Prof. Dr. Tim Conrad

## 7 Project 7

- Deadline: 08.12.2019, 23:59
- All files need to be available through your GIT repository, in the directory "Project 7".
- You can work in teams up to 3 people. Please state in the report the group member names.
- If your code is in python, I must be able to run your code within a Google Colab notebook. If your code is not in Python or R, you
  must provide a manual how to compile and run it on a Linux machine.

## 7.1 Modeling and Simulation of complex biological systems

We will use the PySB library ( <a href="http://pysb.org/">http://pysb.org/</a>) to model and simulate complex biological systems. Your tasks are as follows:

(1) Implement and simulate the Tyson cell division model such that you can reproduce figure 3 from the original paper: "Modeling the cell division cycle: cdc2 and cyclin interactions" by JJ Tyson, PNAS (1991), 88 (16)

(This might help: https://github.com/LoLab-VU/pysb-tutorials/blob/master/tutorials/Tyson\_cycle\_tutorial.ipynb\_)

- (2a) Read the paper "Modeling, Simulating, and Parameter Fitting of Biochemical Kinetic Experiments" by D. Goulet, SIAM REVIEW, 58 (2).
- (2b) Implement and simulate the model described in the Goulet paper in a modeling framework of your choice (e.g. PySB, Copasi, Matlab etc.) such that you can reproduce figures 2 and 6.

## 7.2 Deliverables

Your need to upload all source codes and a report to your GIT repository.

- The report should be about 600-1200 words in length (this is roughly 1-2 pages, depending on your layout).
- The report must be delivered in PDF format using the BMC template (including the abstract as defined in project 5).
- The following sections must be present (you can add more if needed):
  - Tyson Cell Division Model
    - Background and Description of the model (including a brief discussion, why this is a complex model)
    - Brief description of the chosen model implementation and simulation
    - Results (including the produced figures)
  - Receptor Dimerization Model
    - Background and Description of the model (including a brief discussion, why this is a complex model)
    - Brief description of the chosen model implementation and simulation (including a list of the generated model ODEs)
    - Results and Discussion
       (including the produced figures and a discussion about the generated ODE model vs. the ODE model described in the paper)
  - Discussion: How does this project ("Complex Systems") differ from the projects of the previous weeks ("Data Science")? (Or why not?)