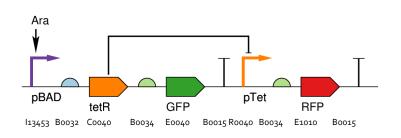
# BBa\_K783067

PART DESCRIPTION: Arabinose-TetR Inverter

This device is specified to act as a logical inverter as a function of [L-arabinose]. GFP expression should increase as L-arabinose concentration increases and RFP expression should decrease. Characterization finds behavior consistent with specification.



## **Contact Information**

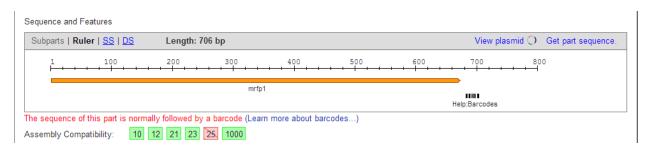
Author(s): Evan Appleton, Monique Freitas, Sonya Iverson

<u>Team</u>: Boston U iGEM 2013 <u>Contact</u>: eapple@bu.edu

<u>Data Collection</u>: Evan Appleton, Monique Freitas <u>Related Parts</u>: BBa\_l13453, BBa\_C0040, BBa\_R0040

Affiliation: Boston University (Densmore Lab) Date: 9/20/2012

# **Standard Design Information**



Chassis:E. coliStrain:Bioline™ α-GoldDevice Name:BBa\_K783067Assembly:BioBricks™

Device Type:InverterProtocol:BU BioBricks assembly protocolSafety Level:Risk Group 1Scars:Yes; 6bp scars between each part

Components: BBa\_13453-BBa\_Boo32-BBa\_Coo4o-BBa\_Boo34- Insertion: Plasmid

BBa\_E0040-BBa\_B0015-BBa\_R0040-BBa\_B0034- Vector: pSB1A3

BBa\_E1010-BBa\_B0015

Additional Comments: SI built device, EA and MF characterized device.

# Restriction Digest and Gel Electrophoresis

## **BASIC INFORMATION**

Purpose: Verify restriction mapping

Data Type: DNA size bands

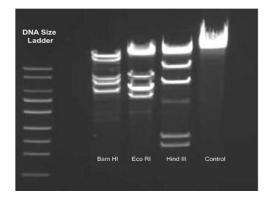
Location: Boston University Center for Advanced Biotech

Data Format:Gel images linkEnzymes:Xbal, PstlLadder:NEB 2Kb ladder

Protocols: Knight Restriction Digest Protocol

Knight Agarose Gel Protocol

Additional Comments: 1% TAE gel; Sybr Safe dye used for staining



### **Growth Curve**

#### **BASIC INFORMATION**

To assess what effect, if any, our genetic parts have on the growth rate of E.coli. Purpose:

Chassis:

Strain: Bioline α-Gold

Protocols: Purdue iGEM Growth Curve Protocol

Date: 7/9/13

#### **GROWTH CONDITIONS**

Media Type: Luria Broth (LB) 10mL Culture Tube Vessel:

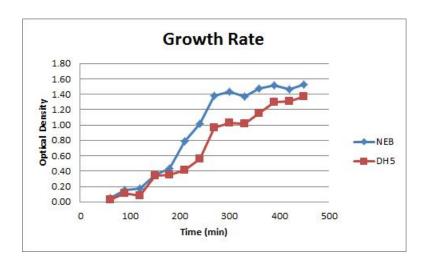
Volume: 5mL

Incubation: 37°C, 250 rpm

#### **MEASUREMENT INFORMATION**

Growth Curve (OD vs Time) Data Type: Location: **Bindley Bioscience Center** 

Machine Name: Time Interval: 30 min **Total Time**: 420 min



Additional Comments: This is where you would write additional comments about the part or its

functionality.

Reference: This is where any references or acknowledgements you need to make will go.

# Flow Cytometry

#### **BASIC INFORMATION**

Measure fluorescence for induction curve data Purpose:

Data-Type: Single-cell fluorescence

Boston University Center for Advanced Biotech Location:

SORP 4B-2YG-1BV, ACDU (FACSAriall) Machine Name:

FCS 3.0 data files link? Data Format:

(option to link to online protocol) Protocol Details: 445nm, 40mW:(515/20nm); Lasers (Filters):

488nm, 50mW: (488/10, 515/20, 545/35, 610/20, 710/50nm);

561nm, 40mW: (610/20, 660/20nm)

## PRE-INDUCTION GROWTH CONDITIONS

Media Type: Luria Broth (LB) Vessel: 5 mL tubes Volume: 2 mL

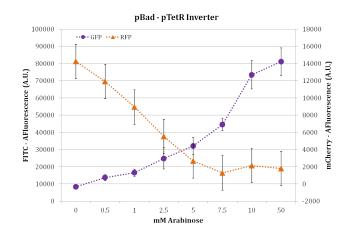
Incubation: 37°C, 300 rpm

Time (min):

#### INDUCTION GROWTH CONDITIONS

Media Type: Luria Broth Vessel: 500 μL tubes 200 μL Volume: Incubation: 37°C, 300 rpm

Time (min): 870



# **Future Work**

As of 7/7/13, the following needs to be completed or added:

- Flow Cytometry Modeling Data and Information
- Promoter Strength
- Induction Curve
- Mass Spectrometry