

iGEM 2016: Pittsburgh
Week 4 Lab Notebook

Monday, June 13

Lab meeting

Kanamycin and ampicillin plates

250 mL each

1:1000 dilution of stock solution

Transformation (Claire and Maddie)

lacZ alpha fragment (part BBa_I732006) Plate 3, 3B

eGFP (part BBa_E0040) Plate 4, 13L

50 uL competent cells + 5 uL resuspended DNA

T7 -- GFP (from last year's team) + 150 uL competent cells

Incubation for 1 hour

lacZ and eGFP initial incubation on ice for closer to 1 hour waiting for T7 -- GFP

Liquid cultures overnight of T7 - RBS and terminator sequences from last week

Mini-prep liquid cultures of T7 - RBS -- amilCP (Maya)

DNA	Concentration (ng/ul)
5:1-1	94.2
5:1-2	114.2
5:1-3	72.3
5:1-4	67.0
5:1-5	99.7
3:1-1	130.1
3:1-2	130.4
3:1-3	93.3
3:1-4	93.4
3:1-5	122.0

Digest and gel electrophoresis check of ligation with SpeI and XbaI of 5:1 and 3:1 samples

Used 100ng DNA

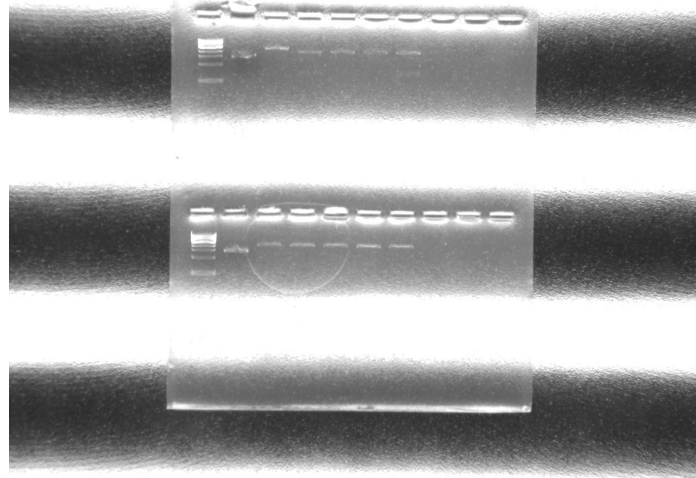
-3-2, 3-5 looked good; 3-3, 3-4 were weird; rest were incorrect

Control is T7-RBS uncut

Ladder is 1kb

Top lane is 3:1 (Ladder, control, 1, 2, 3, 4, 5)

Bottom lane is 5:1 (Ladder, control, 1, 2, 3, 4, 5)



Tuesday, June 14

T7-GFP: many colonies

eGFP: a few colonies

lacZ: one colony in afternoon

Resuspended DNA placed in vial

Concentration = 15.9 ng/ μ L

Liquid cultures of T7-GFP (x2), eGFP (x2), and lacZ (x1) (Claire)

