

# Document 3- R codes for the asymptotic analyses of a stage-structured matrix model for spotted hyenas infected with CDV

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This code presents the stochastic analysis of the matrix model. To account for parameter uncertainty in the calculation and sensitivity analyses of the population's growth rate  $\lambda$  (Fig. 1, Fig. 2) and the basic reproduction number  $R_0$  (Fig. 3) and calculate confidence intervals around the mean abundance of female hyenas projected throughout the study period and predicted beyond (Fig. 5), we used Monte Carlo iterations. This code requires the packages 'popbio' 2.4.4 (Stubben and Milligan 2007) and 'popdemo' 1.3-0 (Stott et al. 2012).

This code is structured as follows:

- 1) Monte Carlo iterations
  - a) Calculating / Plotting the mean  $\pm$  SD of  $\lambda$  (Fig 1) and  $R_0$
  - b) Plotting changes in population abundance - complete model and "no rank" model (Fig 5)
- 2) Sensitivity analysis of  $\lambda$ (Fig 2)
- 3) Sensitivity analysis of  $R_0$  (Fig 3)

Note that parameter and submatrix names may differ between main text and R codes.

- 
- 1) Monte Carlo iterations to calculate the mean  $\pm$  SD of  $\lambda$  and  $R_0$ , plot Fig 1 and Fig 5

First, we create a function to compile the population indicators we are interested in:  $\lambda$ ,  $R_0$ , population size, sensitivity values of  $\lambda$  and  $R_0$ .

```
start_time <- Sys.time()

library(popdemo)

## Warning: package 'popdemo' was built under R version 3.5.1
## Welcome to popdemo! This is version 1.3-0
## Use ?popdemo for an intro, or browseVignettes('popdemo') for vignettes
## Citation for popdemo is here: doi.org/10.1111/j.2041-210X.2012.00222.x
## Development and legacy versions are here: github.com/iaimstott/popdemo

library(popbio)

# First we create the function to store the population indicators we are interested in:

results<-function(Mproj, NGMstoch, popvec, senslambda, theta){

  tabmoy<-matrix(0,2,2)
  rownames(tabmoy)<-c("lambda", "rnot")
  colnames(tabmoy)<-c("mean", "sd")

  # Lambda
```

```

alllambda<-unlist(lapply(Mproj, lambda))
mean_lambda<-mean(alllambda)
sd_lambda<-sd(alllambda)
tabmoy[1,1]<-mean_lambda
tabmoy[1,2]<-sd_lambda

# Population size
popsizeMean <- apply(popvec, 1, mean)
popsizeSD <- apply(popvec, 1, sd)

# Sensitivity of lambda
meansens<-matrix(0, nrow=MCiter, ncol=42)

for (i in 1:MCiter)
meansens[i,]<-t(senslambda[[i]][2])

sens_lambdaMean<-colMeans(meansens)
sens_lambdaSD <- apply(meansens,2,sd)
names(sens_lambdaMean) <- rownames((senslambda[[1]]))
names(sens_lambdaSD) <- rownames((senslambda[[1]]))

# Sensitivity of R0
SensR0<- matrix(data = 0, nrow=ncol(theta), ncol=3)
allrnot<-0
sensi<-0
senslist<-0

if(t == "epidem")
{

allrnot<-unlist(lapply(NGMstoch, lambda))
mean_rnot<-mean(allrnot)
sd_rnot<-sd(allrnot)
tabmoy[2,1]<-mean_rnot
tabmoy[2,2]<-sd_rnot
senslist<-list()
for(r in 1:ncol(theta))
{
senslist[[r]]<- sens_elas_num(r, theta, delta=1e-4)
sensi<- sens_elas_num(r, theta, delta=1e-4)
SensR0[r,2]<-as.numeric(mean(sensi[[2]]))
SensR0[r,3]<-as.numeric(sd(sensi[[2]]))
SensR0[r,1]<-as.character(sensi[[1]])
}
}

return(list(tabmoy, popsizeMean, popsizeSD, sens_lambdaMean, sens_lambdaSD, alllambda, allrnot, means
})

```

Second, we load the text file containing the regression coefficients and the R file MC\_simulations. In this file, we first draw 1000 values from normal distributions with means equal to the regression coefficients of the MECMR model and with standard deviations equal to the standard errors associated with these regression coefficients. To obtain the MECMR parameter estimates and insure that they corresponded to probabilities bounded between 0 and 1, we back-transformed those simulated regression coefficients using the logit-function after accounting for the structural interactions and the temporal additive effects detected on those parameters.

```
data<-read.table("./RegressionCoefficient.txt", header = TRUE)

MCiter<-1000 # Number of Monte Carlo iterations (put 1000)

checkNodisease <- FALSE # this is to use the probabilities of infection estimated for each epidemic per

# Pre-epidemic period (1990-1992)
t<-"pre-epidem"
period <- t
popsize0<-100
Tmax<-3
source('MC_simulations.R')

tabprepidem<-results(Mproj, NGMstoch, popvec,senslambda, theta)

# Epidemic period (1993-1994)
t<-"epidem"
period <- t
Tmax<-3
popsize0<-tabprepidem[[2]][3]
source('MC_simulations.R')
tabepidem<-results(Mproj, NGMstoch, popvec,senslambda, theta)

# Post-epidemic period (1995-1999)
t<-"post-epidem1"
period <- t
Tmax<-6
popsize0<-tabepidem[[2]][3]
source('MC_simulations.R')
tabpost1<-results(Mproj, NGMstoch, popvec,senslambda, theta)

# Period of virus absence in the ecosystem (2000-2010)
t<-"post-epidem2"
period <- t
Tmax<-12
checkNodisease <- TRUE # this sets the probability of infection to 0
popsize0<-tabpost1[[2]][6]

source('MC_simulations.R')
tabpost2<-results(Mproj, NGMstoch, popvec,senslambda, theta)

# Projections in the future (2010 to 2020)
Tmax<-13
checkNodisease <- TRUE
popsize0<-tabpost2[[2]][12]
source('MC_simulations.R')
tabprojection<-results(Mproj, NGMstoch, popvec,senslambda, theta)
```

