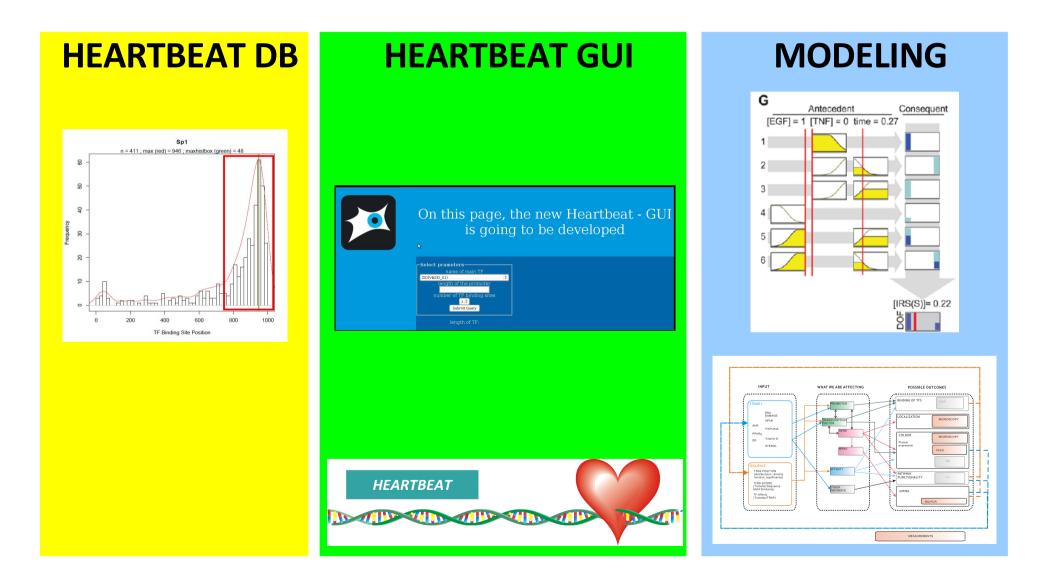
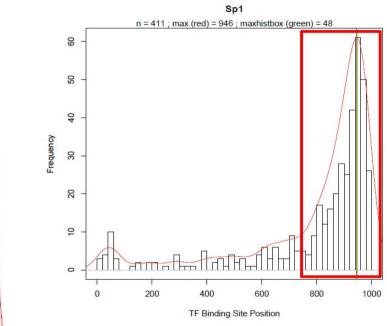
Modeling - Overview

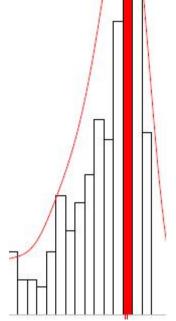


SEQUENCE QUALITY

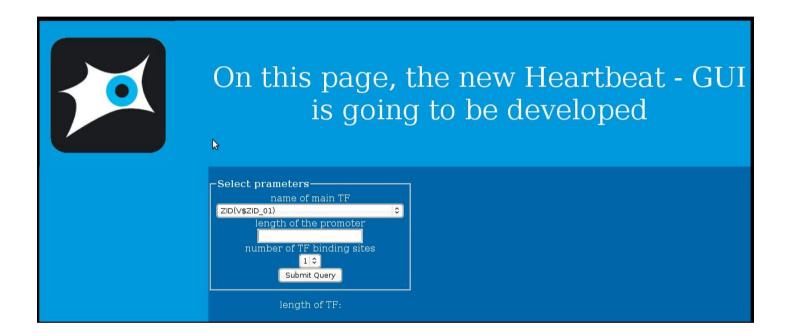
- statistics: characterization & scoring of our data by using:
 - TF affinity (TRAP)
 - density function
 - 20bp-AUC
 - peak amplitude
 - peak width(?)
 - sequence information
 - conserved?
 - position weight matrix
- Documentation



Area under the density function scaled with the amplitude is a measure for the significance of a peak

