

On the use of the Gompertz model to predict microbial inactivation behaviour

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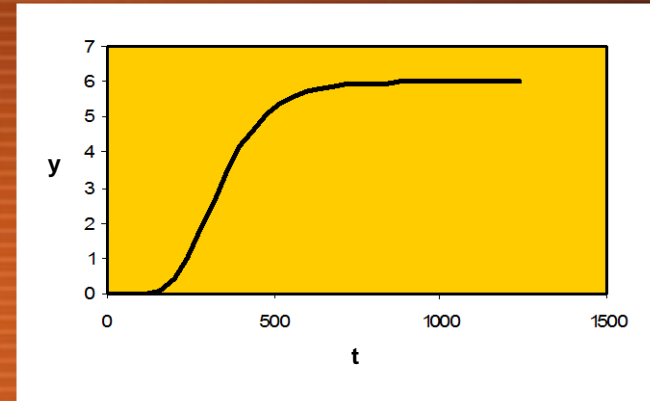
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Objectives

$$y_{\text{orig}}(t) = a \exp(-\exp(b - ct))$$

y population size
t time
a,b,c model parameters



Study of the original Gompertz equation

**modifications in order
to describe microbial
inactivation**

**re-parameterization of
the modified function**

biological meaning

Introduction

Mathematical modelling

- Food microbiologists are giving increased attention to microbial kinetics modelling
- The use of mathematical models that properly describe microbial behaviour under specific environmental conditions is important for predictive purposes and process design
- Mathematical models require the definition of parameters (e.g. the rate constant), which should be precisely estimated

Introduction

Gompertz equation

$$y_{orig}(t) = a \exp(-\exp(b - ct))$$

- Is an empirical sigmoidal relationship (*describe linear and nonlinear curves*)
- Introduced in food microbiology to describe asymmetrical sigmoidal shape of microbial growth
- Can be used to describe microbial inactivation behaviour
(*if properly modified*)

Introduction

 However...

- information related to function mathematical study and to methodology of re-parameterization is lacking



neglecting parameters biological meaning !

- only few works link model parameters to microbiological meaning (**growth/survival rate** and **lag time/shoulder**)

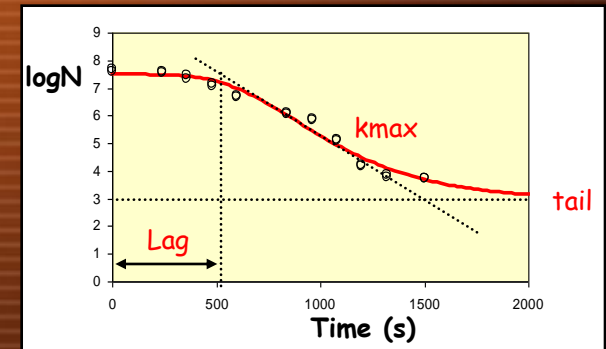


drawbacks in microbial kinetic models

Introduction

Insightful information can be obtained with re-parameterized forms

Microbial survival often presents a **lag**, a linear phase (corresponding to a maximum inactivation rate, k_{max}) and a residual population (**tail**)