a) Calculating / Plotting the mean  $\pm$  SD of  $\lambda$  (Fig 1) and  $R_0$

```
data2 <- matrix(0,MCiter,3)

colnames(data2) <- c("pre-epidem", "epidem","post-epidem1")

data2[,1]<-tabprepidem[[6]]
data2[,2]<-tabepidem[[6]]
data2[,3]<-tabpost1[[6]]

# pdf("Figure1_Lambda.pdf",width=7,height=5) # uncomment this line to get the figure as a pdf

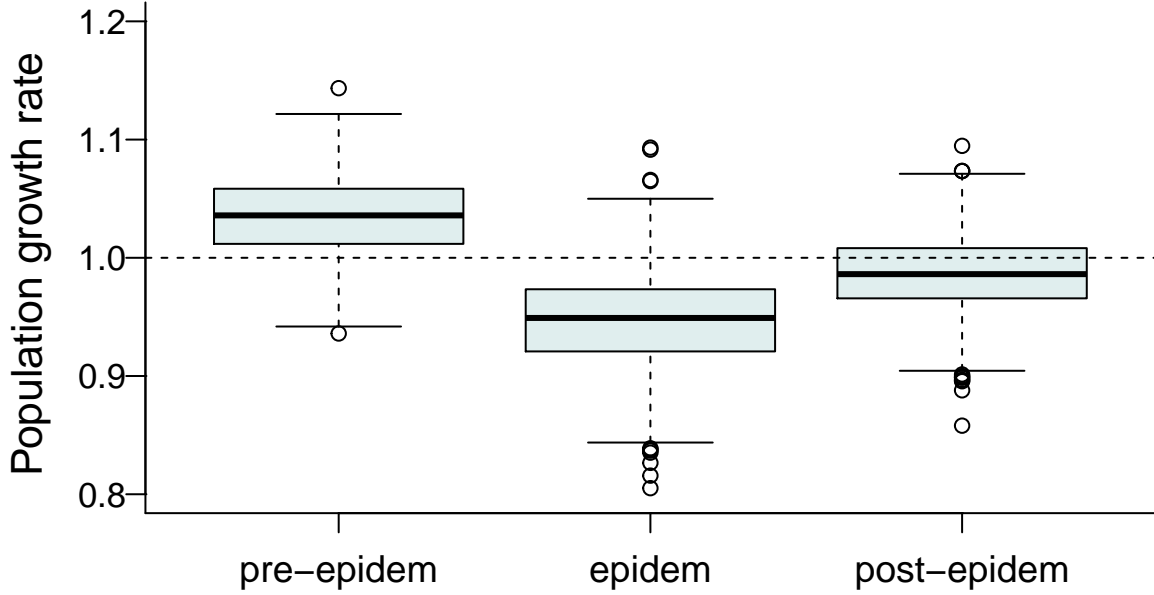
boxplot(data2, las=1, ylab = " ", xlab = "", axes =F, ylim = c(0.80,1.20), col="azure2")

x1 <- c(-0.1,0,1,2,3,4)
y1 <- c(0.70,0.80,0.90,1.00,1.10,1.20,1.30)
axis(1,at=x1,las=1,cex.axis=.6,tck=0, labels=NA) # this is to extend the axis (otherwise axes do not c
axis(2,at=y1,las=1,cex.axis=.6,tck=0, labels =NA)# idem for y

x <- c(1,2,3)
y <- c(0.80,0.90,1.00,1.10,1.20)

axis(1,at=x, las=1,cex.axis=1.2,labels =c("pre-epidem","epidem","post-epidem"))
axis(2,at=y,las=1, cex.axis=1.1,mgp=c(3, 0.5, 0))
mtext(side = 2, text ="Population growth rate ", line = 2.5, cex=1.3)

abline (h=1,lty=2)
```



*#dev.off()# uncomment this line to get the figure as a pdf*

In the following table, we display the basic reproduction number during the epidemic:

Basic reproduction number	Mean estimate	SD
<b>R0</b>	5.6836077	0.6563285

b) Plotting changes in population abundance - complete model and “no rank” model (Fig 5)

We can then project past (1990-2010) and predicted (2011-2020) temporal changes in  $\lambda$  (Figure 5) given parameter uncertainty during the 20 years of survey (1990-2010).

- To describe the past temporal changes during the study period (1990-2010) we calculated 1000 values of  $\lambda$  for each year of the study period accounting for temporal variations in the mean estimates between the pre-epidemic, epidemic and post-epidemic periods. We described with a Markov chain the changes in population size by multiplying the population vector of a given year by  $\lambda$  and reinitialized the population vector to the resulting vector of next year abundance. The initial population vector was defined as the product of 100 individuals and the stable stage distribution.
- To predict abundance of spotted hyenas for 10 years after the end of the study period (i.e. 2011-2020) we considered the 1000 block-matrices  $M$  implemented with the MECMR parameter estimates associated with the second post-epidemic period (2000-2010), and we determined the population vector of the number of individuals in the 22 demographic, social and infection states during the last year of the survey (2010). This vector was defined as the product of the mean abundance estimated in 2010 and the stable stage distribution. We then multiplied the matrices with this population vector to obtain 1000 population vectors and calculate the confidence intervals of the abundance the following year. These population vectors were then multiplied again by the simulated matrices to calculate the

