# Mini-Project in Mathematical and Computational Modeling

École Polytechnique Fédérale de Lausanne, Switzerland

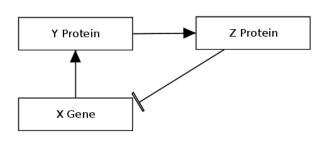
Florian + Dariush

### Introduction

Introduction to the article goes here Introduction to the article goes here

The Model

## Part A - One-Cell Model



(a) One-Cell Model

The gene mRNA X codes for protein Y which, in turn, activates transcriptional inhibitor Z. The resulting model behaves as a three-variable oscillator.

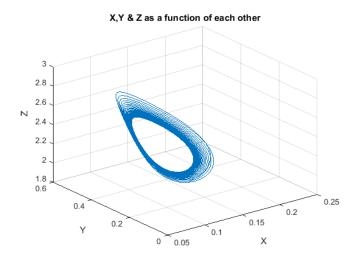
$$\begin{split} \frac{\delta X}{\delta t} &= v_1 \frac{K_1^n}{K_1^n + Z^n} - v_2 \frac{X}{K_2 + X} \\ \frac{\delta Y}{\delta t} &= k_3 X - v_4 \frac{Y}{K_4 + Y} \\ \frac{\delta Z}{\delta t} &= k_5 Y - v_6 \frac{Z}{K_6 + Z} \end{split}$$

translation rate of X $v_1$ degradation rate of X $v_2$ degradation rate of Y $v_4$ degradation rate of Z $v_6$ transcription rate of X $k_3$ transcription rate of Z $k_5$ 

Michaelis constant of X $K_1$ 

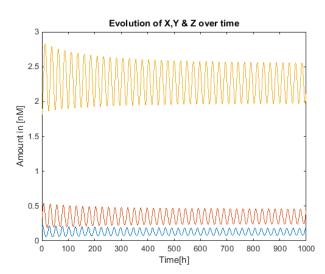
Michaelis constant of Y $K_4$ 

 $K_6$ Michaelis constant of Z



(a) Trajectories

The limit cycle is reached as the variations of X(t), Y(t) and Z(t) become fixed: The trajectories converge, non-lineary (the ellipse (where the blue stripes accumulate)



(b) Frequency spectrum

The amplitude of the three variations stabilize after a few distance between similar trajectories aren't regular) towards an hundred hours. The signal are not in phase but have the same, regular, frequencies.

### Figure 3:

Trajectories of X(t), Y(t) and Z(t) with initial conditions:  $X_0=0.16,\,Y_0=0.33,\,Z_0=1.8$  [nM] We observe on both graphs that Z(t) has the bigger amplitude of variation whereas X(t) and Y(t) have small amplitudes. Additionally, the convergence towards a single loop in (a) indicate that the frequencies of the signals are equal; this is illustrated as well in (b)