Those phenomena should be included
in model parameters

Introduction

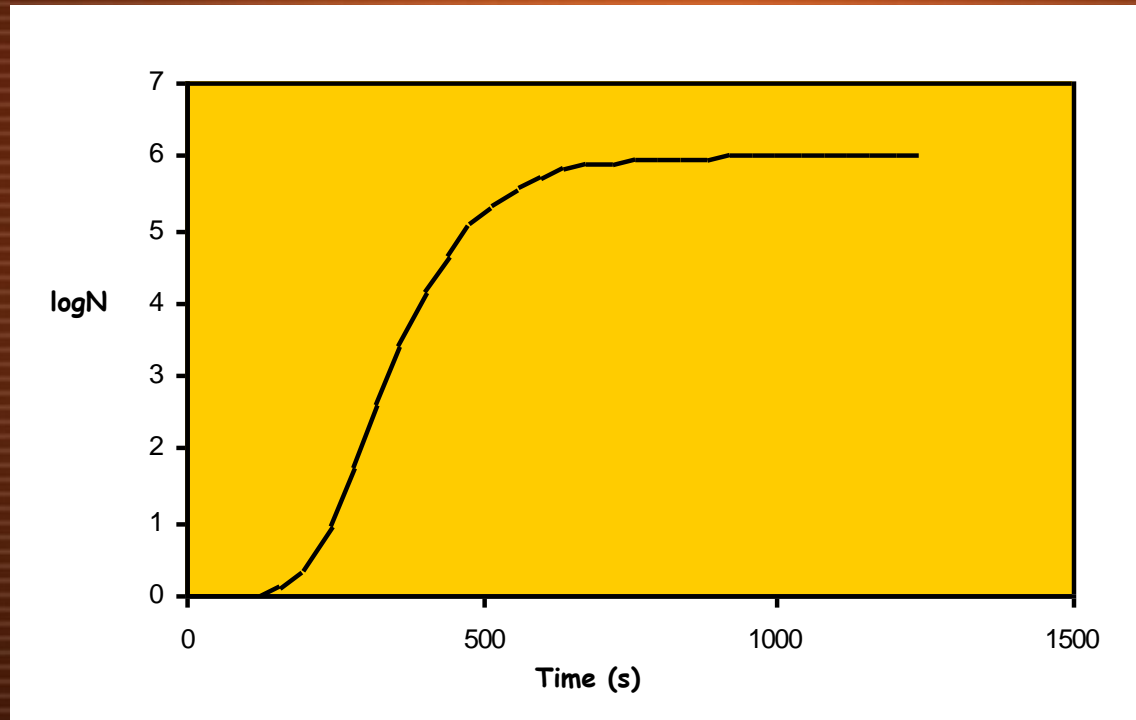
Study aim

modify in order to
describe
inactivation

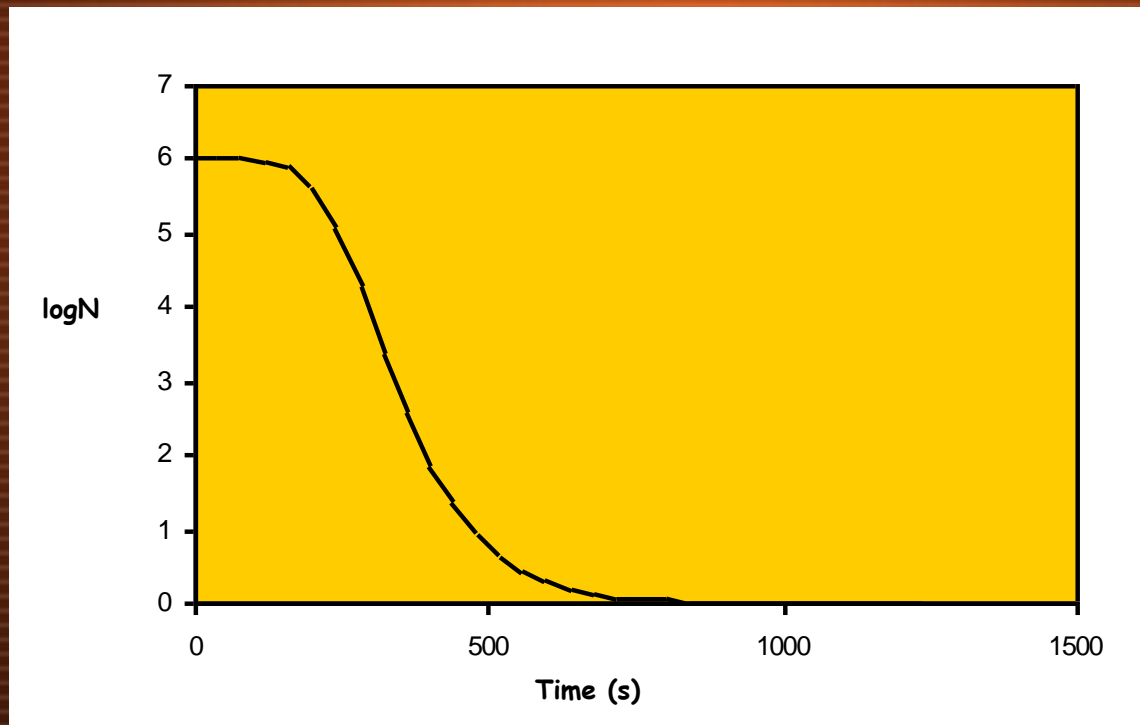


**Original
Gompertz model**

Microbial behaviour



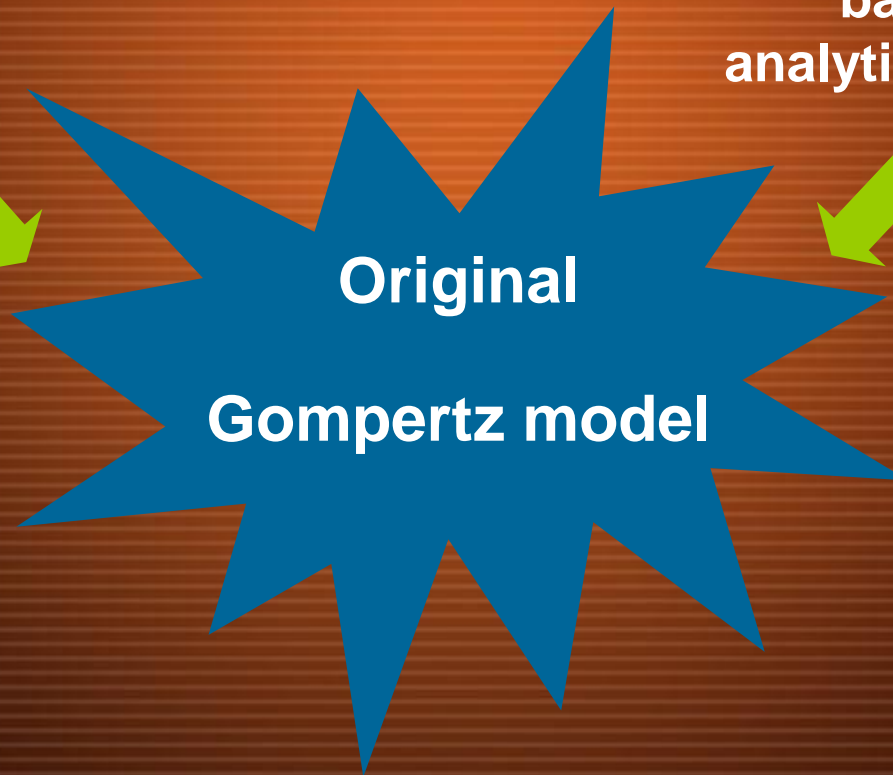
Microbial behaviour



Introduction

Study aim

modify in order to
describe
inactivation



re-parameterization
based on the
analytical study of the
function



Introduction

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modify in order to
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**Original
Gompertz model**

test
drawbacks/advantages
of the re-parameterized
forms



Approaches

Two different approaches were studied in order to obtain Gompertz modified model for inactivation kinetics:

