

# Detecting Heavy Metals in Water

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# Introduction

## *Problem*

- Canada-wide investigations reveal harmful levels of toxic metals are present in tap water [1]
- Make sure public is aware of the quality of their water
- Test water for the heavy metal contaminants, mercury (Hg) and lead (Pb)

## *Solution*

- A prokaryotic vector that is to be placed in a houseplant where fluorescent signals will indicate the presence of a heavy metal
- Yellow will be released when mercury is present, red will be released when lead is present, and yellow, red, and blue will be released when both are present

# The Model

# SBOL and Truth Table

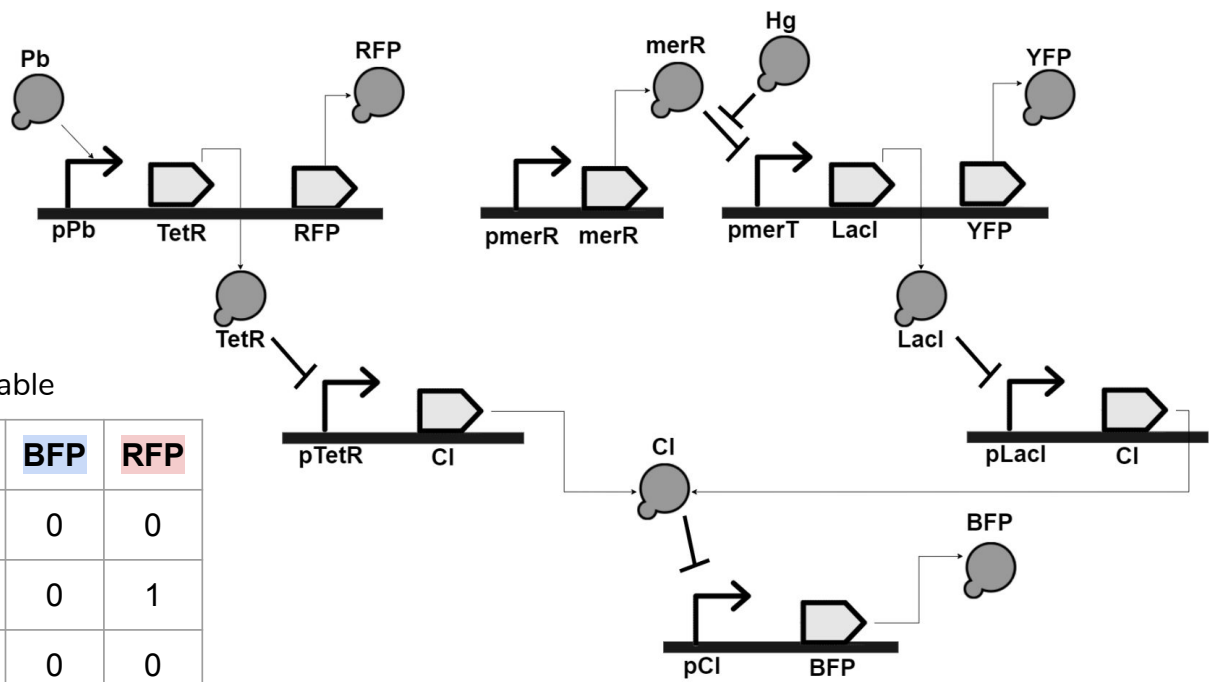


Table 1. Truth table

Pb	Hg	YFP	BFP	RFP
0	0	0	0	0
1	0	0	0	1
0	1	1	0	0
1	1	1	1	1

# Plasmid

Host: *Agrobacterium tumefaciens*

Plasmid: Tumour-inducing (Ti) plasmid (pBBR1MCS)

