

- **Announcements**
- **Pre-lab Lecture**
 - ❖ **Mod3 Concepts**
 - ❖ **Intro to M13 Virus**
 - ❖ **Intro to Solar Cells Materials**
 - ❖ **Today in Lab (M3D1)**

Announcements

- Introducing... Tahoura, TA for Module 3
... and bonus TA, Jackie!
- Quiz next time
- Mod 2 research paper
 - due 11/12 at noon
 - *revision* due 11/26 at 5 pm
 - a few general comments on framing
 - OH: NLL Mon 10-12, SKH Mon 3-5, ANS Tue 3:30-5
 - And Natalie OH by appt (nkuldell)

Ⓟ pic day

Module 3 Foundations

- Biology can interface with nano- and micro-scale materials

cells 1-10⁴ μm
★ viruses 0.01-1 μm
proteins / complexes 1-100 nm

- Nanoscale materials may have improved or even emergent properties

-elec / mag
-optical
-catalytic
...

★ benefits
+ risks ★

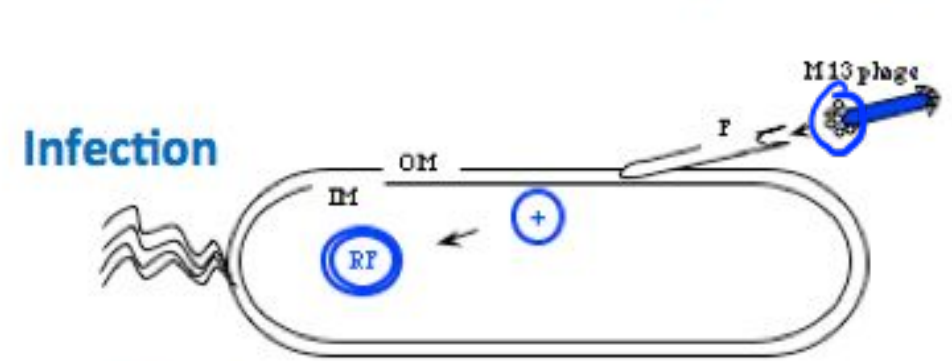
tough to predict
e.g. upon assembly

- Our nanomaterial is a phage!

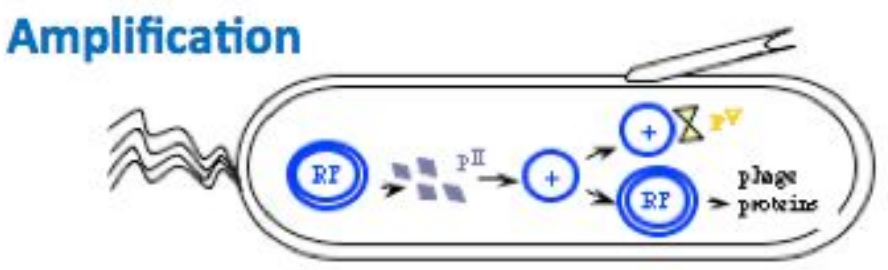
OOM



M13 phage life cycle (life cycle)