- the logarithm of the microbial load ($\log N$) → **approach 1**



$$y_{inact}(t) = \log(N) = const - y_{orig}(t) \quad (1)$$

- normalized to the initial load ($\log N/N_0$) → **approach 2**



$$y_{inact}(t) = \log\left(\frac{N}{N_0}\right) = -y_{orig}(t) \quad (2)$$

Re-parameterization

Analytical study of the functions

➤ Function limits

time $\rightarrow \infty$ tailing effect

time $\rightarrow 0$ initial value

➤ Derivatives



Equations re-written

Parameters with microbiological meaning

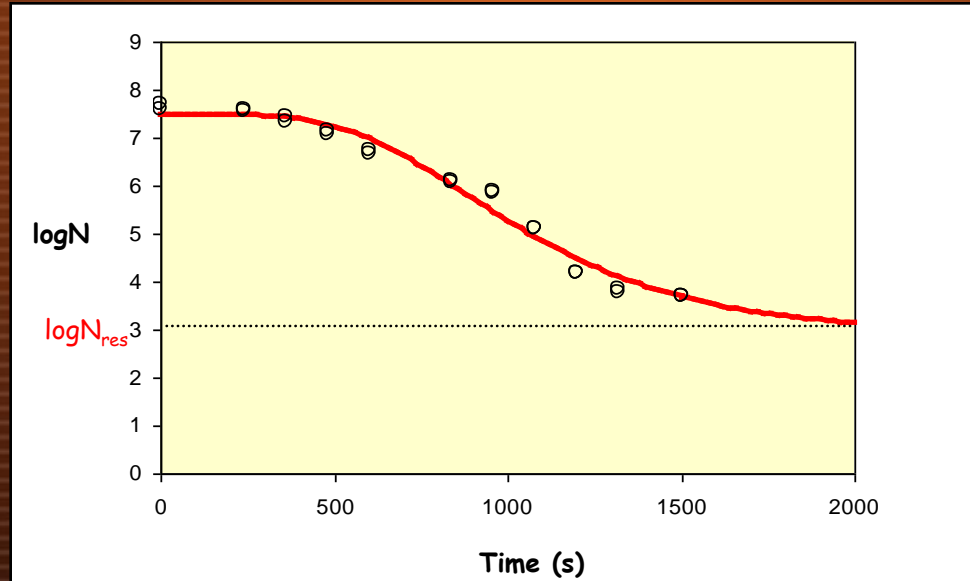
Re-parameterization

⚡ Model parameters

Tail

($\log N_{res}$)

asymptote of the function

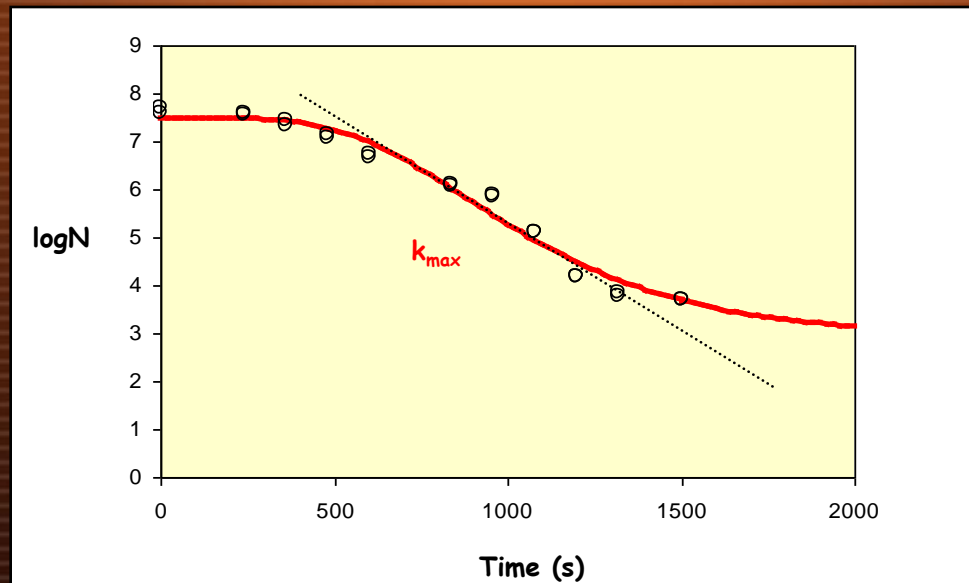


Re-parameterization

⚡ Model parameters

Maximum inactivation rate
(k_{max})