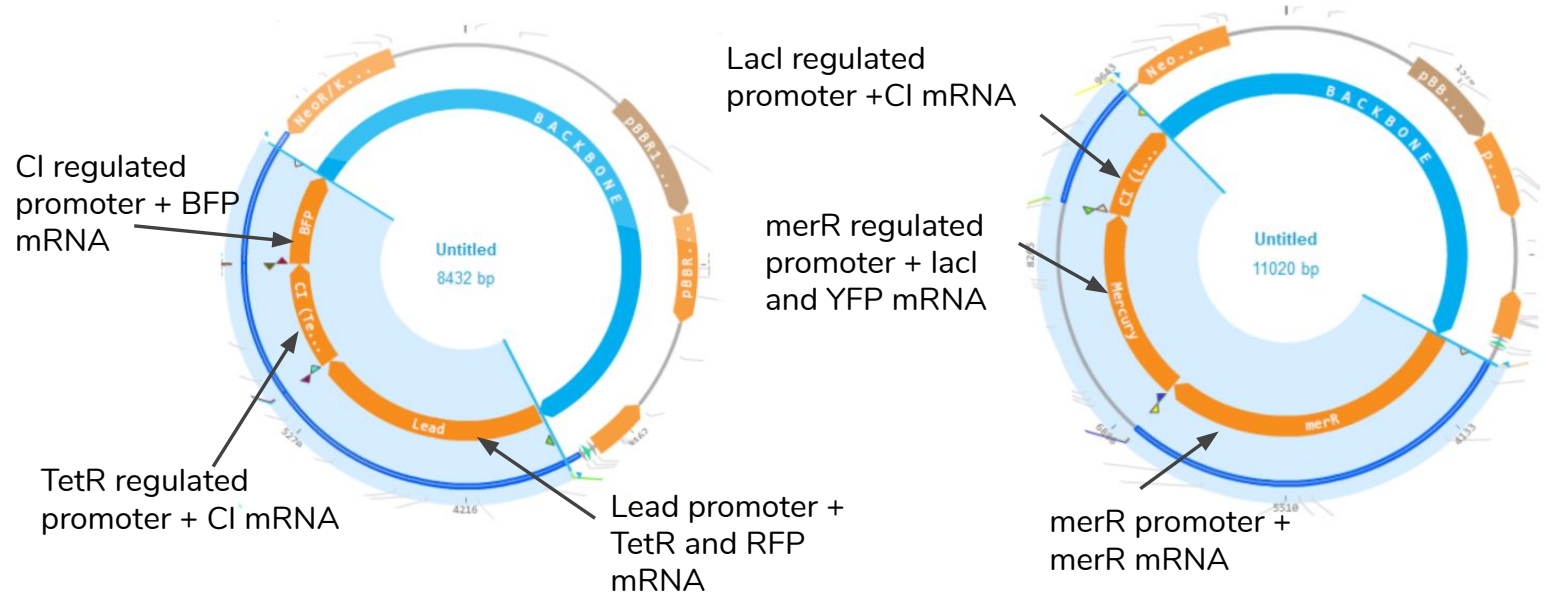


Figure 1. Tumour-inducing plasmid containing various parts that allow the visual response to mercury and/or lead. The left plasmid contains the system for lead detection (RFP) and detection of both metals (BFP) while the right is



# Assumptions

## Zero Rate Assumption

- mRNA dilution  $\ll$  mRNA decay; mRNA loss dictated by mRNA decay
- Protein decay  $\ll$  protein dilution; protein loss dictated by protein dilution

## Constant Rate Assumption

- Applies to decay, dilution, translation
- Processes occur at a constant rate ( $\propto k$ )

## Mass Action Kinetics

- For merR + Hg binding
- Reaction rate proportional to product of reactants ( $\propto k[\text{merR}][\text{Hg}]$ )