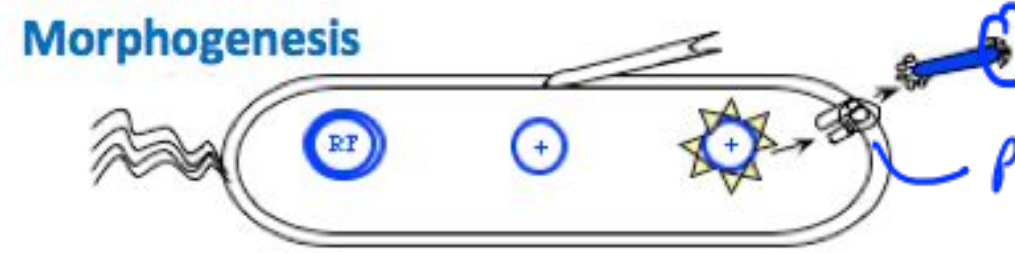


p3 p6 entry at TolA of the F pilus of E coli



p2, 5, 10 replication in DS form DS = double-stranded

packaged in SS form → coated by p8



p7/p9 exit p4, 11, 1 - make pore

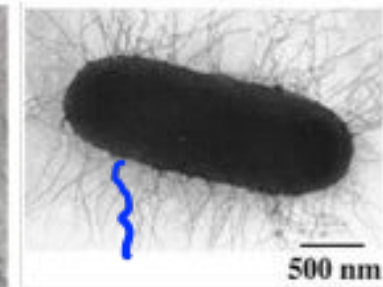
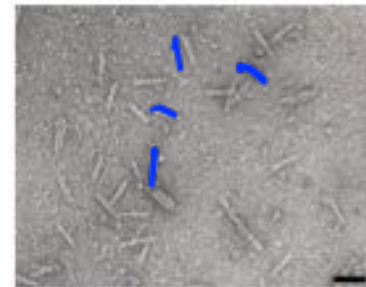
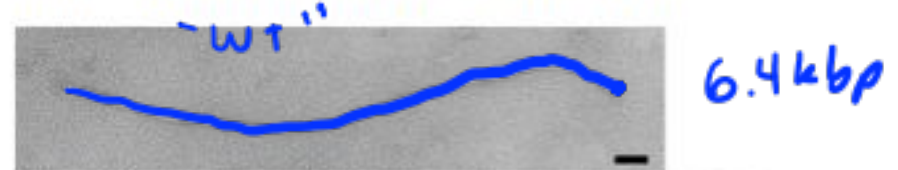
initial ϕ w/in 10^1 after infection

Image from Fall 2007 wiki. RF = replicating form

M13 as engineering substrate

- Length of DNA (to be packaged) dictates phage size... w/in limits
- Surface proteins for functional peptide display
- Method: (1) design library (2) binding assay screen

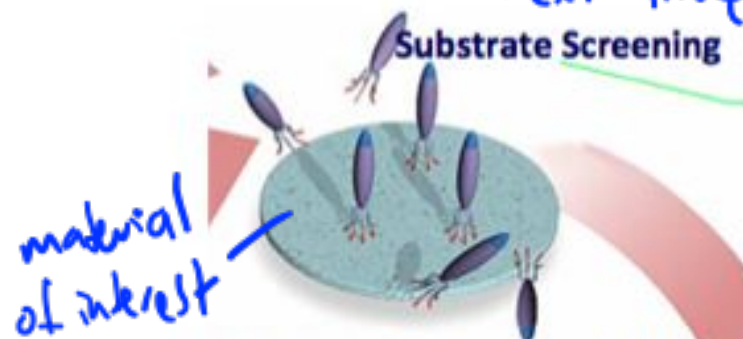
Micrographs from 20.109 wiki



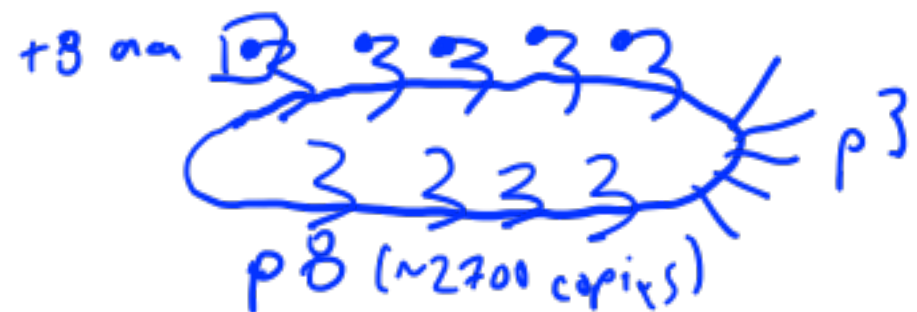
200 bp

113 kbp (cant exit)