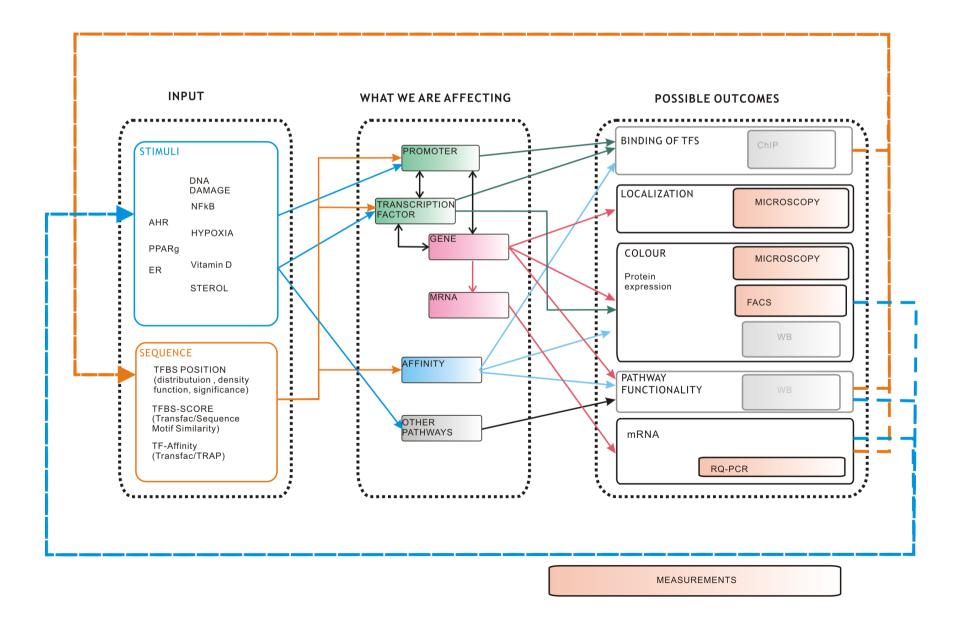


HEARTBEAT – GUI



- http://igem.boquant.uni-heidelberg.de/embperl/main.epl
- Embperl-based web page (perl code embedded in html)
- Design will correlate with the wiki's design

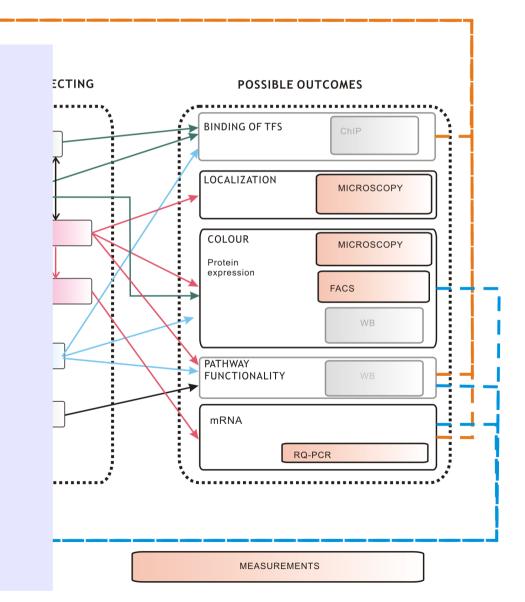
Modeling Our Network



Modeling Our Network

- during this part of the presentation -

- Please take a look at our basic network and collect ideas and available informations
 - experimental setup
 - available data/output
 - general ideas
 - hypothesis
- I will collect the handouts after the presentation & brainstorming :)

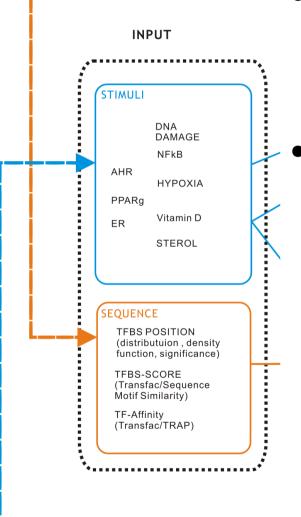


What modeling is good for...

so far we came up with the ideas...