5 mL LB + 5 μ L ampicillin

Miniprep liquid cultures of T7 RBS and terminator sequences

Concentrations: T7-RBS 1: 127.6 ng/uL

T7-RBS 2: 92.1 ng/uL

Terminator 1: 56.5 ng/uL

Terminator 2: 91.4 ng/uL

Restriction digests

T7-RBS: 1 uL SpeI

1 uL PstI

1 ug T7-RBS 1 = 7.84 uL DNA

2 uL 10x buffer

7.16 uL H₂O

Terminator: 1 uL EcoRI

1 uL XbaI

1 ug Terminator 2 = 10.94 uL DNA

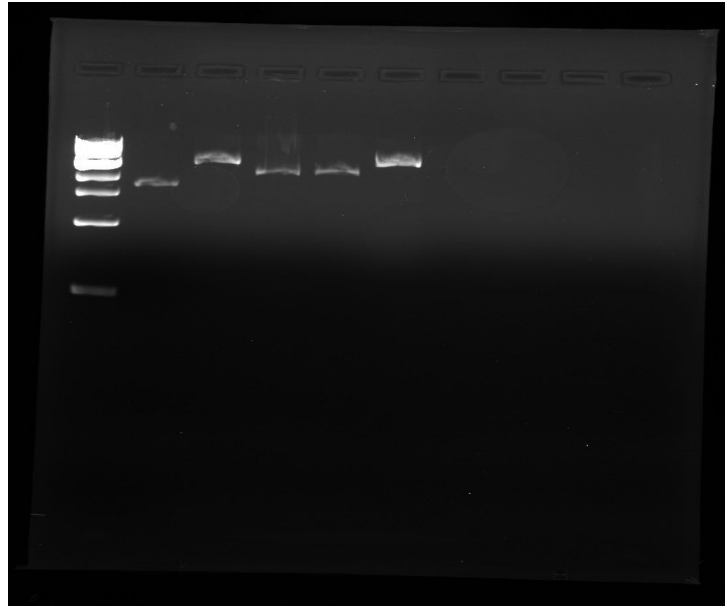
2 uL 10x buffer

5.06 uL H₂O

Request full lacZ (BBa_K1444017) from iGEM, 2014 Submission Plates

*sequence not confirmed

Linearized 3:1 (2,3,4,5) with EcoRI using 1ul DNA (6/14 gel electrophoresis image)



1kb Ladder

T7-RBS uncut control

Ladder, control, 2, 3, 4, 5

2 and 5 migrated the least, these probably have T7-RBS-amiCP

3 and 4 migrated in between control and 2/5: Cut, but did not have insert

Contact TECBio and DiSCoBi0

Wednesday, June 15

Glycerol stocks: T7-GFP (MP), eGFP (MP), lacZ (Claire)

Mini-prep all liquid cultures (Claire)

Concentrations

T7-GFP MP: 120.8 ng/ul

T7-GFP CC: 110.9 ng/ul

eGFP MP: 160.6

eGFP CC: 188.5

lacZ: 103.8

restriction digests (Praneeth)

lacZ: 1 ul Xbal

1 uL PstI

1 ug DNA = 9.63 ul lacZ

2 uL 10x buffer

6.37 uL H₂O

eGFP: 1 ul Xbal

1 uL PstI

1 ug DNA = 9.63 ul eGFP CC

2 uL 10x buffer

10.69 uL H₂O
eGFP: 1 ul XbaI
1 uL PstI
1 ug DNA = 6.23 ul eGFP MP
2 uL 10x buffer
9.77 uL H₂O

Incubate at 37 degC 30 min.

Incubate at 80 degC 20 min.

Gel electrophoresis

Lane	1	2 ul ladder, 2 ul loading dye
	2	uncut T7, 2 ul dye
	3	1 ul double digest T7 (SpeI/PstI)
	4	uncut terminator, 2 ul dye
	5	1 ul double digest terminator (EcoRI/XbaI)
	6	1 ul eGFP CC uncut, 2 ul loading dye
	7	all double digest eGFP CC (XbaI/PstI), 2 ul dye
	8	all double digest eGFP MP (XbaI/PstI), 2 ul dye
	9	1 ul lacZ uncut, 2 ul dye
	10	all double digest lacZ (XbaI/PstI), 2 ul dye

Purify T7 and terminator (Maya)

Concentrations:	T7:	69.9 ng/uL
	Terminator:	66.5 ng/uL

Gel extraction of eGFP (2 lanes) and lacZ (Praneeth)

Concentrations:	eGFP CC:	11.9 ng/ul
	eGFP MP:	15.8 ng/ul
	lacZ:	10 ng/ul

Ligation: T7 – reporter (eGFP or lacZ) (Claire)

