#### Lab book

#### Saul Pierotti

July 9, 2019

#### 02/07/2019

- 12 am: I prepared 1L of M9 medium whith Anuradha, who prepared another 1L
- 3 pm: streaked P. fluorescens SBW25 with Anuradha from the stock at -80°C in position IIIA3-29
  - I streked 1 LB plate and 1 M9 plate
  - Anuradha streaked 1 LB plate and 1 M9 plate
  - Plates put at 28  $^{\circ}\mathrm{C}$

#### 03/07/2019

- 9.30 am: streaked *E. coli* K12 MG1655 from the stock at -80°C in position IIIA1-22 and \* E. coli\* B REL606 form the stock in position IIIA1-14 with Gunda
  - I streked 2 LB plates and 2 M9 plates (1 for each strain)
  - Gunda streaked 2 LB plates and 2 M9 plates (1 for each strain)
  - Plates put at 28 °C
- I tried to write the primers for discrimitating the strains by myself

## 04/07/2019

- 9.00 am: Put all the cultures in 4 mL of LB/M9 at the appropriate temperature (28/37°C, 200rpm)
  - Labeled all the dilution plates for tomorrow, put glass beads inside
  - Put glycerol in vials for tomorrow, vials labeled
  - Put plates with streaks of the strains at 4°C
- Checked primers with Gocke: need to change something because I did not check for partial matches
- Afternoon: I have rewritten the primers and sent them to Gocke

# 05/07/2019

- Medium controls for the O/N cultures look all clean, cultures are all well grown except for SBW25-1 M9 (nothing visible) and SBW25-2 M9 (really faint)
- 9.00 am: inoculated all the glycerol stocks except for SBW25-1 M9, which did not show any growth
  - Stocks put at -80°C in the bottom shelf
  - Overnight cultures put at 4°C
  - SBW25-1 M9 put back at  $28^{\circ}\mathrm{C}$
- 10.15 am: meeting with Arne
  - We decided to talk to other theory people to discuss the modelling part
- 11.30 am: inoculation of the long term cultures (4 mL of LB or M9) with 4  $\mu$ L of the overnight cultures
  - -05/07/19 is my day 0 (D0) of the long term experiment)
  - Long term cultures are labeled as strain(SBW25/MG1655/REL606)-sample(1-3, 1-6 for REL606) medium(LB/M9)

- \* LB is labeled in black, M9 in blue
- Long term cultures put at 28/37°C as appropriate
- Overnight cultures put back at 4°C
- SBW25-1 M9 seems to show an incredibly faint cloudiness, I decided to use it to inoculate the respective long term culture
  - \* I will wait this afternoon for making the glycerol stock
- Long term cultures put at the 28/37°C with MCs, SBW25-1 M9 put at 28°C
- 5 pm: SBW25-1 M9 shows visible growth
  - Glycerol stock inoculated and put at -80°C
  - Stocks are labeled with the date (05/07/19), medium, strain, sample number, day of the experiment (D0), my initials (SP)
  - LB vials labeled in black and M9 vials in blue
- Inoculated all the dilution plates (52 plates) with 100  $\mu$ L taken from the respective dilution of the O/N cultures in Ringer's (D0 dilution plates)
  - There are 2 plates for each sample, at different dilutions (10^-7 and 10^-6 for LB, 10^-6 and 10^-5 for M9)
  - 3 SBW25 LB samples (1-3), 3 MG1655 LB samples (1-3), 6 REL606 LB samples (1-6)
  - 3 SBW25 M9 samples (1-3), 3 MG1655 M9 samples (1-3), 6 REL606 M9 samples (1-6)
  - Medium controls for LB and M9, at 28°C and 37°C (4 in total)

## 06/07/2019

• Counted the dilution plates for E. coli, and put at 4°C

## 07/07/2019

- Counted the dilution plates for P. fluorescens, and put at 4°C
- Plate SBW25-2 LB D0 10<sup>-</sup>7 contaminated with green mould
  - Put parafilm around it to avoid spreading of spores
- Created spreadsheets for glycerol stock and dilution plates
- SBW25 did not produce many colonies in LB (10<sup>-7</sup> and 10<sup>-6</sup> dilutions)
  - Next time I will try with 10<sup>-6</sup> and 10<sup>-5</sup>
- CFU/mL in the D0 O/N cultures used for inoculating the long term cultures

Sample	$\mathrm{CFU/mL}$
SBW25-1 LB	$1,00*10^{8}$
SBW25-2 LB	$7,00*10^7$
SBW25-3 LB	$5,00*10^7$
SBW25-1 M9	$7,60*10^7$
SBW25-2 M9	$8,70*10^7$
SBW25-3 M9	$1,41*10^8$
MG1655-1 LB	$7,60*10^8$
MG1655-2 LB	$7,80*10^8$
MG1655-3 LB	$6,10*10^8$
MG1655-1 M9	$1,04*10^9$
MG1655-2 M9	$9,00*10^{8}$
MG1655-3 M9	$7,00*10^8$
REL606-1 LB	$3,40*10^{8}$
REL606-2 LB	$2,80*10^8$
REL606-3 LB	$3,40*10^{8}$
REL606-4 LB	$3,70*10^{8}$
REL606-5 LB	$3,50*10^{8}$

Sample	$\mathrm{CFU/mL}$
REL606-6 LB	$1,12*10^9$
REL606-1 M9	$1,45*10^{8}$
REL606-2 M9	$4,70*10^{8}$
REL606-3 M9	$7,40*10^{8}$
REL606-4 M9	$5,80*10^8$
REL606-5 M9	$5,90*10^{8}$
REL606-6 M9	$1,14*10^9$

## 08/07/2019

- Dilution plates of D0 checked with Jenna and then trashed
- I have labeled the plates, cryovials and tubes for D5, which will be on Wednesday 10/07
- Primers for telling apart SBW25, MG1655 and REL606 ordered with Anuradha
  - $\ SBW25\_unique1\_fw: \ 5'-ATACTACGACTCCAGAGCGATGG-3'$
  - $\ SBW25\_unique1\_rv: \ 5'-GTTCAGCGTCTGCGTGGCTTG-3'$
  - SBW25 expected product size: 1024 bp
  - REL606\_unique1\_fw: 5'-CAGTGGATTGTGGTTTGTTGCC-3'
  - $\ REL606\_unique1\_rv: \ 5'-GGCTGGTACTTTTCAGGTCGG-3'$
  - REL606 expected product size: 1138 bp
  - $\ MG1655\_unique1\_fw: \ 5'-CTGAATCGGTCATGATGATGGGGACTG-3'$
  - MG1655\_unique1\_rv: 5'-TTCAGGCGGACTTACTATCCCG-3'
  - MG1655 expected product size: 1241 bp