Identification of cell signaling pathways based on biochemical reaction kinetics repositories

Student: Gustavo Estrela

Advisor: Marcelo da Silva Reis (Butantan Institute)

May 2019

Instituto de Matemática e Estatística Centro de Toxinas, Resposta-imune e Sinalização Celular (CeTICS) Laboratório Especial de Ciclo Celular, Instituto Butantan

This project receives funding from FAPESP

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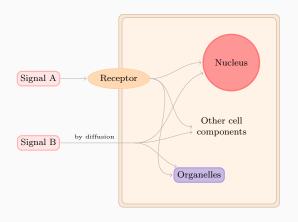
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Understanding the functioning of cell signaling is important in many biological areas.



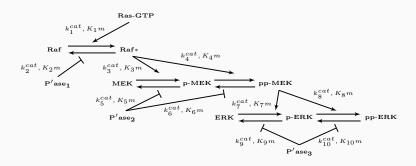
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We call the path of a signal a cell signaling pathway.

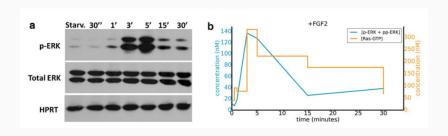
Cell Signaling Pathways

A cell signaling network can be characterized by a sequence of chemical reactions



Dynamic of Cell Signaling Pathways

We can summarize the state of the cell signaling pathway with measurements based on the concentration of some chemical species.



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A possible output to the problem is composed by:

- a model composed by a set of chemical reactions that are relevant for the biological experiment;
- information about the reaction rate constants of the model.

A Model for Cell Signaling Pathways

We model cell signaling pathways with a system of ordinary differential equations. These equations represent the dynamic of concentrations of chemical species.

Mathematical Models of Signaling Networks

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we can write the following equation:

$$\frac{d[C]}{dt} = k[A][B];$$

where k is a reaction rate constant.

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It is then desirable to systematically propose new models.

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On her work, the problem of identification of cell signaling pathways is treated as a feature selection problem.

Feature Selection Problem

The feature selection problem can be defined in the following way:

Given a set of features S and a error function err(), find subset $X \subseteq S$, with minimum error err(X).

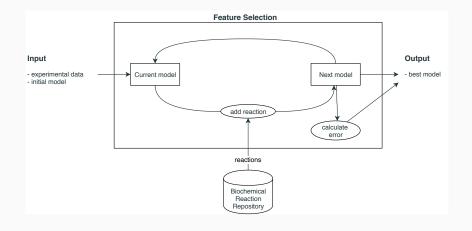
Feature Selection for Identification of Signaling Pathways

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Dynamic of Cell Signaling Pathways



Results of Wu's Methodology

However, the methodology worked satisfactorily only when the Kernel model was similar to a correct model.

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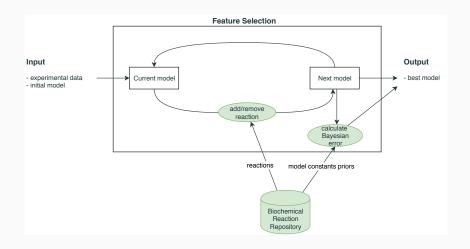
We can point three aspects of Wu's work that could explain its limitations.

- the database of interactions used could be more nearly complete;
- the search algorithm could also consider removing interactions;
- the cost function could implement a proper penalization of models;

What we Propose on this Project

We propose to create a methodology that uses a feature selection approach for identification of signaling pathways, tackling the difficulties encountered by Wu.

What we Propose on this Project



Objectives of this Project

• Build a database of interactions.

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- Create a cost function for models of signaling pathways.
- Formulate systematic modifications to a model as the search space of a feature selection model.
- Test the methodology on known signaling pathways and also on pathways of interest in our lab.

Bayesian Ranking of Models

SigNetMS

To perform model ranking, we created a Python package called ${\sf SigNetMS}.$

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Given a experimental data, a model and its constants priors, the software can calculate an estimative of p(D|M). This value is called the marginal likelihood.

SigNetMS

 ${\sf SigNetMS}$ is an open source software and it is available at:

https://github.com/gustavoem/SigNetMS

Experiments on Model Selection

Model Ranking Experiment

We tested SigNetMS on the ranking of models.

The experiment is based on the following procedure:

• Create 4 candidate models.

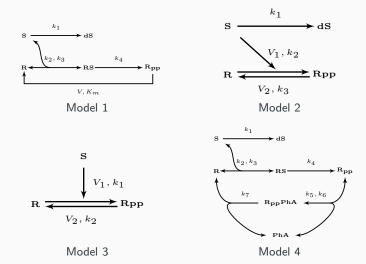
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- Rank the four models.

Model Ranking Experiment



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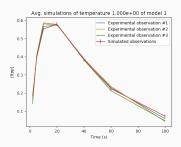
Note that:

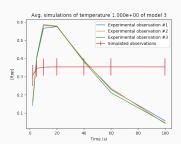
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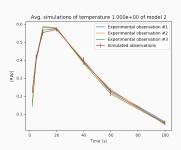
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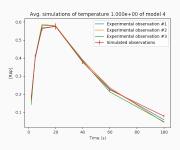
Note that:

- the correct model is ranked first;
- overly complex models are ranked worse than simpler models.





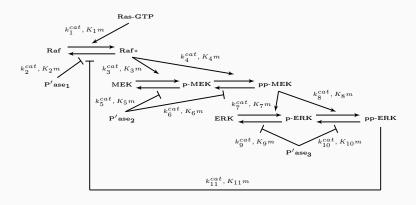


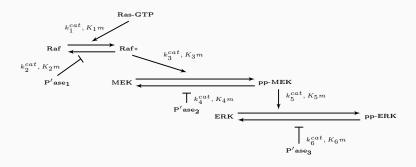


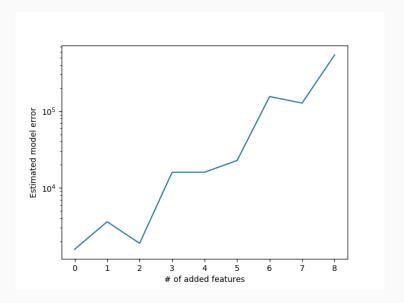
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The scores are calculated in respect to experimental data generated by a model that is present in the search space.







Future Work

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More activities are expected to be completed in this project, mainly the follow:

 Creation of a relational database of chemical interactions focused on our further applications.

Future Work

 Apply the method in ERK signaling pathways of tumor cell lines Y1 and HEK293. Thank you!