## Prior\_production\_IM

karine Durand

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## production des priors IM

## mon\_script

```
library(stats)
library("KScorrect", lib.loc="~/R/x86_64-pc-linux-gnu-library/3.3")
#####partie locus
#variables locus
#-L=taille du gène
\#-t=theta
\#-r=rho
#-delta=taille du track recombinant
#boucle de 10 000 iterations(10 000 tirages demographique)
demo<-NULL
locus<-NULL
tbs<-NULL
#####TIRER un prior locus dans une distribution uniforme de bornes
L<-scan("/home/kadurand/partage_windows/Xylella/analyses_genomiques/ABC/msms/lenght")#bound_taille du g
t<-runif(997,1E-6, 0.01) #bound_theta=[0-0.0003] bornes vrai pour 13pauca_multiplex augmenter la borne s
r<-runif(997,0,0.01)#bound_rho=[0-0.0003]bornes vrai pour 13pauca_multiplex augmenter la borne sup à 0.
\#delta < -round(runif(997, 10, 1000))\#bound = [10-1000]
\#print(L, t, r, delta)
m_locus=matrix(c(L,t*L,r*L),ncol=3)
m_locus=as.data.frame(m_locus)
for (i in 1:1000){#tirage des priors demographiques
  #variables demographique modéle SI
  ##Param_demo (5) = Ts N1, N2, M12, M21
  Ts<-rlunif(1,1E+4,1E+7)#bound=[1,100,1E+7]de 100ans a 100 000 ans
  N1 < -rlunif(1,100,1E+6) \#bound = [100,1E+6]
  N2 < -rlunif(1,100,1E+6) \#bound = [100,1E+6]
  Na<-rlunif(1,100,1E+6)#Bound=[100,1E+6]
  M12 < -runif(1,0.01,30) \#bound = [0.01-30]
  M21 < -runif(1,0.01,30) \#bound = [0.01-30]
  #print( Ts N1, N2, M12, M21)
m_demo=matrix(c(Ts,N1,N2,Na,M12,M21),ncol=6)
m_demo=as.data.frame(m_demo)
locus<-cbind(m locus,m demo)</pre>
path <- "/home//kadurand/partage_windows/Xylella/analyses_genomiques/ABC/msms/Priors_IM_10000_mod/IM"
write.table(locus,file= paste(path,i, sep="-"),col.names=FALSE,row.names =FALSE)
```