derived by calculating the 1st
derivative at the curve inflexion
point



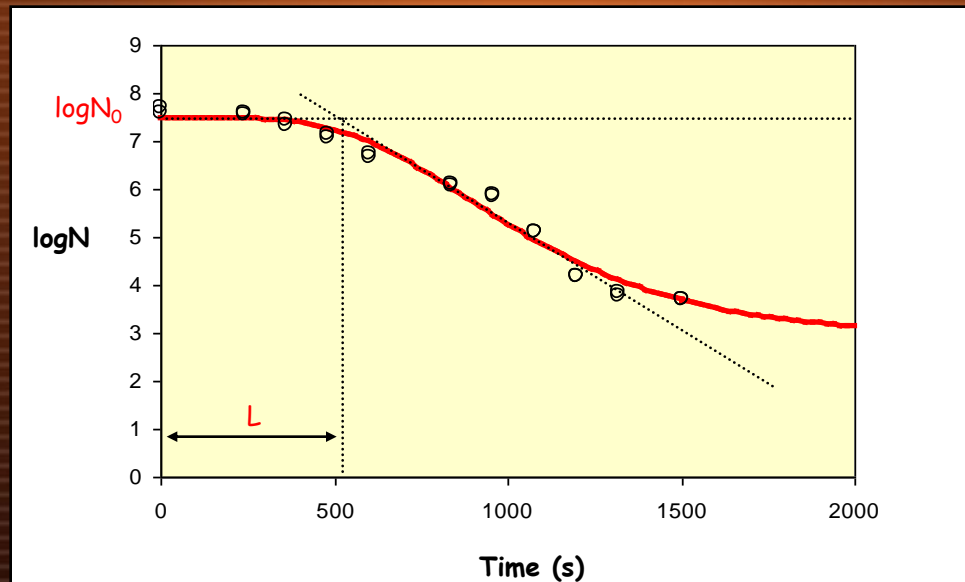
Re-parameterization

⚡ Model parameters

Shoulder

(L)

determined by the interception of the extrapolated tangent line with initial value (or time-axis - approach 2)



Re-parameterized forms

Approach 1

$$y_{inact}(t) = \log(N) = const - a \exp(-\exp(b - ct))$$



$$a = \log\left(\frac{N_0}{N_{res}}\right)$$

$$c = -\frac{k_{max} \exp(1)}{a}$$

$$b = L c + 1$$

4 parameters

$$y_{inact}(t) = \log(N_0) + \log\left(\frac{N_{res}}{N_0}\right) \exp\left(-\exp\left(\frac{k_{max} e}{\log\left(\frac{N_{res}}{N_0}\right)}(L - t) + 1\right)\right)$$

Re-parameterized forms

Approach 2

$$y_{inact}(t) = \log\left(\frac{N}{N_0}\right) = -a \exp(-\exp(b - ct))$$



$$a = -\log\left(\frac{N_0}{N_{res}}\right)$$

$$c = -\frac{k_{max} \exp(1)}{a}$$

$$b = L c + 1$$

3 parameters

$$y_{inact}(t) = \log\left(\frac{N_{res}}{N_0}\right) \exp\left(-\exp\left(\frac{k_{max} e}{\log\left(\frac{N_{res}}{N_0}\right)} (L - t) + 1\right)\right)$$

Case study

Studied approaches were validated on the basis of experimental thermal inactivation data of *Listeria innocua*