- error checking / proof of concept
 - expectation vs experimental outcome
- *in silico* simulation
 - "assuming we have a promoter X with TFBS Y and Z… how would the outcome look like, depending on factor A, input B and stimulus C?"
- exclusive pathway activation
 - assume several synthetic promoters
 - combine single activation scenarios

model: basic concept



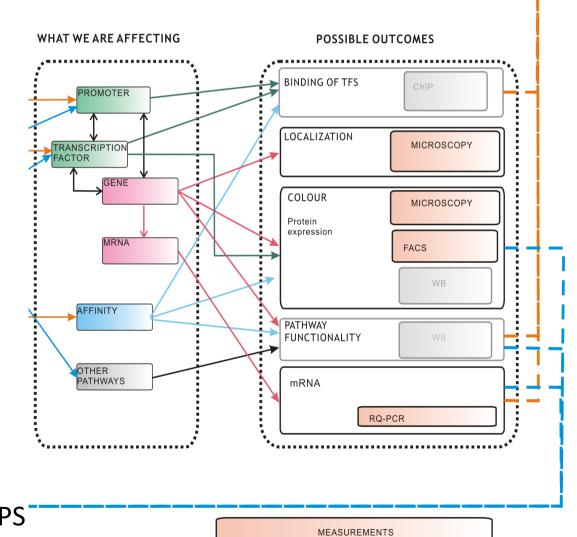
 build up model from available "reference data" (CMV/JeT) and also literature

possible INPUTS:

- STIMULI
 - stimulant type (drug?) / concentration
 - experimental setup
 - cell type
- SEQUENCE
 - TFBS position
 - TFBS sequence
 - TF affinity

model: basic concept

- "What we are affecting"
 - promoter
 - transcription
 - TF
 - mRNA level
 - other pathways
- OUTPUT
 - WHAT: colour, localization, promoter strength, mRNA concentration, dynamics (?)
 - HOW: microscopy, flow cytometry, RQ-PCR, (also POPS measurements(?))



modeling: first outcomes – and then?

- model training / model improvement
- expectation/simulation vs. experimental result
 - error checking:
 - what's wrong with our sequence?
 - will the right pathway be activated?
 - pathway induction
 - are there any cross-activation of pathways?
 - exclusiveness
 - are our promoters really exclusive?
- simulate several *in silico* / exp. relevant scenarios

Modeling: Tool

• Basic concept:

build up network topology – simulation – visualization

• we will use MATLAB Fuzzy Logic Toolbox

Basic Concepts

- list up what is available (data, pathway information)
- select data to use for building up the model
- define fuzzy rules
- select network subset (if whole network will be too complex)
- we will/can introduce you to fuzzy logic in an additional meeting

Modeling – what we need from you

- Promoter
 - which core / proximal promoter
 - sequence
 - natural vs. random synthetic vs. synthetic
- Data
 - values: FACS // Microscopy // qRT-PCR // PoPS (?)
 - experimental setup: stimulant, pathway... "SCENARIOS"
 - ! REFERENCE measurements !
- General
 - target pathways
 - literature

Modeling – what we need from you

- Promoter
 - which core / proximal promoter
 - sequence

basically we need everything and every small piece of information will help us! (but well documented, please ;-)

- what experimental setup: stimulant, pathway
- ! REFERENCE measurements !
- General
 - target pathways
 - literature

Computer Team – TODO

HEARTBEAT DB

- Improve localization of TF-binding site maxima
- Detailed
 documentation of
 HEARBEAT work flow

HEARTBEAT GUI

- Development of an interactive web-based
 GUI (in progress)
- Implementation of an algorithm for automatic sequence construction
- Documentation of the web-page

MODELING

- activate MATLAB licence
- build up model:
- collect sequences
- COLLECT DATA!
- simulate, simulate, simulate...