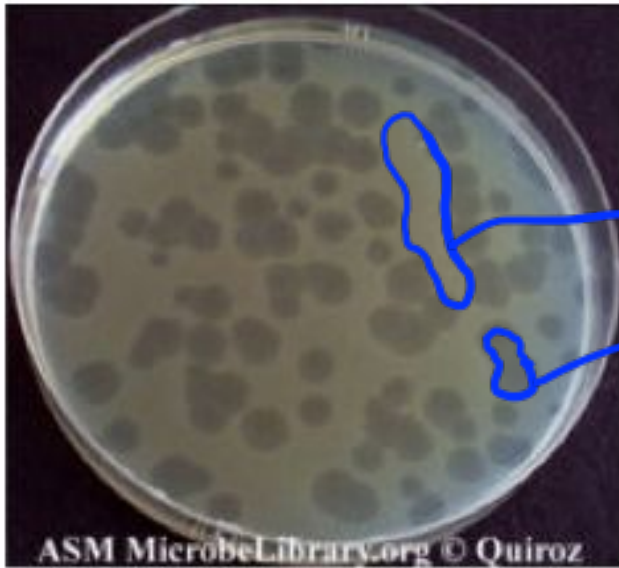
move next time!



Schematic from A. Belcher



Phage titer: plaque assay or spec.



By plating:

Phage slow *E. coli* growth upon infection

"lawn" - opaque = bacteria (saturated w/)

"plaque" clear = less dense bacteria,
∴ infected by ϕ

PFU (cf CFU)

By spectroscopy:

etc.

biol. content

A₂₆₀ = D/RNA peak

A₂₈₀ = protein peak

6×10^{16} (A₂₆₉ - A₃₂₀) → background

phage particles =

DNA bases in phage genome

for given Abs, genome size ↑ means # particles ↓ (cf blb:ins)

(SWNT-)Au/TiO₂ nanocrystal approach

- Begin today: react phage w/gold
 - Au ↑ usable light collection → your exp't
 - SWNT ↑ e- collection efficiency → TA exp't
- Why bother with phage?
 - surfactant for SWNT
 - bring TiO₂ proximate to SWNT or Au
- *Vary size of Au nanoparticles*
- Next time react w/Ti(OCH(CH₃)₂)₄
- Eventually...
 - TEM observation
 - solar cell assembly