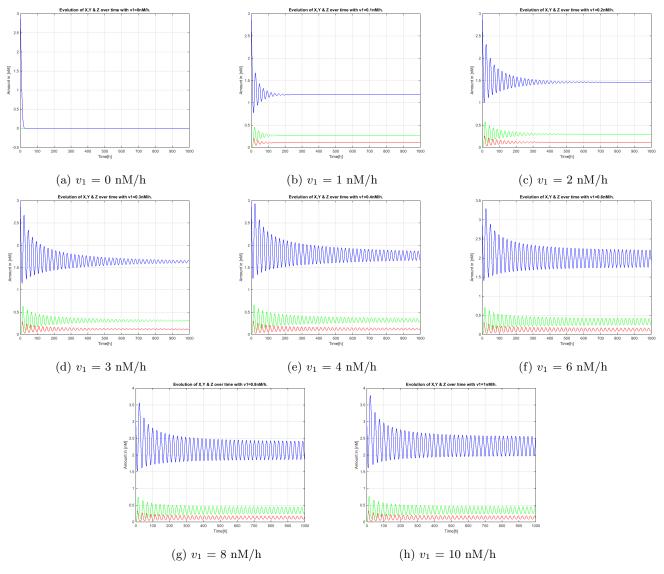


Figure 4: X(t), Y(t) and Z(t) with initial conditions  $X_0 = 0.16$ ,  $Y_0 = 0.33$ ,  $Z_0 = 1.8$  [nM] The first signal to fade is Y(t) and its oscillatory stability predicts stability of the system. We also observe that the signals converge towards null or the limit cycle in a non-linear fashion. At the opposite, it is rather difficult to predict the threshold value of  $v_1$  using those plots?

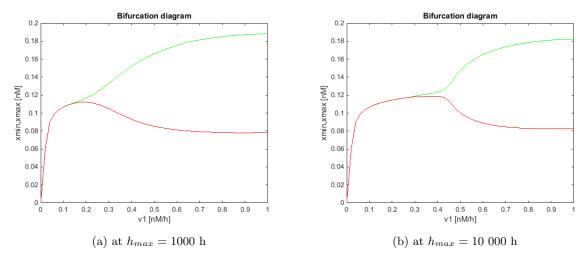


Figure 5: Bifurcation Diagram:  $X_{min}$  and  $X_{max}$  plotted at time intervals [9/10; 1] of  $h_{max}$ A limit cycle might be reached when  $X_{min} \neq X_{max}$ . However, the system needs to be run for enough time for the cycle to be reached, as the (a) suggests. (b) illustrates the non-linear convergence of the system; also the threshold for  $v_1$  seems to be around 4.5

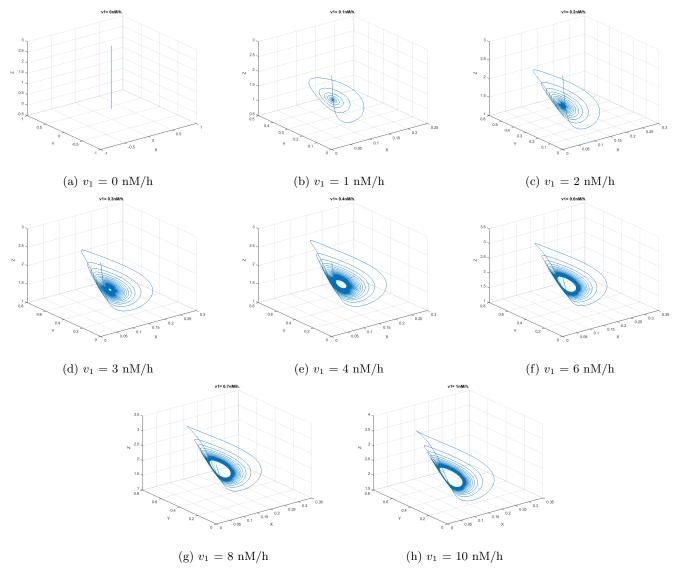


Figure 6: Trajectories when varying  $v_1$  with initial conditions  $X_0 = 0.16$ ,  $Y_0 = 0.33$ ,  $Z_0 = 1.8$  [nM]  $v_1$  has to reach a certain value for X(t) to be able to compensate its inhibition by Z(t) and therefore for the system to reach a limit cycle. We observe that this value is slightly greater than 4 nM/h, as the trajectories still converge to null in (e); there is an 'eye', even though it is smaller than in (f) and (g), since the timescale is not big enough to let the system dissipate completely.

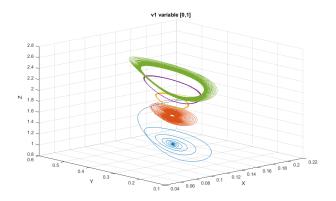


Figure 7: Superimposed trajectories at late timepoints with initial conditions  $X_0 = 0.16$ ,  $Y_0 = 0.33$ ,  $Z_0 = 1.8$  [nM] and  $v_1 = 0.1/0.3/0.5/0.7/0.9$  nM/h. We observe here that Z(t) tends to reach greater concentration stability with increasing  $v_1$ .

