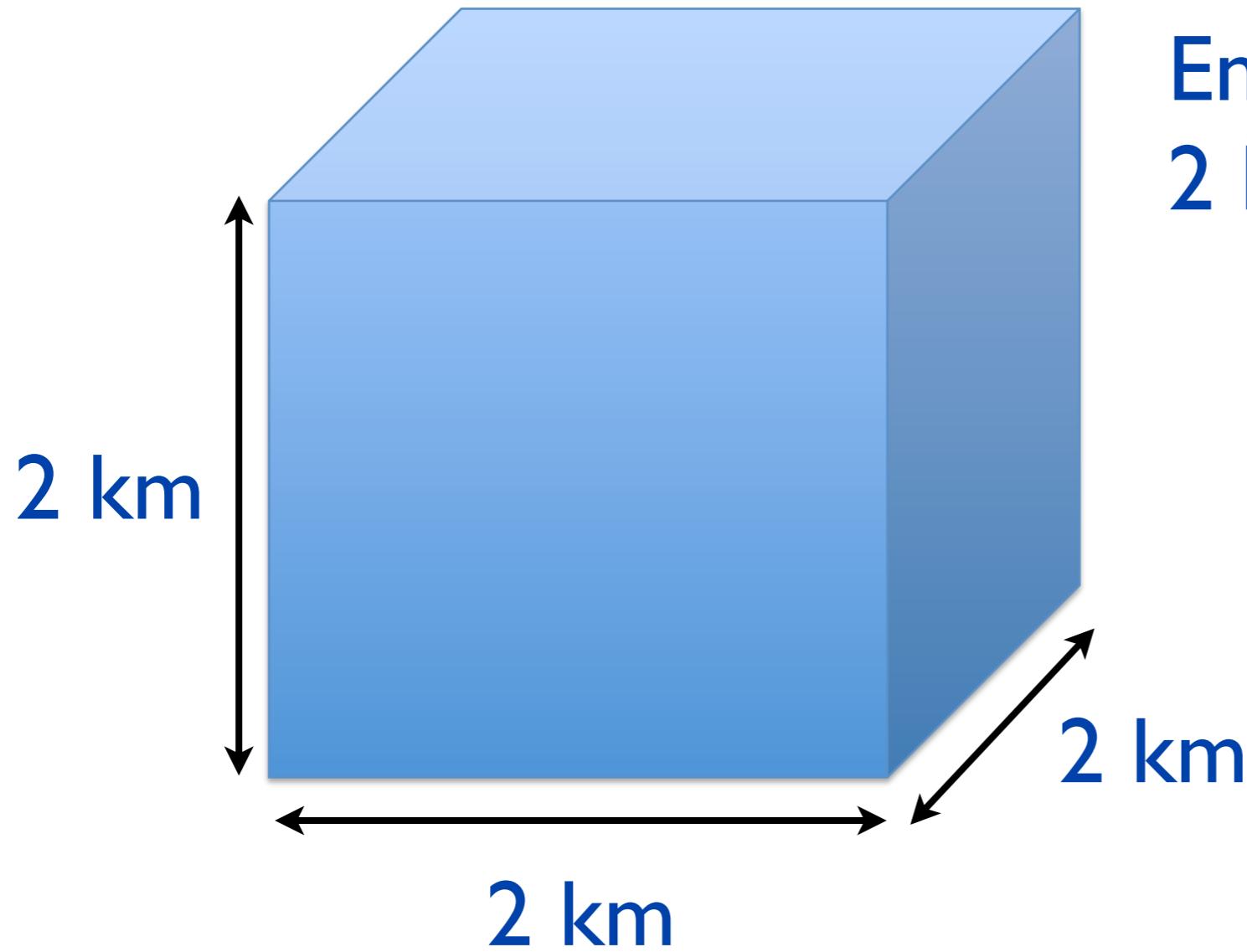
**Dimers Remain Same Concentration**

**Sunny Sample**

**Dimers Doubled**

# DNA as a functional material

Now much DNA is there?

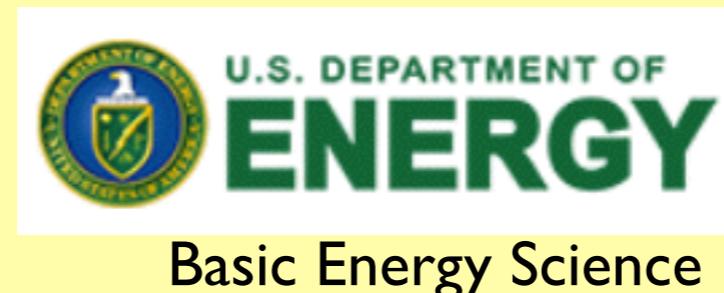


Enough to fill a cube  
2 kilometers on a side

Enough to build 200 cities the size of New York

# Summary

- Dynamic Clustering when flux in ( $\rho_+$ ) > flux out
- DNA is a great structural material
  - specific, controllable, reversible, or permanent bonds
- 1<sup>st</sup>? Artificial Self-replicating system with:
  - design flexibility
  - autonomous offspring
  - no enzymes
  - exponential growth (great way to make zillions of nanodevices)
  - uses only temperature and light mimicking daily cycles
  - replicates information and structure
  - 1: 7,500,000 and growing
- Next:
  - evolution
  - without nucleic acids



Keck Foundation  
NYU NSF MRSEC  
GORDON AND BETTY  
**MOORE**  
FOUNDATION