mean abundance and its associated confidence interval in the following year. In such case the range of abundance increases with time as we use all the population vectors from a previous year (and not only the mean one) to calculate the range of population sizes the next year, accounting for the uncertainty around the parameter estimates. This Markov chain in which the population vectors of the next year only depend on the population vectors of the current year and of the simulated projection matrices was then reiterated for 10 years.

```

lambdaseq <- c(rep(tabprepidem[[1]][1,1], 3), rep(tabepidem[[1]][1,1], 2) , rep(tabpost1[[1]][1,1], 5),
lambdase <- c(rep(tabprepidem[[1]][1,2], 3), rep(tabepidem[[1]][1,2], 2) , rep(tabpost1[[1]][1,2], 5),r

popproj<-rep(0,31)
popproj[1]<-100
varmoins<-rep(0,31)
varplus<-rep(0,31)
varplus[1]<-100
varmoins[1]<-100

# loop on the retrospective (1990 to 2010)

for(t in 2:21)
{
  popproj[t]<-popproj[t-1] * lambdaseq[t]
  varplus[t]<-popproj[t-1] * (lambdaseq[t] + lambdase[t])
  varmoins[t]<-popproj[t-1] * (lambdaseq[t] - lambdase[t])
}

# projections 2010-2020

p<-1
for(t in 22:33)
{
  p<-p+1
  popproj[t]<- tabprojection[[2]][p]
  varplus[t]<- tabprojection[[2]][p] + tabprojection[[3]][p]
  varmoins[t]<- tabprojection[[2]][p] - tabprojection[[3]][p]
}

# To get a smoother curve, we delete here the meaningless abundances obtained at the 2 first years of t
popproj<-popproj[-c(22,23)]
varplus<-varplus [-c(22,23)]
varmoins<-varmoins[-c(22,23)]

mat2 <- cbind(popproj,varmoins,varplus)

mat2<-as.data.frame(mat2)
mat2$Model<-rep("normal", 31)
mat2$Time<-1:31
mat2$Time1 <-c(1990:2020)
mat2

##      popproj  varmoins   varplus  Model Time Time1
## 1  100.00000 100.00000 100.00000 normal    1  1990
## 2  103.59755 100.29873 106.89636 normal    2  1991
## 3  107.32451 103.90702 110.74200 normal    3  1992

```

```
## 4 101.73940 97.40575 106.07305 normal 4 1993
## 5 96.44493 92.33680 100.55306 normal 5 1994
## 6 95.15359 92.04800 98.25919 normal 6 1995
## 7 93.87954 90.81553 96.94355 normal 7 1996
## 8 92.62254 89.59956 95.64553 normal 8 1997
## 9 91.38238 88.39987 94.36489 normal 9 1998
## 10 90.15882 87.21625 93.10140 normal 10 1999
## 11 91.92250 89.36115 94.48384 normal 11 2000
## 12 93.72067 91.10922 96.33212 normal 12 2001
## 13 95.55402 92.89149 98.21656 normal 13 2002
## 14 97.42324 94.70862 100.13786 normal 14 2003
## 15 99.32902 96.56129 102.09674 normal 15 2004
## 16 101.27208 98.45021 104.09394 normal 16 2005
## 17 103.25315 100.37608 106.13021 normal 17 2006
## 18 105.27297 102.33962 108.20632 normal 18 2007
## 19 107.33230 104.34158 110.32303 normal 19 2008
## 20 109.43192 106.38269 112.48116 normal 20 2009
## 21 111.57261 108.46373 114.68150 normal 21 2010
## 22 114.70994 106.24725 123.17263 normal 22 2011
## 23 117.34259 105.98808 128.69710 normal 23 2012
## 24 120.12617 105.58485 134.66748 normal 24 2013
## 25 123.08377 105.09226 141.07528 normal 25 2014
## 26 126.21910 104.52257 147.91564 normal 26 2015
## 27 129.53854 103.87668 155.20040 normal 27 2016
## 28 133.05033 103.14841 162.95224 normal 28 2017
## 29 136.76349 102.32762 171.19936 normal 29 2018
## 30 140.68797 101.40149 179.97445 normal 30 2019
## 31 144.83465 100.35490 189.31440 normal 31 2020
```

In the next step we repeat this procedure for the model “no rank”.

```
data<-read.table("./RegressionCoefficientNORANK.txt", header = TRUE)

checkNodisease <- FALSE
t<-"pre-epidem"
period<-t
popsize0<-100
Tmax<-3
source('MC_simulations.R')
tabprepidemNR<-results(Mproj, NGMstoch, popvec, senslambda, theta)

t<-"epidem"
period<-t
Tmax<-3
popsize0<-tabprepidemNR[[2]][3]
source('MC_simulations.R')
tabepidemNR<-results(Mproj, NGMstoch, popvec, senslambda, theta)

t<-"post-epidem1"
period<-t
Tmax<-6
popsize0<-tabepidemNR[[2]][3]
source('MC_simulations.R')
tabpost1NR<-results(Mproj, NGMstoch, popvec, senslambda, theta)
```

```

t<-"post-epidem2"
Tmax<-12
period<-t
checkNodisease <- TRUE
popsize0<-tabpost1NR[[2]][6]
source('MC_simulations.R')

tabpost2NR<-results(Mproj, NGMstoch, popvec, senslambda, theta)

Tmax<-13
period<-t
checkNodisease <- TRUE
popsize0<-tabpost2NR[[2]][12]
source('MC_simulations.R')
tabprojectionNR<-results(Mproj, NGMstoch, popvec, senslambda, theta)

```

We store in a data frame the population vectors describing variations in expected abundance over 30 years, in both the models with and without rank effect.

```

lambdaseq <- c(rep(tabprepidemNR[[1]][1,1], 3), rep(tabepidemNR[[1]][1,1], 2) , rep(tabpost1NR[[1]][1,1], 1))

lambdase <- c(rep(tabprepidemNR[[1]][1,2], 3), rep(tabepidemNR[[1]][1,2], 2) , rep(tabpost1NR[[1]][1,2], 1))

popproj<-rep(0,31)
popproj[1]<-100
varmoins<-rep(0,31)
varplus<-rep(0,31)
varplus[1]<-100
varmoins[1]<-100

# loop on the retrospective (1990 to 2010)

for(t in 2:21) # starts at 2 because t-1 if t=1 is 0
{
  popproj[t]<-popproj[t-1] * lambdaseq[t]
  varplus[t]<-popproj[t-1] * (lambdaseq[t] + lambdase[t])
  varmoins[t]<-popproj[t-1] * (lambdaseq[t] - lambdase[t])
}

# Projections 2010-2020

p<-1
for(t in 22:33)
{
  p<-p+1 # to move to the next value
  popproj[t]<- tabprojectionNR[[2]][p]
  varplus[t]<- tabprojectionNR[[2]][p] + tabprojectionNR[[3]][p]
  varmoins[t]<- tabprojectionNR[[2]][p] - tabprojectionNR[[3]][p]
}

popproj<-popproj[-c(22,23)]