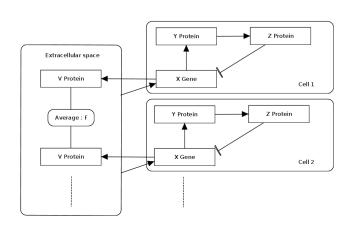
## Part B - Multiple Cells Model

 $v_1$ 

 $v_2$ 

 $v_8$ 

 $k_3$ 



(a) Multiple Cells Model The gene X codes for protein Y which, in turn, activates transcriptional inhibitor Z. In addition, gene X activates a positive feedback loop through the mean concentration of

extracellular protein  ${\cal V}$ 

$$\begin{split} \frac{\delta X}{\delta t} &= v_1 \frac{K_1^n}{K_1^n + Z^n} - v_2 \frac{X}{K_2 + X} + v_c \frac{KF}{K_c + KF} \\ \frac{\delta Y}{\delta t} &= k_3 X - v_4 \frac{Y}{K_4 + Y} \\ \frac{\delta Z}{\delta t} &= k_5 Y - v_6 \frac{Z}{K_6 + Z} \\ \frac{\delta V_i}{\delta t} &= k_7 X_i - v_8 \frac{V_i}{K_8 + V_i} \\ \end{split}$$
 where  $F = \frac{1}{N} \sum_{i=1}^N V_i$ 

translation rate of X degradation rate of X degradation rate of Y degradation rate of Z degradation rate of V transcription rate of X transcription rate of Z

 $k_7$  transcription rate of V  $k_1$  transcription rate of X  $K_4$  Michaelis constant of Y  $K_6$  Michaelis constant of Z  $K_8$  Michaelis constant of VK Coupling Constant

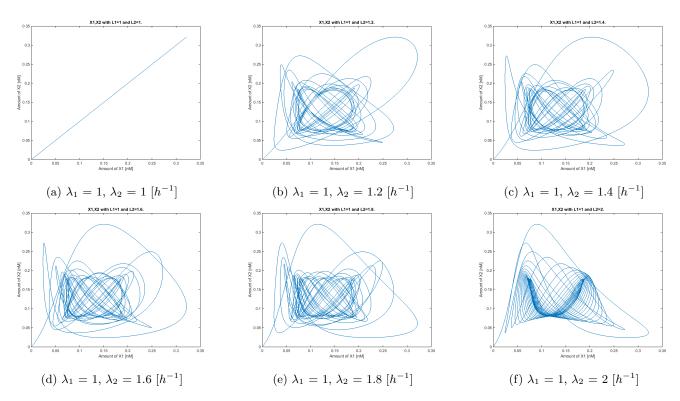


Figure 10:  $X_1$  and  $X_2$  trajectories with varying  $\lambda_i$  in a two-cells Model I don't really know what to say except 'wow it's cool' + square is max/min of Xs

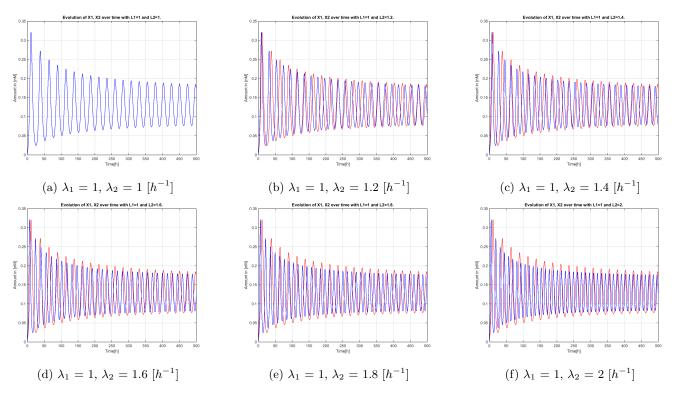


Figure 11:  $X_1(t)$  and  $X_2(t)$  trajectories in a two-cells Model We observe that