TSBYE



52.5
55.0
57.5
60.0
62.5
65.0



TSAYE



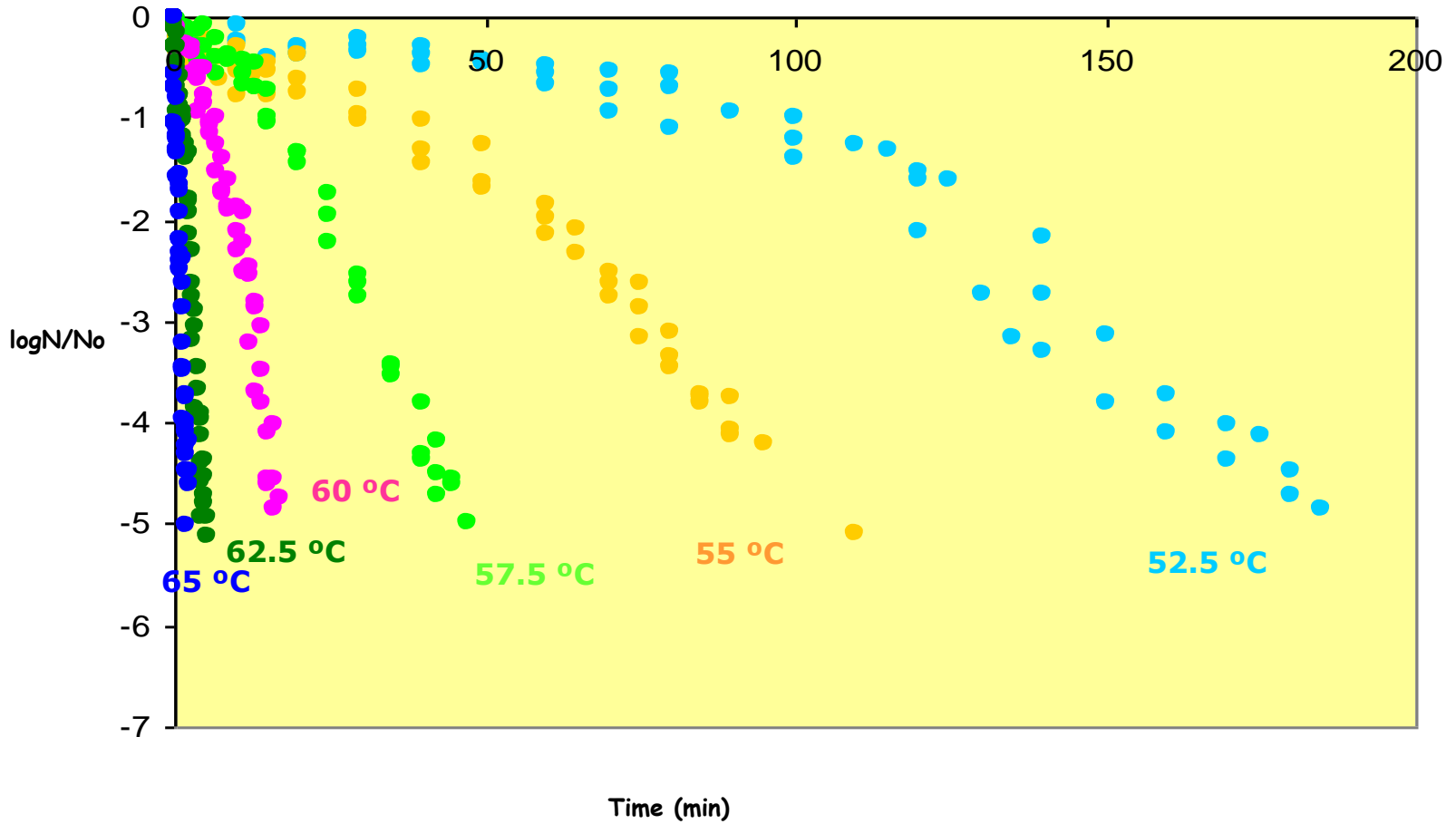
Methodology

➤ Kinetic parameters were estimated by non-linear regression analysis of isothermal data