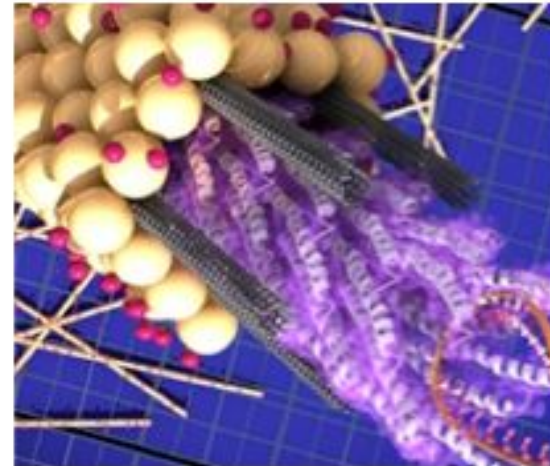


Image: Matt Klug

Today in Lab (M3D1): Workflow

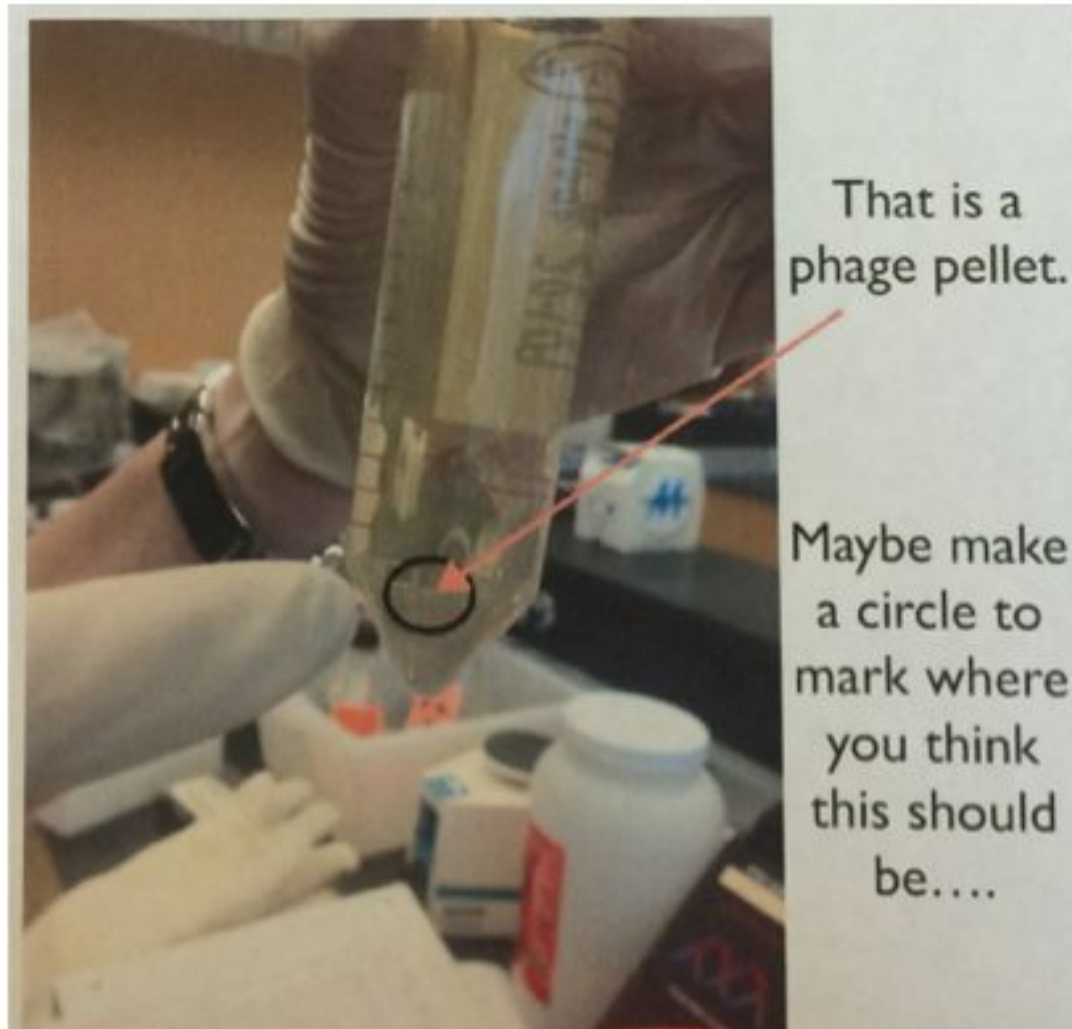
- Prepare phage by precipitation with PEG/NaCl
 - Incubations/spins *alone* are almost 2 h
 - At the end, phage are in the supernatant!! (in final steps, they are pellet)
 - Pellet is *bacteria*
- Obtain viral titer
 - take care with quartz cuvettes!
- React phage w/gold
- Downtime: calculation sheet, ~~reflections~~, FNT, etc.
in advance (2)

Today in Lab (M3D1): Samples and Steps

Group (T/R)	AuNP Size (nm)	Group (W/F)	AuNP Size (nm)
Red Cherries	5	Green	20
Tiger	50	Blue	5
Yellow	20	Pinkle	50
Green	20		
Blue	5		
Pink	20		
Purple	50		

- Measure # phage/mL
- Calculate volume Au needed
 - stocks given in g/mL
- Goal: 1.45×10^{-17} g gold/phage
- Mix in glass scintillation vial
- Store in fridge

Today in Lab (M3D1): Key Tip from Shannon



★ orient to know pellet location ★