# Lab Notebook Entry 40

### **Date Performed**

30-May

# **Procedure**

2.0 PCR of pTlpA36-wasabi and of previous gel extracted pTlpA

## **Details**

Purpose: See reasons above, but I screwed up and ruined our PCR product by adding the wrong buffer.

#### **Protocol**

Use the protocol for PCR in the 2017 Protocol file, BUT THIS TIME DILLUTE 2uL OF THE TEMPLATE IN 18 uL OF PCR WATER IN A SEPERATE TUBE (label the tube and store in freezer box for future), THEN USE 1uL OF THAT AS TEMPLATE. Use "switch\_and\_gibsonFOR" and "switch\_and\_gibsonREV" as primers. Use the following heat cycle program: 95C for 30secs, 30 cycles of [95C for 10secs, 60C for 30secs 72C for 3mins], 72 for 2mins, 4C for infinity (just put in 99 hours or whatever the max is)