

# Red Modeling

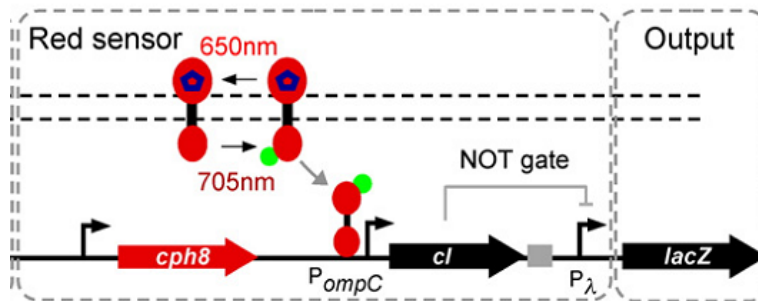
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## 1 Background Knowledge

*Ref: Multichromatic Control of Gene Expression in Escherichia coli*

Transcription from the output promoter of the previously constructed red light sensor (PompC) is inversely proportional to the intensity of red light. But we want an activating one, so a genetic inverter was placed between the red light sensor and lacZ. The CI repressor from phage  $\lambda$  is expressed as the output of PompC, and lacZ is expressed under the control of a CI repressible promoter. Dark exposure therefore results in high-level production of CI repressor and repression of lacZ transcription, while exposure to red light relieves this repression.



In our iGEM project, however, we will use RFP instead of lacZ as final reporter.

## 2 Modeling

### 2.1 Abstract

The passway is clear: Sensor – Inverter – RFP

The sensor is about the light-intensity induced chemical reaction. The Inverter is the  $\lambda$  one.

The promoter of red-light sensing protein *cph8* is constant. When exposed to different wave-length light, the *chp8* protein will be in different ground state. In the non-red ground state, the protein will pass a chemical group to *OmpR*, which then binds and activates transcription from  $P_{OmpR}$ , which will lead to the expression of repressor of RFP's promoter.

### 2.2 Definations

NULL

### 2.3 Equations

1. The total amount of *chp8* protein = production - degradation
2. When exposed to different wave-length light, the ratio between different ground state of protein *chp8*
3. The passing effect from *chp8* to *OmpR*, which should be a Hill equation.
4. The amount of *OmpR*
5. The expression rate of  $P_{OmpR}$
6. The production of  $\lambda$  repressor
7. The repressing effect to the promoter  $\lambda$
8. The amount of RFP