

**iGEM TU/e 2017**  
Biomedical Engineering

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## Protein Expression

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# 1 Protein Expression

**Estimated bench time:** 60 minutes

**Estimated total time:** 17 hours

**Purpose:** Protein expression of the bacteria.

It is essential to work near the Bunsen burner at all times.

## 1.1 Materials

- 1.5 ml cuvettes
- LB-medium
- Aluminum foil
- Antibiotic stock(s)
- Arabinose (20%)
- Cell Density Meter (OD600)
- Fresh culture of bacteria containing the right plasmid(s).
- Incubator
- IPTG (1 M)
- Pipettes and tips
- Sterile culture tubes

## 1.2 Setup & Protocol

- Prepare a culture tube containing 7 ml LB and 7  $\mu$ l of both antibiotic stocks (overnight)
- Transfer small culture to large culture with same ratios in medium and antibiotics as the small culture.
- Grow the bacteria in the incubator at 37 °C and 160-250 rpm (depends on size culture).
- After 90 minutes: measure the OD600.  
OD measurement requires a blank measurement with 1 ml 2 LB.  
Pipette 1 ml of the culture in the cuvette and measure the OD600.
- Put the culture back in the incubator (37 °C and 250 rpm). Regarding the fact that a cell division cycle takes around 20 minutes, calculate the amount of time the culture needs to obtain an OD600 of 0.6. (the OD600 doubles after  $\pm$ 20 minutes)
- After the additional time: measure the OD600 again. Pipette 1 ml of the culture in the cuvette and measure the OD600.
- The amount left in the culture should be 5 ml. When the OD600=0.6 wrap the culture tube in aluminum foil (only if your protein is sensitive to light) and add
  - IPTG (0.5 mM working concentration in our case)<sup>1</sup>
  - 56.24  $\mu$ l arabinose (20%)<sup>1</sup>
  - If applicable non-natural amino acid (10 mM)<sup>1</sup>
 This makes the final concentration in the culture tube:
  - 0.5 mM IPTG
  - 0.2% arabinose
  - 1mM unnatural amino acid.
- Perform protein expression of  $\pm$ 15 hours at 18 °C and 160-250 rpm.

<sup>1</sup> It depends on your plasmid(s) what you need to add to initiate protein expression.