# Hill Kinetics + Hg-merR Dissociation

Repression of pTetR:	$\frac{d[mRNA]}{dt} = \frac{K_{trsc}}{1 + \left(\frac{[TetR]}{K_{hill\ repression}}\right)^H}$	Repression of pmerT:	$\frac{d[mRNA]}{dt} = \frac{K_{trsc}}{1 + \left(\frac{[merR]}{K_{hill\ activation}}\right)^H}$	$H = 1$
				$K_{hill\ repression} = 1\mu M$
Repression of pLacI:	$\frac{d[mRNA]}{dt} = \frac{K_{trsc}}{1 + \left(\frac{[LacI]}{K_{hill\ repression}}\right)^H}$	Activation of pPb:	$\frac{d[mRNA]}{dt} = \frac{K_{trsc} * \left(\frac{[Pb]}{K_{hill\ activation}}\right)^H}{1 + \left(\frac{[Pb]}{K_{hill\ activation}}\right)^H}$	$K_{hill\ activation} = 1\mu M$
				$K_{trsc} = 5.56 \times 10^{-14} \text{ M/s}$
Repression of pCI:	$\frac{d[mRNA]}{dt} = \frac{K_{trsc}}{1 + \left(\frac{[CI]}{K_{hill\ repression}}\right)^H}$	Hg-merR Dissociation:	$\frac{d[merR]}{dt} = -K_1 * [merHg]$	$K_1 = 4.5 \times 10^4 \text{ mM}^{-1}\text{s}^{-1}$

Figure 2. The Hill kinetics equations for the system, in which H is the hill coefficient, and K is a coefficient representing either activation/repression or transcription.



# Translation & Dilution Rates

Protein translation:  $\frac{d[protein]}{dt} = k_{trsl} * [mRNA]$

mRNA degn:  $\frac{d[mRNA]}{dt} = k_d * [mRNA]$

Protein dilution:  $\frac{d[protein]}{dt} = k_{d\ prot} * [protein]$

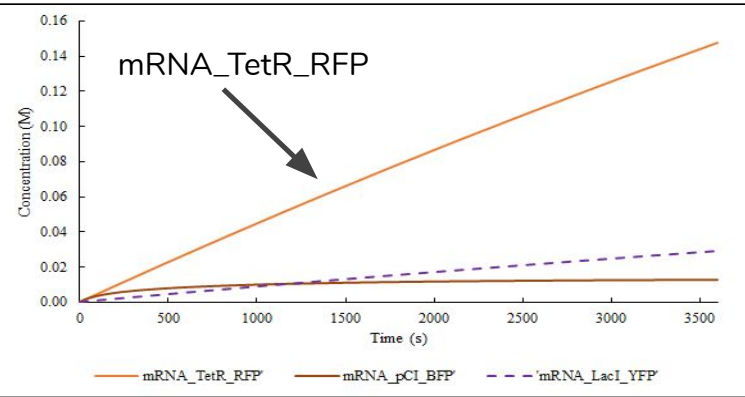
Figure 3. General equations for translation and dilution of the proteins used.



