

Week of Oct 7

WEDNESDAY, 10/10/2018

Origin Tests: *S. mel*, *L. lac*, *C. glut*

1. Transformed 200 ng of RSF1010, pBBR1 and pWV01 plasmid into all 3 strains
 - a. transform by electroporation;
 - I. 2.5 kV for *S. mel* and *C. glut*
 - II. 2 kV for *L. lac*
 - b. recovered in plain LB for *S. mel*; universal medium + glucose for *C. glut*; universal medium + sucrose for *L. lac*; all in 30C shaking incubator
 - I. 3 hrs for *S. mel*
 - II. 2 hrs for *L. lac*
 - III. 1 hr for *C. glut*
2. Plated *L. lac* and *C. glut* on both plain and Kan50 LB plates; also plated *L. lac* onto m17 + Kan50 plate; *S. mel* plated on plain and Neo100 LB agar plate
3. All strains in 30C incubator;

THURSDAY, 10/11/2018

FRIDAY, 10/12/2018

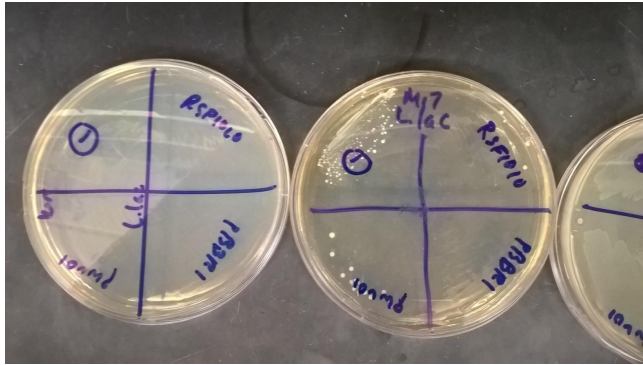
Results of Origin Tests

L. lac

1. Some growth on Kan+M17 Agar plate; growth on plain LB agar plate; no growth on Kan plate



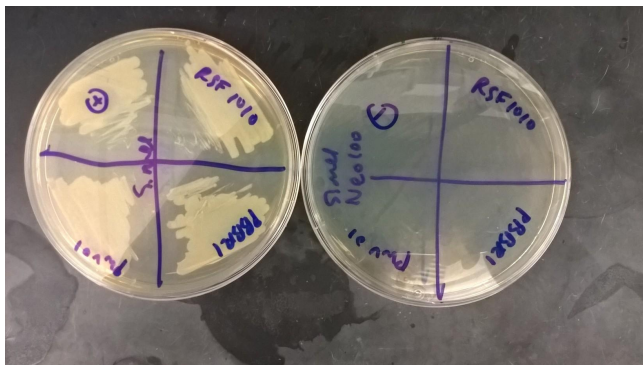
Origin testing L. lac Final 2.jpg



S. mel

1. Some growth plain LB plate; no growth on Neo plate

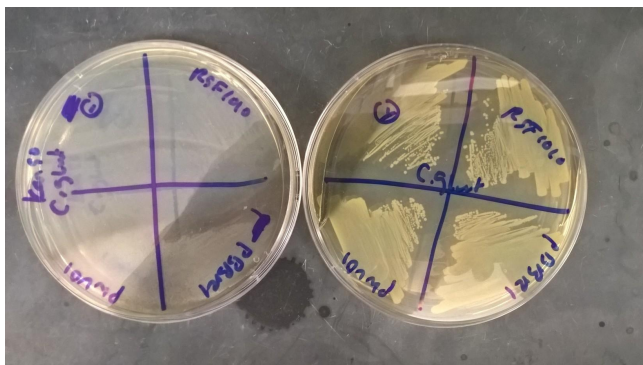
Origin testing S. mel final.jpg



C. glut

1. Growth on plain agar plate; dried out Kan plate

Origin testing C. glut Final.jpg



Origin Testing: Acinetobacter baylyi

1. Picked colony of A. b. from plate Anna made into 4 mL of LB
2. Let grow overnight in 30C shaking incubator

SATURDAY, 10/13/2018

Origin Testing: *Acinetobacter baylyi*

1. Put 1 mL of LB, 70 μ L of *A. b.*, and 100 ng of either RSF1010, pWV01, or pBBR1 into Falcon tube
2. Incubated tube for 3 hrs
3. Plated 25 μ L of each transformation + non-transformed *A. b.* onto Kan50 and plain LB plate; spread with sterile inoculation loop
4. Put in 30C incubator overnight

SUNDAY, 10/14/2018

Origin Testing Results: *Acinetobacter baylyi*

1. pWV01 on Kan50 plate grew; no other growth on Kan50 plate; all transformants grew on plain LB plate
2. Weird; was supposed to work with RSF1010; is our RSF1010/pBBR1 DNA bad?
 - a. test w/ *E. coli*?