```



```
varplus<-varplus [-c(22,23)]
varmoins<-varmoins[-c(22,23)]
```

```
matNR <- cbind(popproj,varmoins,varplus)
matNR<-as.data.frame(matNR)
matNR$Model<-rep("NR", 31)
```

```
matNR$Time<-1:31
matNR$Time1 <-c(1990:2020)
```

```
matNEW <-rbind(mat2,matNR)
matNEW
```

##	popproj	varmoins	varplus	Model	Time	Time1
## 1	100.00000	100.00000	100.00000	normal	1	1990
## 2	103.59755	100.29873	106.89636	normal	2	1991
## 3	107.32451	103.90702	110.74200	normal	3	1992
## 4	101.73940	97.40575	106.07305	normal	4	1993
## 5	96.44493	92.33680	100.55306	normal	5	1994
## 6	95.15359	92.04800	98.25919	normal	6	1995
## 7	93.87954	90.81553	96.94355	normal	7	1996
## 8	92.62254	89.59956	95.64553	normal	8	1997
## 9	91.38238	88.39987	94.36489	normal	9	1998
## 10	90.15882	87.21625	93.10140	normal	10	1999
## 11	91.92250	89.36115	94.48384	normal	11	2000
## 12	93.72067	91.10922	96.33212	normal	12	2001
## 13	95.55402	92.89149	98.21656	normal	13	2002
## 14	97.42324	94.70862	100.13786	normal	14	2003
## 15	99.32902	96.56129	102.09674	normal	15	2004
## 16	101.27208	98.45021	104.09394	normal	16	2005
## 17	103.25315	100.37608	106.13021	normal	17	2006
## 18	105.27297	102.33962	108.20632	normal	18	2007
## 19	107.33230	104.34158	110.32303	normal	19	2008
## 20	109.43192	106.38269	112.48116	normal	20	2009
## 21	111.57261	108.46373	114.68150	normal	21	2010
## 22	114.70994	106.24725	123.17263	normal	22	2011
## 23	117.34259	105.98808	128.69710	normal	23	2012
## 24	120.12617	105.58485	134.66748	normal	24	2013
## 25	123.08377	105.09226	141.07528	normal	25	2014
## 26	126.21910	104.52257	147.91564	normal	26	2015
## 27	129.53854	103.87668	155.20040	normal	27	2016
## 28	133.05033	103.14841	162.95224	normal	28	2017
## 29	136.76349	102.32762	171.19936	normal	29	2018
## 30	140.68797	101.40149	179.97445	normal	30	2019
## 31	144.83465	100.35490	189.31440	normal	31	2020
## 32	100.00000	100.00000	100.00000	NR	1	1990
## 33	102.23532	99.57277	104.89786	NR	2	1991
## 34	104.52060	101.79853	107.24266	NR	3	1992
## 35	97.85228	94.16247	101.54209	NR	4	1993
## 36	91.60939	88.15499	95.06380	NR	5	1994
## 37	89.29581	86.88226	91.70936	NR	6	1995

## 38	87.04066	84.68806	89.39326	NR	7	1996
## 39	84.84246	82.54928	87.13564	NR	8	1997
## 40	82.69978	80.46451	84.93505	NR	9	1998
## 41	80.61121	78.43239	82.79002	NR	10	1999
## 42	81.47686	79.90866	83.04505	NR	11	2000
## 43	82.35180	80.76676	83.93683	NR	12	2001
## 44	83.23614	81.63408	84.83819	NR	13	2002
## 45	84.12998	82.51072	85.74923	NR	14	2003
## 46	85.03341	83.39676	86.67006	NR	15	2004
## 47	85.94655	84.29232	87.60077	NR	16	2005
## 48	86.86949	85.19750	88.54148	NR	17	2006
## 49	87.80234	86.11240	89.49228	NR	18	2007
## 50	88.74521	87.03712	90.45330	NR	19	2008
## 51	89.69821	87.97178	91.42464	NR	20	2009
## 52	90.66144	88.91647	92.40641	NR	21	2010
## 53	91.17895	86.54329	95.81461	NR	22	2011
## 54	92.06503	85.73942	98.39064	NR	23	2012
## 55	93.00552	84.88132	101.12971	NR	24	2013
## 56	94.00604	84.00318	104.00890	NR	25	2014
## 57	95.06625	83.11270	107.01980	NR	26	2015
## 58	96.18626	82.21230	110.16022	NR	27	2016
## 59	97.36664	81.30116	113.43212	NR	28	2017
## 60	98.60816	80.37686	116.83945	NR	29	2018
## 61	99.91175	79.43615	120.38735	NR	30	2019
## 62	101.27855	78.47526	124.08184	NR	31	2020

