



# **Bioflux Analyses: Modelling**

In 1 collection

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#### **ABSTRACT**

Biofilm formation under shear flow conditions was monitored using the Bioflux1000 device (Fluxion Biosciences, Inc.). In short, Candida albicans overnight cultures were washed in pre-warmed RPMI medium. Cells were seeded for 2-5 sec from the outlet well into the channels of Bioflux1000 flow chambers, which were primed before with warm medium. The cells were allowed to adhere to the channels for 90 min without any flow, followed by removal of non-adherent cells by flowing fresh, pre-warmed RPMI medium for 5 sec. Shear flow was set for time series experiments over 24 h biofilm formation and images were captured every 20 min. Two channels were investigated in parallel having a 10 × magnification to allow a direct comparison between a mutant and a reference (wild-type) strain. Image capturing and stacks to movies was performed using the MetaMorph® Software (Molecular Devices).

An ODE model reflecting the logistic growth as well as the lag phase was fitted to the individual experiments. Fitting was carried out by minimising a cost function (unweighted least-squares-based) using the *Nelder-Mead* algorithm. Growth rate time series generated from the fitted model were used to compare wild type and mutant regarding the maximum observed growth rates at their respective time points.

All computations were performed using the programming language python (version 3.6.9) and the additional packages numpy (version 1.16.2), pandas (version 0.25.0), and scipy (version 1.3.1).

The modelling presented here uses a simple ODE model fitted to the individual Bioflux Ipix time series data sets (Fig. 1-A) in order to estimate the height as well as the time point of maximum observable growth (Fig. 2-B).

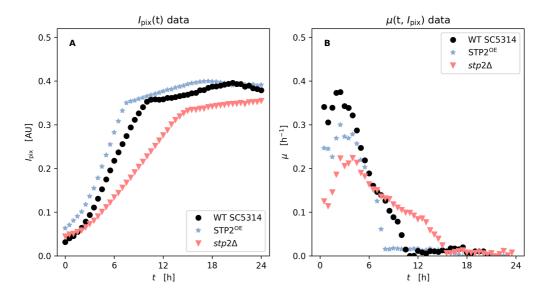


Fig. 1. Example of Bioflux time series data sets; A: medium pixel intensity; B: calculated growth rate.

Table 1 contains symbols and abbreviations used within the modelling approach and parameter estimation.

Symbol	Description	Unit
Latin		
$I_{\rm pix}$	Medium pixel intensity	AU
$I_{\mathrm{pix},0}$	Pixel intensity at start of experiment	AU
$I_{ m pix,max}$	Maximum of medium pixel intensity	AU
J	Cost function	_
t	Time	h
$t_{ m i}$	Lag time constant	h
$t_{ m d}$	Adherence time (difference between seeding and start of experiment)	h
Greek		
$\lambda$	Wolf-equation (to reflect lag phase)	_
$\mu$	Specific growth rate	$\mathrm{h}^{-1}$
$\mu_{\mathrm{max}}$	Maximum specific growth rate	$h^{-1}$
$\theta$	Vector of all possible parameter combinations	
χ	Verhulst-equation (to reflect logistic growth)	_
Ω	Vector of estimated optimum parameter values	
Indices		
data	Measured value	
model	Modelled value	

Table 1. Symbols and abbreviations used within the modelling approach and parameter estimation.

## 1.1 Ordinary differential equation for discontinuous bioprocesses

The modelling presented here was carried out using the ordinary differential equation for discontinuous bioprocesses with respect to the mean pixel intensity calculated according to the protocol 'Bioflux Analyses: Image Preprocessing'.

$$rac{dI_{ ext{pix}}}{dt} = \mu_{ ext{max}} \cdot \lambda(t) \cdot \chi(I_{ ext{pix}}) \cdot I_{ ext{pix}}$$

### **Growth Kinetics**

The growth of the culture was described using the following kinetics:

• logistic equation (Verhulst-equation; employed to modell the capacitive limit of the biomass accumulation)

$$\chi(I_{
m pix}) = 1 - rac{I_{
m pix}}{I_{
m pix,max}}$$

• lag phase (Wolf-equation; employed to modell the initial lag phase of the culture due to adaptation)

$$\lambda(t) = 1 - \exp\left(-rac{t+t_d}{t_i+t_d}
ight)$$

The model parameters  $\mu_{\text{max}}$ ,  $I_{\text{pix,max}}$ ,  $t_{\text{i}}$  as well as the start condition  $I_{\text{pix,0}}$  is estimated within Step No. 2. The adherence time  $t_{\text{d}}$  represents the time difference between the seeding of the cells and the start of the experiment/simulation.  $t_{\text{d}}$  was set to 1.5 h for all simulations in accordance to the experimental set-up.

