

# Lab Log

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**18/10/2017**

- Autoclaved the soil again
- Autoclaved 1 L of DI millicule water
- Autoclaved 1 L of 0.6% soft agar
- Autoclaved a spoon to weigh out soil
- Got some *SBW25φ2* from Floh. Around 1 mL at  $7.26 \times 10^8$ .
  - Grow up overnight
  - Transfer 60  $\mu$ L of bacteria and 10  $\mu$ L of phage (around  $7 \times 10^6$  pfus) into 6 mL of KB agar
  - Should give a concentration of around  $10^8$  phage/mL
  - Done this in triplicate
- Grow up *lacZ* and *WT* strains overnight. Should give concentration of around  $\sim 10^8$  cells in 60  $\mu$ L.
- Do these in triplicate
- Added 60  $\mu$ L of frozen overnight culture from first experiment (18/08/2017 *lacZ* and *WT*)

**Retrospectively work out density of the overnight stocks and phage**

**19/10/2017**

- Put 80g of soil into each 10cm x 10cm microcosm
  - Used autoclaved spoon
  - Placed scale in laminar flow hood (cleaned with ethanol before and after)
- Placed 5 mL ( $\sim 200$   $\mu$ L per microcosm) of *lacZ* and *WT* into separate 12 mL centrifuge tubes
  - Centrifuged for 15 minutes at max speed ( $\sim 4500$  r.p.m) on big centrifuge
  - Want to get to 5 mL per microcosm for inoculating ( $\sim 125$  mL in total)
  - Resuspended pellet into 2250  $\mu$ L, vortexed and placed 620  $\mu$ L, 620  $\mu$ L and 810  $\mu$ L into three different falcon tubes
  - Filled these three falcon tubes up to 40 mL, 40 mL and 45 mL respectively
    - \* This guaranteed the same concentration of sample in each falcon tube
  - Placed 5 mL of *lacZ* or *WT* strain into each microcosm
  - Froze ( $-80$   $^{\circ}$ C) 900  $\mu$ L of inoculate in 900  $\mu$ L of glycerol (25% final concentration)
- In the no phage treatments, we added 5 mL of M9
- Added 5 mL of phage to phage treatments
  - Place 900  $\mu$ L of bacteria + phage into three centrifuge tubes
  - Add 100  $\mu$ L (10%) chloroform into each tube (under fume hood)
  - Vortex rigorously
  - Centrifuge for 2-3 minutes at full speed (minifuge)
  - Take out supernatant and placed in a single tube (took out 800  $\mu$ L of each tube)
  - Put 40 mL of M9 into 6 tubes
  - Added 400  $\mu$ L into each tube ( $\sim 100$  fold dilution from the initial stock)
  - Shake each tube and add 5 mL into each microcosm
- Place microcosms into the 26  $^{\circ}$ C incubator (Level 1 incubator room)

**22/10/2017**

- setup 2 *WT* microcosms up for spot assays of phage
- used a crystal from  $T_0$
- autoclaved 2 L of KB agar. Put in small autoclave