Then we plot Figure 5:

```
library(ggplot2)
```

```
## Warning: package 'ggplot2' was built under R version 3.5.1
```

```
p <- ggplot(data=matNEW, aes(x=Time1, y=popproj, ymin=varmoins, ymax=varplus)) +
```

```
  geom_rect(aes(xmin=1992, xmax=1994, ymin=-Inf, ymax=Inf), fill="#FFCC99",alpha=0.5) +
```

```
  geom_line(data=matNEW[matNEW$Model=="normal", ],aes(colour="myline1",x=Time1, y=popproj),size=1.5) +
  geom_ribbon(data=matNEW[matNEW$Model=="normal", ],alpha=0.7, color= "white",fill = "#CC79A7") + xlab("Year")
```

```
  geom_line(data=matNEW[matNEW$Model=="NR", ],aes(colour="myline2",x=Time1, y=popproj),size=1.5) +
  geom_ribbon(data=matNEW[matNEW$Model=="NR", ],alpha=0.7, color= "white",fill = "#56B4E9") + xlab("Year")
```

```
  scale_colour_manual(name="Model",values=c(myline1="#CC79A7", myline2="#56B4E9"),
    labels = c("COMPLETE MODEL", "NO RANK")) +
  ylim(50, 192.5) +
```

```
  scale_x_continuous(breaks = round(seq(min(matNEW$Time1), max(matNEW$Time1), by = 4),1))
```

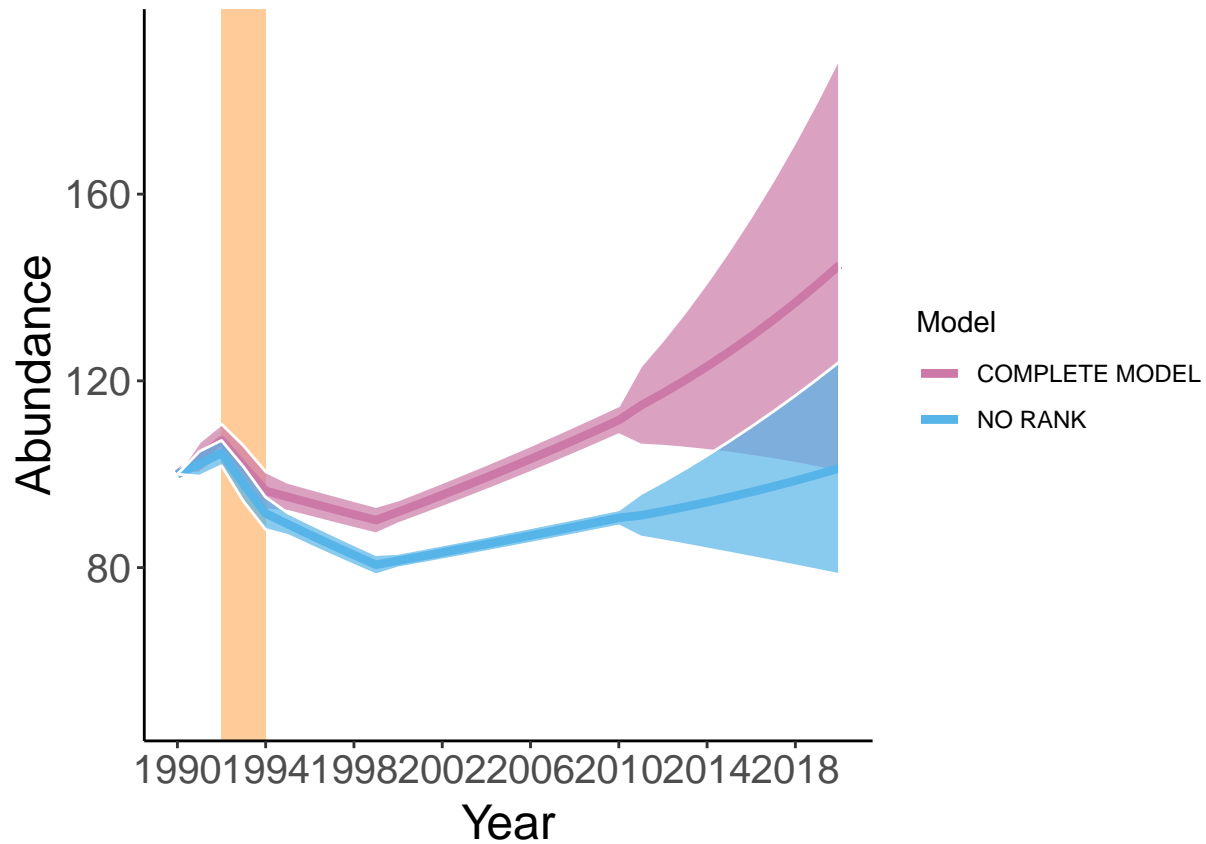
```
p + theme_bw() +
```

```
  theme(
    axis.line = element_line(colour = "black"),
    axis.text = element_text(size=15),
    panel.grid.major = element_blank(),
    panel.grid.minor = element_blank(),
```

```

panel.border = element_blank(),
panel.background = element_blank(),
axis.title.x = element_text(colour="black",size=18,angle=0,hjust=.5,vjust=0,face="plain"),
axis.title.y = element_text(colour="black",size=18),
plot.margin = unit(c(7.5, 12.5, 5.5, 5.5), "points")
)

```



## 2) Sensitivity analysis of the population's growth rate ( $\lambda$ )(Fig 2)

To determine which parameters contributed most to( $\lambda$ ) and predict the results of future changes in parameter estimates, we performed a sensitivity analysis. When elements of a population matrix are composed of several vital rates, the classical first order sensitivity analysis is not recommended, as it does not allow disentangling the effects of demographic, social and infection parameters. Therefore, we conducted lower-level sensitivity analyses for ( $\lambda$ ). In the source R file **MC\_simulations** we applied the function 'vitalsens' from the R package 'popbio' which evaluate the expression of the matrix projection using all parameter values of the MonteCarlo iterations.

```

figtab<-list((tabprepidem[[8]]), (tabepidem[[8]]), (tabpost1[[8]])) # removes the character names and
names(figtab)<-c("sens_preepidem", "sens_epidem", "sens_postepidem1")

colnames(figtab[[1]])<-names(tabepidem[[5]])
colnames(figtab[[2]])<-names(tabepidem[[5]])
colnames(figtab[[3]])<-names(tabepidem[[5]])

meanfigtab<-rbind(colMeans(tabprepidem[[8]]), colMeans(tabepidem[[8]]), colMeans(tabpost1[[8]]))

