

1 Fitting a growth curve

```
library(babar)

B092_1.file <- system.file("extdata", "B092_1.csv", package = "babar")
B092_1.data <- read.csv(B092_1.file, header=TRUE, sep="," ,
                        na.strings=c("ND", "NA"))
plot(B092_1.data, ylim=c(0,10))
```

Run the line to perform this command. This may take a minute or so to complete. Some typical output is shown below; this will appear in the command window.

```
t <- results_Bar4par$fit.t; y <- results_Bar4par$fit.ymean
lines(t,y,col="black",lwd=2)
```

Run this next line to perform the analysis with a different noise level, extracting and plotting the results as before, to see how the results change - you could try plotting using a different colour, making it easier to compare your results.

```
set.seed(11) ## for reproducibility
results_Bar4par <- Bayesfit(B092_1.data,model="Bar4par",inf.sigma=FALSE,
                           sigma=0.5)
```

Try changing the noise level manually to see how the results change again.
Repeat the analysis with the noise level inferred to see the difference.

```
set.seed(11) ## for reproducibility
results_Bar4par <- Bayesfit(B092_1.data,model="Bar4par")
```

1.2 Extension: fitting undetected data points

Next, we will look at fitting some data with undetected points; these are points with bacterial concentration below the level (or “threshold”) that we were able to detect using the experimental method.

Run these lines to import and plot curve *B119_5.csv*, which contains undetected points, choosing the threshold value 1.3.

```
B119_5.file <- system.file("extdata", "B119_5.csv", package = "babar")
B119_5.data <- read.csv(B119_5.file, header=TRUE, sep="," ,
                      na.strings=c("ND","NA"))
plot(B119_5.data,ylim=c(0,10))
threshold = 1.3
for (i in 1:nrow(B119_5.data)){
  if (is.na(B119_5.data[i,2])) {
    points(B119_5.data[i,1], threshold, pch=16)
  }
}
```

Run this line to perform the analysis including undetected points with threshold 1.3. Extract and plot the results as before.

```
set.seed(11) ## for reproducibility
results_Bar4par <- Bayesfit(B119_5.data, model="Bar4par", inc.nd=TRUE,
                           threshold=1.3)
```

Repeat the above steps with a different threshold value to see how this affects the results.

Model name	Parameters
linear	y_0, μ_{max}
logistic	y_0, y_{max}, μ_{max}
Bar3par	y_0, μ_{max}, h_0
Bar4par	$y_0, y_{max}, \mu_{max}, h_0$
Bar6par	$y_0, y_{max}, \mu_{max}, \lambda, \nu, m$

Table 1: The bacterial growth models available in the function. $y_0 = \log_{10}(x_0)$, where x_0 is the initial bacterial concentration, $y_{max} = \log_{10}(x_{max})$ where x_{max} is the maximum of the bacterial concentration, μ_{max} is the maximum specific growth rate, λ is the lag time, $h_0 = \mu_{max}\lambda$, and ν and m control the curvatures from the lag to exponential phase and exponential to stationary phase respectively.

Bayes' factor	Strength of evidence for/against first model or hypothesis
10 to ∞	Strong for
3 to 10	Substantial for
1 to 3	Barely worth mentioning for
1/3 to 1	Barely worth mentioning against
0.1 to 1/3	Substantial against
$-\infty$ to 0.1	Strong against

Table 2: The regions of Jeffreys' scale for interpreting the Bayes' factor.

Note: We can also use the model comparison techniques to calculate the Bayes' factor for the noise level inferred versus prescribed.