1.2 Import of the required packages.

```
import numpy as np
import pandas as pd
import scipy.integrate
from scipy.optimize import fmin
```

1.3 Definition of the growth kinetics.

```
# logistic equation (Verhulst)
def chi(Ipixmax,Ipix):
    return 1 - Ipix / Ipixmax

# description of lag phase (Wolf-term)
def wolf(ti,td,t):
    return 1 - np.exp( - ( t + td ) / ( ti + td ) )

# growth kinetics
def mu(mumax,Ipixmax,ti,td,Ipix,t):
    return mumax * chi(Ipixmax,Ipix) * wolf(ti,td,t)
```

1 4 Definition of the ODE model.

```
# Model
def eq(indep_par,exp_par,start_t,end_t,incr):
    initial cond = indep par[0],0
    mumax,Ipixmax,ti = indep par[1:6]
    td = exp par
   t = np.linspace(start_t, end_t,incr) # time grid
    def funct(y,t):
                                            # ODE system
       Ipix = y[0]
                                            # biomass
        tin = y[1]
                                            # internal t (needed for Wolf-term)
        dIpixdt = mu(mumax,Ipixmax,ti,td,Ipix,tin) * Ipix
        dtindt = 1
        return [dIpixdt,dtindt]
    ds = scipy.integrate.odeint(funct,initial cond,t) #integrate
   return (ds[:,0],t)
```

## Parameter Estimation

2 Based on the ordinary differential equation above, the experimental Ipix time series data sets were simulated individually for the respective experiment. The time series were simulated for an experiment time of 24 h. Numerical integration was carried out employing Isoda from ODEPACK (scipy.integrate.odeint).

The objective of the parameter estimation was to identify the parameter values for the respective experiment. The model parameters  $\mu_{\text{max}}$ ,  $I_{\text{pix,max}}$ ,  $I_{\text{pix,$ 

$$J( heta) = \sum_{i=1}^n ig(I_{ ext{pix,data}} - I_{ ext{pix,model}}ig)^2$$
 $J(\Omega) = \min_{ heta} J( heta)$ 

2.1 Definition of the scoring function (least-squares).

```
# definition of scoring function (least-squares beased)
def score(parms):

# score difference between model and data points
ss = lambda data,model:(( data - model )**2 ).sum()

# get solution to system
Ipix_model,t_model = eq(parms,exp_rates,0,time,100)

# model index to compare to data
mt=np.linspace(0,time,100)
findindex=lambda x:np.where(mt>=x)[0][0]
Ipix_index=list(map(findindex,data.time))
data.Ipix_model = Ipix_model[Ipix_index]

return ss(data.Ipix,data.Ipix_model)
```

2.2 Execution of the parameter estimation.

```
initials = [Ipix_zero,mumax,ti,Ipix_max]
exp_rates = 1.5 # [h]
time = 24 # [h]
answ = scipy.optimize.fmin(score, rates, full_output=1, retall=1, maxiter=1000)
estimates = answ[0]
score_value = answ[1]
```

3 Based on the parametrised model Ipix and growth rate time series were simulated for each experiment (Fig. 2-A and B). The subsequent evaluations were carried out using the modelled time series.

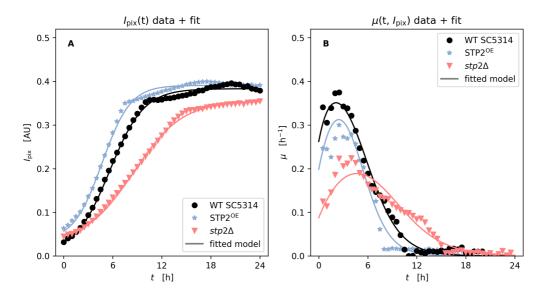


Fig. 2. Example of the fitted Bioflux time series data sets; A: medium pixel intensity; B: growth rate.

According to Fig. 2-B, the height as well as the time point of the maximum observable growth rate is recorded. These values are used for the comparison between the wild type an their respective mutant.

## Case Study

In order to illustrate the differences in the biofilm formation between wild type and mutant strains, the observed maximum growth rates are plotted against their respective time points (Fig. 3.1)-A. The differential plot (Fi. 3.1-B), conferring to subfigure A, presents these differences between the mutant strains (arrow head) and their respective wild types (arrow tail).

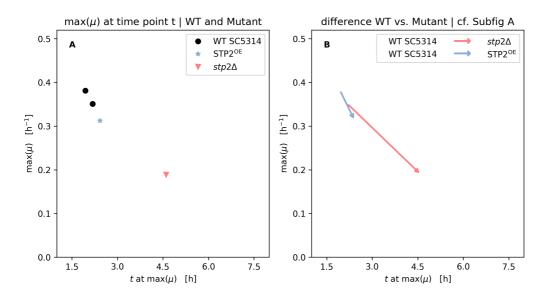


Fig. 3.1 Maximum observable growth rates  $max(\mu)$  at their respective time points for wild type and mutant strains; A: Scatter plot of  $max(\mu)$  at time point t from two examples (wild type vs. mutant); B: Difference plot for  $max(\mu)$  at time point t, wild type (arrow tail) to respective mutant (arrow head).

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