least squares estimation

using STATISTICA TM v 6.0 software

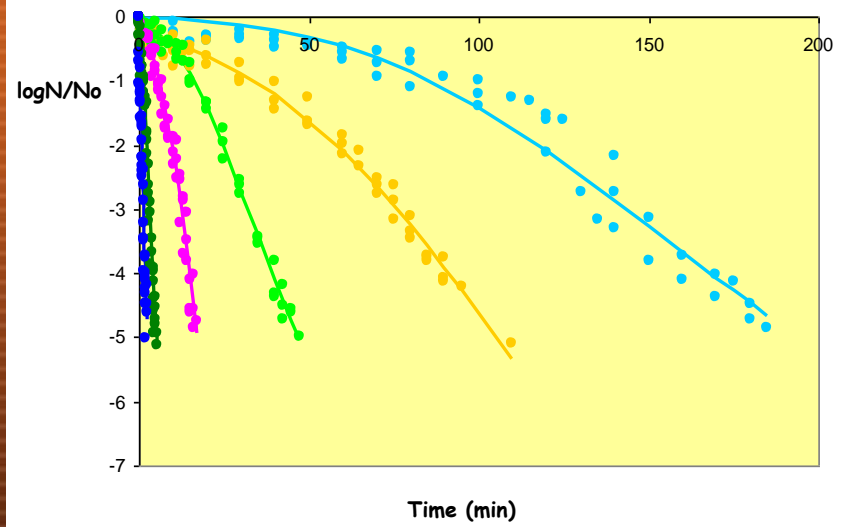
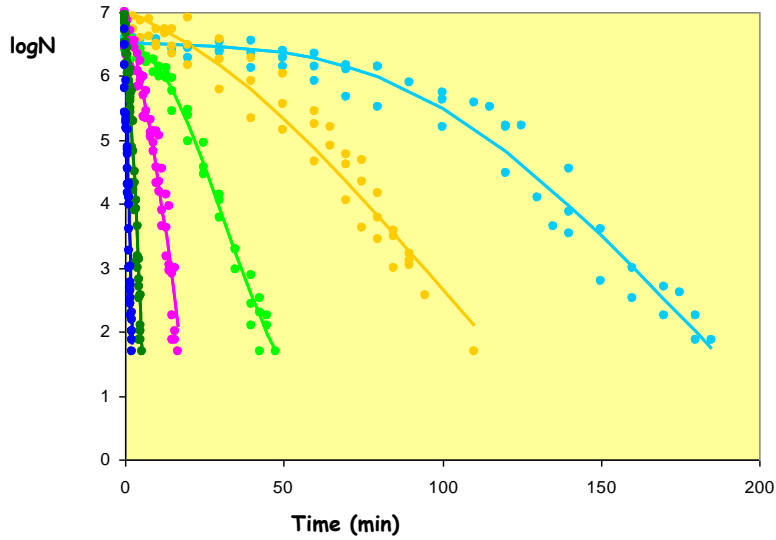
Results



Results

approach 1

approach 2



Results

- Gompertz modified model allowed accurate predictions of *Listeria innocua* inactivation in the temperature range considered
- Both approaches are quite similar

Results

Approach 1

Approach 2

$$y_{inact}(t) = \log(N) = const - a \exp(-\exp(b - ct))$$

$$y_{inact}(t) = \log\left(\frac{N}{N_0}\right) = -a \exp(-\exp(b - ct))$$

Regression Analysis

R_{adj}^2	0.876
Normality	√
Randomness	√

R_{adj}^2	0.970
Normality	√
Randomness	√

Parameter

Error (%)

Parameter

Error (%)

Parameter's Precision

$\log N_{res}/N_0$	26 - 37
k_{max}	11 - 17
L	43 - 563

$\log N_{res}/N_0$	14 - 28
k_{max}	7 - 13
L	16 - 63

Results

Approach 1

Approach 2

$$y_{inact}(t) = \log(N) = const - a \exp(-\exp(b - ct))$$

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$$error = \frac{95\% \text{ confidence interval}}{2} \times \frac{1}{\text{parameter estimate}}$$

Conclusions

- Gompertz allowed accurate predictions of *Listeria innocua* inactivation in the range considered
- Both approaches are quite similar
- Parameters were estimated with precision
- The quality of regression and parameters estimation were improved if normalized data is used
- The use of accurate models is a considerable tool to predict target pathogen's survival

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Thank You!