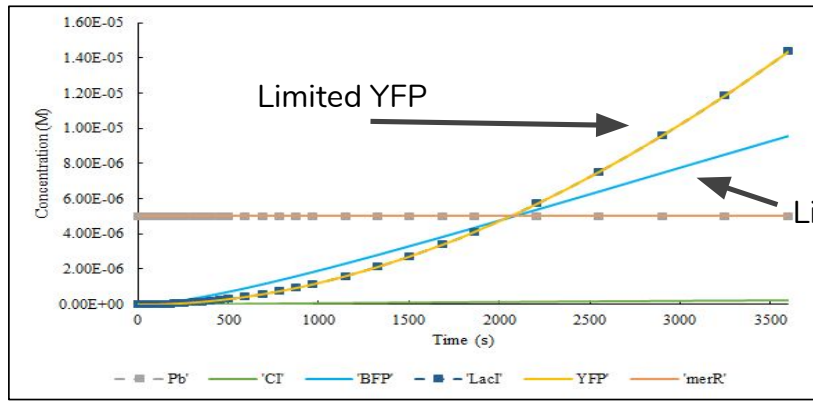
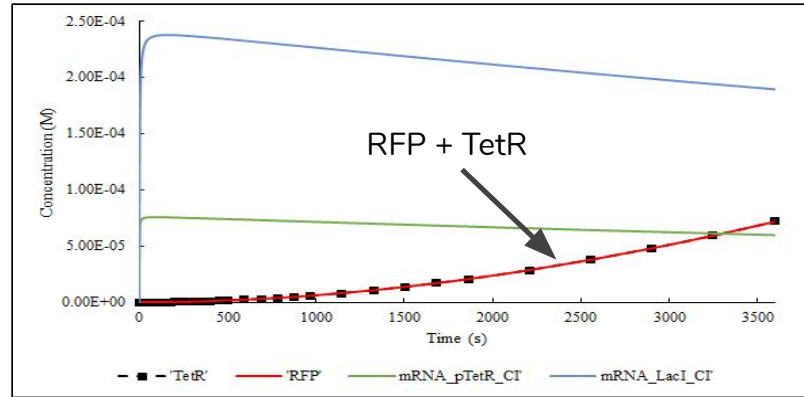
# Results and Discussion

# Simulation Output: Pb Present, RFP output

**Figure 4**



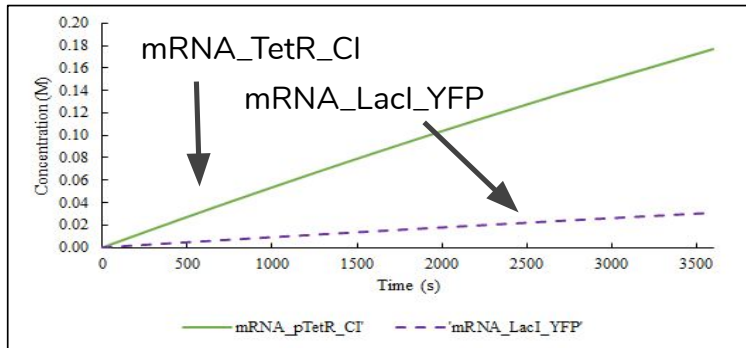
**Figure 5**



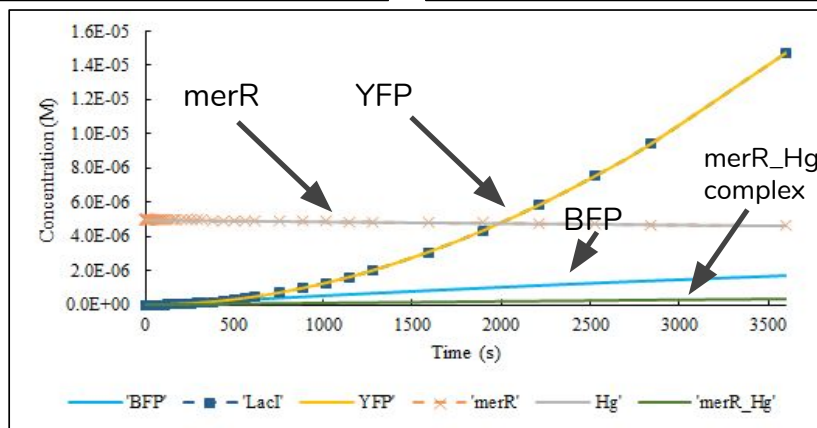
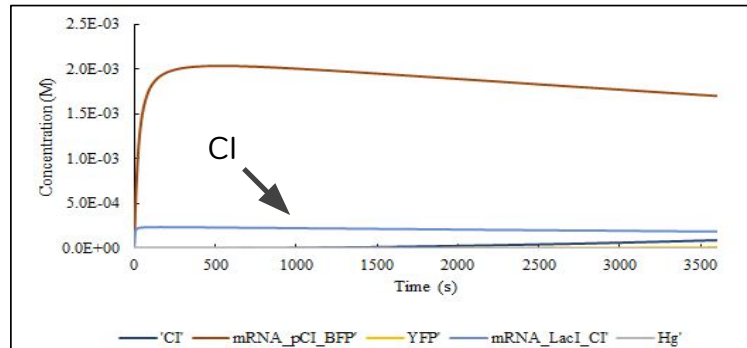
**Figure 6**