```

```

row.names(meanfigtab)<-c("sens_preepidem", "sens_epidem", "sens_postepidem1")
colnames(meanfigtab)<-names(tabepidem[[5]])

meanfigtab<-data.frame(t(meanfigtab))

# 2.a) Order the preepidem table by decreasing importance
sens1<-subset(meanfigtab, abs(meanfigtab$sens_preepidem) >= 0.10)
sensN1<-figtab[[1]][, rownames(sens1)[order(abs(sens1[,1]),decreasing=TRUE)]] # Order from highest im

# to check the sequence of parameters later included in the "titre":
sens1_order <- sens1[order(abs(sens1[,1]),decreasing =TRUE),]

# 2.b) Order the epidem table by decreasing importance
sens2<-subset(meanfigtab, abs(meanfigtab$sens_epidem) >= 0.10)
sensN2<-figtab[[2]][, rownames(sens2)[order(abs(sens2[,2]),decreasing=TRUE)]] # Order from highest im

sens2_order <- sens2[order(abs(sens2[,2]),decreasing =TRUE),]

# 2.c) Order the post-epidem table by decreasing importance
sens3<-subset(meanfigtab, abs(meanfigtab$sens_postepidem) >= 0.10)
sensN3<-figtab[[3]][, rownames(sens3)[order(abs(sens3[,3]),decreasing=TRUE)]] # Order from highest im

sens3_order <- sens3[order(abs(sens3[,3]),decreasing =TRUE),]

# prepare the labels and colors
titre<-c(expression(psi["BH"]), expression(psi["NBH"]), expression(psi["SAH"]), expression(r["H"]), e

titre2<-c(expression(psi["BH"]), expression(psi["NBH"]), expression(phi["NBHR"]), expression(r["H"]))

titre3<-c(expression(psi["BH"]), expression(psi["NBH"]), expression(phi["NBHR"]), expression(r["H"]))

color = rep(NA, length=ncol(sensN1))

color[which(colnames(sensN1)== "bSH")] = "#CC79A7"
color[which(colnames(sensN1)== "bBH")] = "#CC79A7"
color[which(colnames(sensN1)== "bNBH")] = "#CC79A7"

color[which(colnames(sensN1)== "bSL")] = "#F0E442"
color[which(colnames(sensN1)== "bBL")] = "#F0E442"
color[which(colnames(sensN1)== "bNBL")] = "#F0E442"

color[which(colnames(sensN1)== "rHH.R")] = "#CC79A7"
color[which(colnames(sensN1)== "rLL.R")] = "#F0E442"

color[which(colnames(sensN1)== "phiBHR")] = "#CC79A7"
color[which(colnames(sensN1)== "phiNBHR")] = "#CC79A7"

color[which(colnames(sensN1)== "phiNBLR")] = "#F0E442"

```

```

color2 = rep(NA, length=ncol(sensN2))

color2[which(colnames(sensN2)=="bSH")] = "#CC79A7"
color2[which(colnames(sensN2)=="bBH")] = "#CC79A7"
color2[which(colnames(sensN2)=="bNBH")] = "#CC79A7"

color2[which(colnames(sensN2)=="bSL")] = "#F0E442"
color2[which(colnames(sensN2)=="bBL")] = "#F0E442"
color2[which(colnames(sensN2)=="bNBL")] = "#F0E442"

color2[which(colnames(sensN2)=="rHH.R")] = "#CC79A7"
color2[which(colnames(sensN2)=="rLL.R")] = "#F0E442"

color2[which(colnames(sensN2)=="phiBHR")] = "#CC79A7"
color2[which(colnames(sensN2)=="phiNBHR")] = "#CC79A7"

color2[which(colnames(sensN2)=="phiBLR")] = "#F0E442"
color2[which(colnames(sensN2)=="phiNBLR")] = "#F0E442"

color3 = rep(NA, length=ncol(sensN3))

color3[which(colnames(sensN3)=="bSH")] = "#CC79A7"
color3[which(colnames(sensN3)=="bBH")] = "#CC79A7"
color3[which(colnames(sensN3)=="bNBH")] = "#CC79A7"

color3[which(colnames(sensN3)=="bSL")] = "#F0E442"
color3[which(colnames(sensN3)=="bBL")] = "#F0E442"
color3[which(colnames(sensN3)=="bNBL")] = "#F0E442"

color3[which(colnames(sensN3)=="rHH.R")] = "#CC79A7"
color3[which(colnames(sensN3)=="rLL.R")] = "#F0E442"

color3[which(colnames(sensN3)=="phiBHR")] = "#CC79A7"
color3[which(colnames(sensN3)=="phiNBHR")] = "#CC79A7"

color3[which(colnames(sensN3)=="phiBLR")] = "#F0E442"
color3[which(colnames(sensN3)=="phiNBLR")] = "#F0E442"

# pdf("Figure2SensLambdaNew.pdf",width=7,height=5) #uncomment this line to get the figure in pdf

nf <- layout(matrix(c(0,1,0,0,2,0,0,3,0),3, byrow = TRUE),
               widths= c(1.2,2,1.2), heights=c(2,2,2)) # widths for columns,heights for rows
#layout.show(nf)

par(mar =c(3.3,3,2,0))

# x1 and y1 to draw the axes