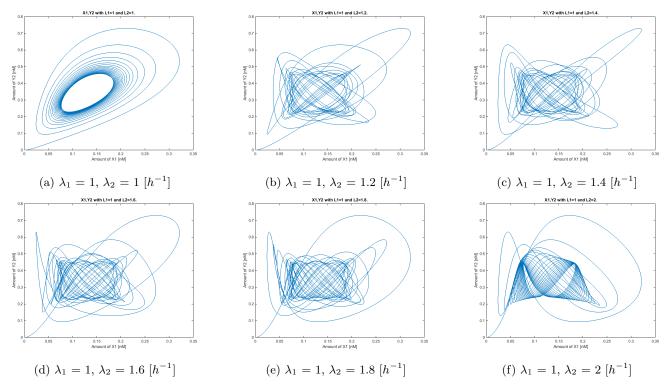


Figure 12:  $X_1$  and  $Y_2$  trajectories

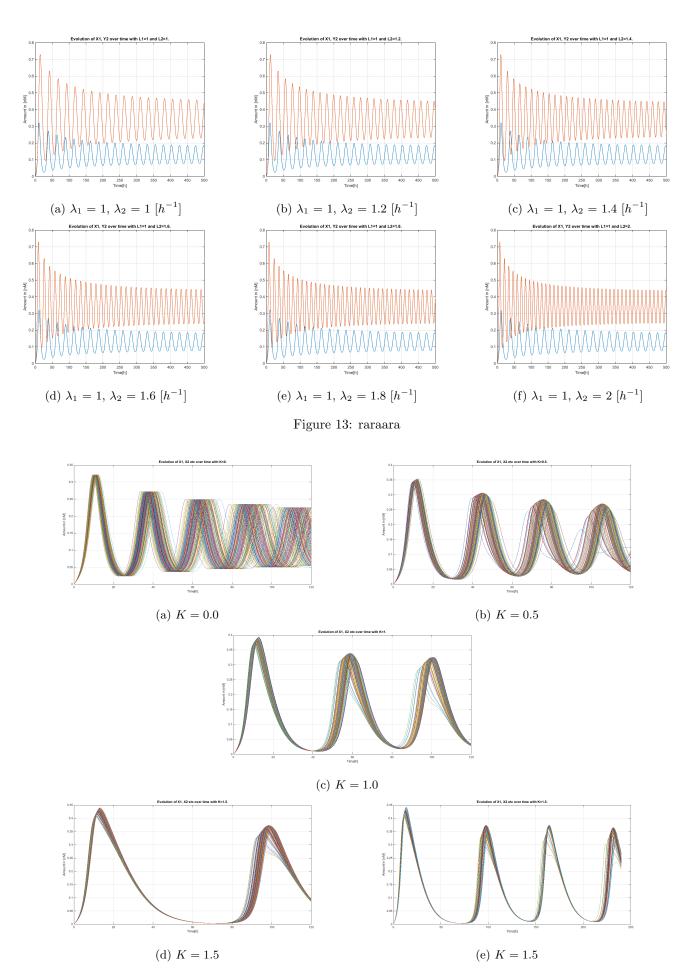
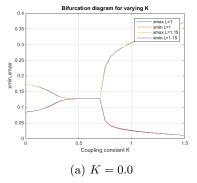
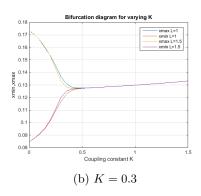


Figure 14: raraara





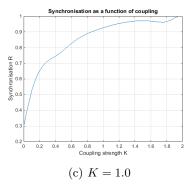


Figure 15: raraara