Control:	1 ul T7
	0 ul insert
	2 ul buffer
	16 ul H ₂ O
	1 ul T4 DNA ligase
3:1 eGFP:	1 ul T7
	50 ng eGFP = 3.16 uL eGFP MP
	2 ul buffer
	12.84 ul H ₂ O
	1 ul T4 DNA ligase
7:1 eGFP:	1 ul T7
	115 ng eGFP = 9.66 uL eGFP CC
	2 ul buffer
	10.34 ul H ₂ O
	1 ul T4 DNA ligase
3:1 lacZ:	1 ul T7

50 ng eGFP = 5 uL lacZ

2 ul buffer

11 ul H₂O

1 ul T4 DNA ligase

Incubate at room temperature for 10 min

Incubate at 65 degC for 10 min

Transformation of ligation reactions (Aife)

50 uL Cheryl's competent cells

5 uL reaction

Ice 20 min

42 degC 2 min

Ice 5 min

Add 200 uL SOC medium

37 degC shaker for 1 hr 4 min

Plate 100 uL onto chloramphenicol plates

Sent out 3:1 (2), 3:1 (5), EGFP, and Terminator to GeneWiz for sequencing (Maya)

Used VF2 and VR both for 3:1 plasmids

For terminator and EGFP used VF2

Thursday, June 16

Liquid cultures of transformations (Aife)

3:1 lacZ (x2)

3:1 eGFP (x5)

7:1 eGFP (x1)

Friday, June 17

7:1 eGFP didn't actually contain a colony

Mini-prep yesterday's liquid cultures (Claire)

Concentrations: lacZ 3-1: 193.3 ng/uL

lacZ 3-2: 170.8 ng/uL

eGFP 3-1: 144.3 ng/uL

eGFP 3-2: 151.1 ng/uL

eGFP 3-3: 198.6 ng/uL

eGFP 3-4: 182.8 ng/uL

eGFP 3-5: 208.7 ng/uL

Restriction digest: check for insert (Praneeth)

For all mini-preps, 1 uL EcoRI

0.5 uL DNA

1 uL buffer

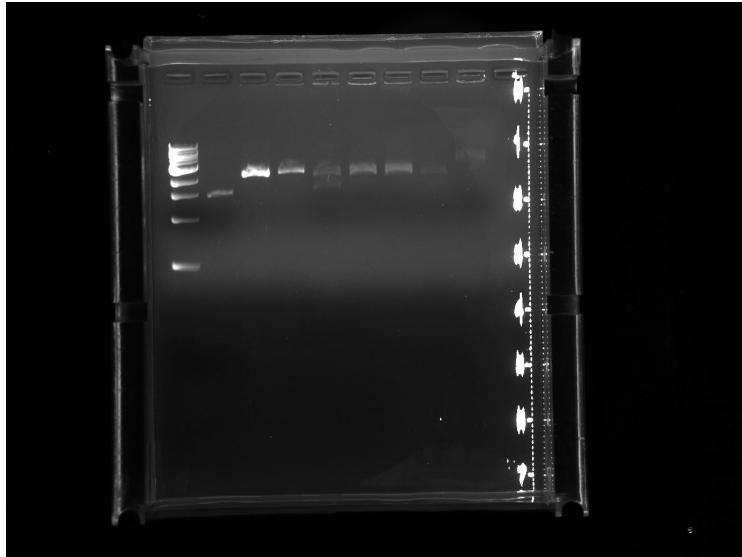
7.5 uL H₂O

Incubate at 37 degC for 30 min; 65 degC for 20 min

Gel electrophoresis

Lane 1 ladder

- 2 uncut T7-RBS
- 3 T7 - lacZ 3-1
- 4 T7 - lacZ 3-2
- 5 T7 - eGFP 3-1
- 6 T7 - eGFP 3-2
- 7 T7 - eGFP 3-3
- 8 T7 - eGFP 3-4
- 9 T7 - eGFP 3-5



lacZ 1 and 2; eGFP 2, 3, and 4 good