```

```

x1 <- c(-0.1:13)
y1 <- c(-0.82,-0.50,-0.25,0,0.25,0.50,0.75,1)

# x2 and y2 to add labels and tickmarks
x2 <- c(1:11)
y2 <- c(-0.50,0,0.5,1)

# pre-epidemic period:

boxplot(sensN1, col=color, ylim=c(-0.75,1), axes=F)
mtext(side = 3,text=expression(paste(bold("a"))),adj=0, line =0.4,cex = 1.2)

axis(1,at=x1,las=1,tck=0, labels=NA)
axis(2,at=y1,las=1,tck=0, labels =NA)

axis(1,at=x2,las=1,cex.axis=0.85, labels=titre)
axis(2,at=y2,las=1,cex.axis=1.4)

mtext(side = 2, text =expression(paste("Sensitivity ", lambda)), line = 2.9, cex=1.3)

abline(h=0, lty=2)

legend(8,1.5, legend=c("high-ranking", "low-ranking"),
fill=c("#CC79A7", "#F0E442"), cex=0.8, xpd = T)

# epidemic period
x2 <- c(1:12)

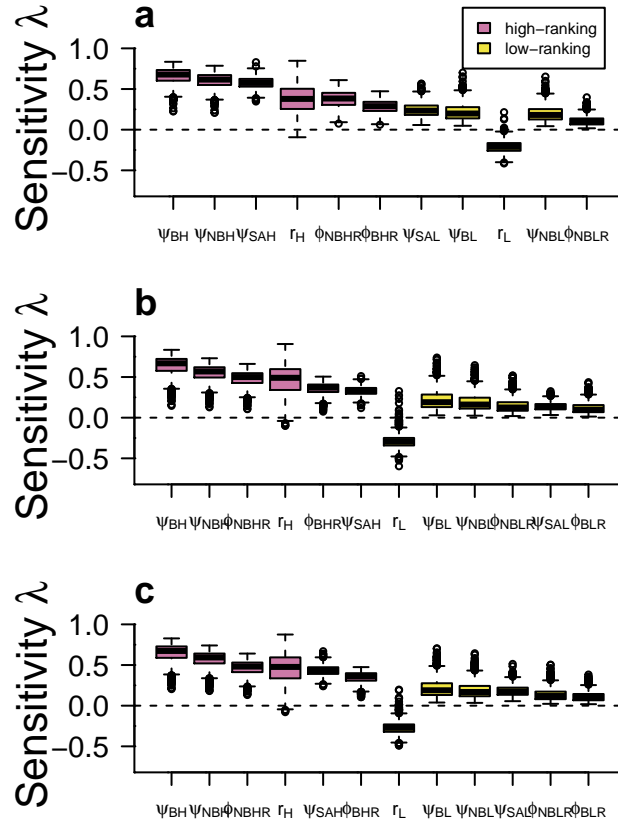
boxplot(sensN2, axes=F, col=color2, ylim=c(-0.75,1))
mtext(side = 3,text=expression(paste(bold("b"))),adj=0, line =0.4,cex = 1.2)
axis(1,at=x1,las=1,tck=0, labels=NA)
axis(2,at=y1,las=1,tck=0, labels =NA)
axis(1,at=x2,las=1,cex.axis=0.85, labels=titre2)
axis(2,at=y2,las=1,cex.axis=1.4)
mtext(side = 2, text =expression(paste("Sensitivity ", lambda)), line = 2.9, cex=1.3)
abline(h=0, lty=2)

# post-epidemic period

boxplot(sensN3, axes=F, col=color3, ylim=c(-0.75,1))
mtext(side = 3,text=expression(paste(bold("c"))),adj=0, line =0.4,cex = 1.2)

axis(1,at=x1,las=1,tck=0, labels=NA)
axis(2,at=y1,las=1,tck=0, labels =NA)
axis(1,at=x2,las=1,cex.axis=0.85, labels=titre3)
axis(2,at=y2,las=1,cex.axis=1.4)
mtext(side = 2, text =expression(paste("Sensitivity ", lambda)), line = 2.9, cex=1.3)
abline(h=0, lty=2)

```



```
# dev.off() # uncomment this line if you want to get only the figure in pdf
```

### 3) Sensitivity analysis of $R_0$ (Fig 3)

Here we determine which parameters contributed most to variation in  $R_0$  in order to predict future disease dynamics with changes in parameter ( $\theta$ ). Because we did not have a symbolic expression of the next generation matrix, we could not apply the vitalsens function as we did for calculating the sensitivity of lambda. Instead, we performed the sensitivity analysis of  $R_0$  by coding the following equation  $\delta(R_0)/\delta(\theta)$ .

Now we plot the sensitivity values of  $R_0$ :

```
fulltab<-matrix(0, nrow=MCiter, ncol=23)
colnames(fulltab)<-1:23
for(i in 1:23)
{
  fulltab[,i]<-tabepidem[[10]][[i]]$sensR0
  colnames(fulltab)[i]<-tabepidem[[10]][[i]]$param
}
```

```
fulltab<-as.data.frame(fulltab)
```

```
# 2) Order by decreasing importance
```

```
# here we first extract the mean values of the sensitivity to  $R_0$ , we select those which absolute value
```

```

figtab<-cbind(as.numeric(tabepidem[[9]][,2]), as.numeric(tabepidem[[9]][,3])) # extract the mean sens

colnames(figtab)<-c("sens_epidem","sd_epidem")

figtab<-as.data.frame(figtab)

figtab$sens_epidem[12:15]<- 1 - figtab$sens_epidem[12:15]

figtab$param<- as.character(tabepidem[[9]][,1])

#----- Epidem
# 1) Select mean sensitivity values whose abs. value is higher than 10%
sensNew<-subset(figtab, abs(figtab$sens_epidem) >= 0.10)

# 2) Order by decreasing importance
sensNew1<-fulltab[order(abs(sensNew$sens_epidem),decreasing=TRUE),]

sens<-NULL
for(i in 1:20){
sens<-c(sens,which(colnames(fulltab) == sensNew$param[order(abs(sensNew$sens_epidem),decreasing=TRUE)]))

sensN2<-fulltab[,sens] #
# SENSITIVITY TO INFECTION (1 - sensitivity to the transition from Susceptible to Susceptible)
sensN2[,3]<-1-sensN2[,3]
sensN2[,2]<-1-sensN2[,2]
sensN2[,8]<-1-sensN2[,8]
sensN2[,9]<-1-sensN2[,9]

titreR0<-c("Breeding_BH", "Infection_SA&B&NBH", "Infection_SA&B&NBL", "Breeding_NBH", "Staying_H", "S

titrefig<-c (expression(psi["BH"]), expression(beta["H"]), expression(beta["L"]), expression(psi["NBH"]
expression(sr),expression(phi["SAIR"]))

colnames(sensN2)<-titreR0
# -- Plot; decreasing importance

par(mar=c(5.1,5.1,4.1,2.1))

color = rep(NA, length=length(sensNew1$param))

# Survival
# cubs
color[which(colnames(sensN2)=="Survival_CHS")] = "#CC79A7"

