## 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/1368oRTHOLOGUES\_summarystats/locality and the summary states and the summary states are summary states are summary states are summary states and the summary states are summary states are summary states are summary states and the summary states are summary states and the summary states are summary states are summary states and the summary states are summa
t<-runif(1,1E-11, 0.001) #bound_theta=[0-0.0003] bornes vrai pour 13pauca_multiplex augmenter la borne s
r<-runif(1,0,0.0001)#bound_theta=[0-0.0003]bornes vrai pour 13pauca_multiplex augmenter la borne sup à
delta<-round(runif(1,10, 1000))#bound=[10-1000]
\#print(L, t, r, delta)
m_locus=matrix(c(t,r,delta),ncol=3)
m_locus=as.data.frame(m_locus)
for (i in 1:10000){#tirage des priors demographiques
    #variables demographique modéle SI
    ##Param demo (5) = Ts N1, N2, M12, M21
    Ts<-rlunif(1,100,1E+8)#bound=[1,100,1E+8]
    \#N1 < -rlunif(1, 100, 1E+6) \#bound = [100, 1E+6]
    N2 < -rlunif(1, 1E-3, 1E+3) \#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]
    Tsc < -rlunif(1,10,Ts) \#bound = [0-100]borne sup < Ts
    TS2=Ts+(Ts/1E+6)
    #print( Ts N1, N2, M12, M21)
m_demo=matrix(c(Ts,N2,Na,M12,M21,TS2),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/fastSimBac_linux/Priors_IM_100
write.table(locus,file= paste(path,i, sep="-"),col.names=FALSE,row.names =FALSE)
```