# Simulation Outputs: Hg Present, YFP output

**Figure 7**



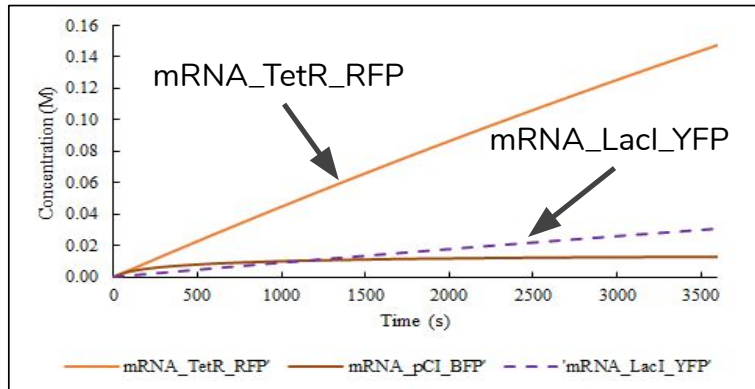
**Figure 8**



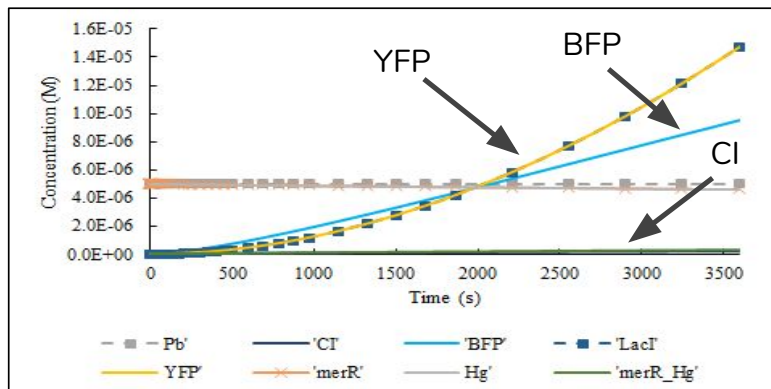
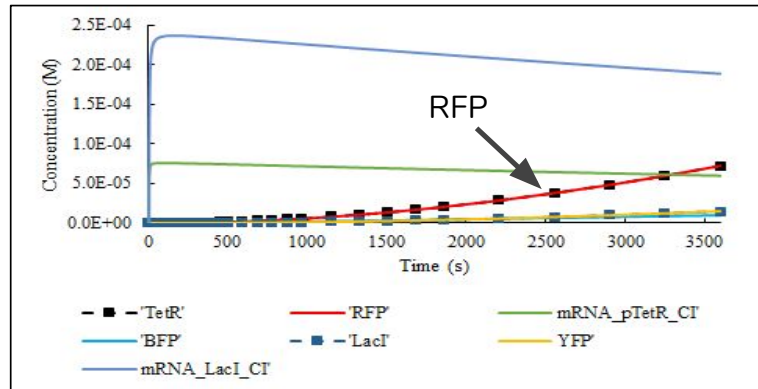
**Figure 9**

# Simulation Outputs: Both Metals Present, All Colours output

**Figure 10**



**Figure 11**



**Figure 12**

# Limitations and Conclusion



## Limitations

- Cannot sense the specific concentration of mercury or lead
- Colour prominence



## Conclusion

- Give a visual alert to homeowners through colour outputs
- Can detect lead whether lead and/or mercury is present

## Future Work

- Sensing contaminant concentration
- Detect additional contaminants



# References

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