```



```

color[which(colnames(sensN2)=="Survival_CLS")] = "#F0E442"

# subadults
color[which(colnames(sensN2)=="Survival_SAHS")] = "#CC79A7"
color[which(colnames(sensN2)=="Survival_SAI&R")] = "#999999"

color[which(colnames(sensN2)=="Survival_SALS")] = "#F0E442"

#breeders
color[which(colnames(sensN2)=="Survival_B")] = "#999999"

#Nonbreeders
color[which(colnames(sensN2)=="Survival_NB")] = "#999999"

#rank transitions
color[which(colnames(sensN2)=="Staying_H")] = "#CC79A7"
color[which(colnames(sensN2)=="Staying_L")] = "#F0E442"

# Infection proba
color[which(colnames(sensN2)=="Infection_CH")] = "#CC79A7"
color[which(colnames(sensN2)=="Infection_CL")] = "#F0E442"

color[which(colnames(sensN2)=="Infection_SA&B&NBH")] = "#CC79A7"
color[which(colnames(sensN2)=="Infection_SA&B&NBL")] = "#F0E442"

# Breeding
color[which(colnames(sensN2)=="Breeding_BH")] = "#CC79A7"
color[which(colnames(sensN2)=="Breeding_NBH")] = "#CC79A7"
color[which(colnames(sensN2)=="Breeding_SAH")] = "#CC79A7"

color[which(colnames(sensN2)=="Breeding_BL")] = "#F0E442"
color[which(colnames(sensN2)=="Breeding_NBL")] = "#F0E442"
color[which(colnames(sensN2)=="Breeding_SAL")] = "#F0E442"

#Sex ratio
color[which(colnames(sensN2)=="SexRatio")] = "#999999"

# pdf("SensitivityRO.pdf",width=7,height=5)

colnames(sensN2)<-NULL
boxplot(sensN2, col=color, ylim=c(-7,5), axes=F)

# x1 and y1 to draw the axes
x1 <- c(-0.3:20)
y1 <- c(-8,5)

# x2 and y2 to add labels and tickmarks
x2 <- c(1:20)
y2 <- c(-6,-4,-2,0,2,4)

```

```

axis(1,at=x1,las=1,tck=0, labels=NA)
axis(2,at=y1,las=1,tck=0, labels =NA)

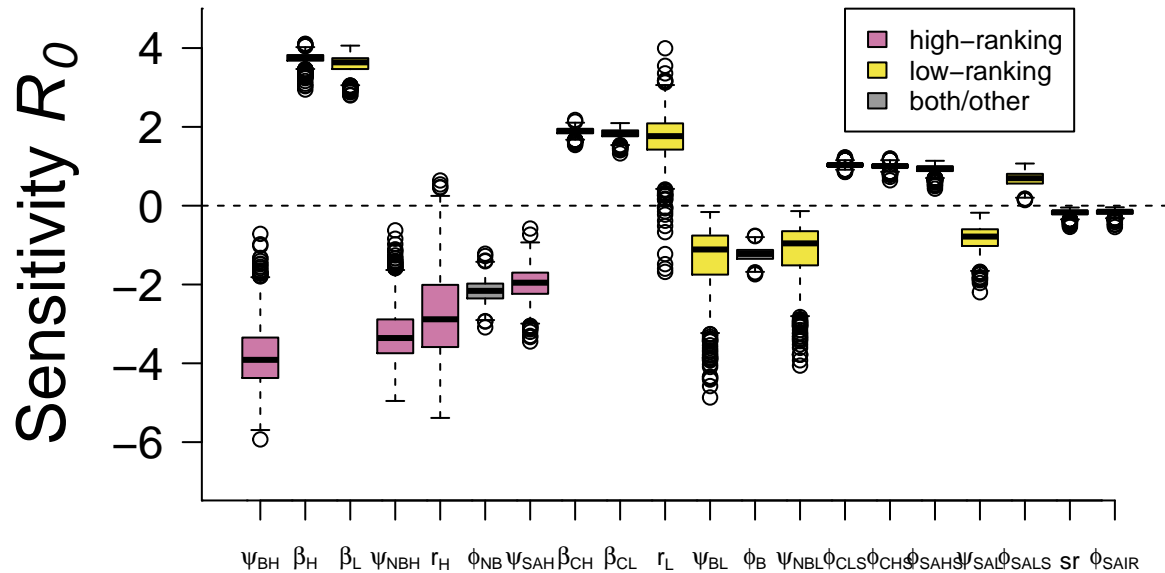
axis(1,at=x2,las=1,cex.axis=0.75, labels=titrefig)
axis(2,at=y2,las=1,cex.axis=1.4)

abline(h = 0, lty =2)

mtext(side = 2, text =expression(paste("Sensitivity ", italic(R["0"]))), line = 3, cex=2.1)

legend(14,5, legend=c("high-ranking", "low-ranking", "both/other"),
fill=c("#CC79A7", "#F0E442", "#999999"), cex=0.8)

```



```

#dev.off()

end_time <- Sys.time()
run_time <-end_time-start_time
round(run_time,digits = 2)

## Time difference of 11.59 hours

```

## References (R packages)

Stubben, C. & Milligan, B. Estimating and analyzing demographic models using the popbio package in R. *J. Stat. Softw* 22, 1-23 (2007).

Stott, I., Hodgson, D. J. & Townley, S. popdemo: an R package for population demography using projection matrix analysis. *Methods Ecol. Evol.* 3, 797-802 (2012).