Prior_production_IM

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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-11, 0.001 )#bound_theta=[0-0.0003]bornes vrai pour 13pauca_multiplex augmenter la borne
r<-runif(997,0,0.0001)#bound_theta=[0-0.0003]bornes vrai pour 13pauca_multiplex augmenter la borne sup
delta<-round(runif(997,10, 1000))#bound=[10-1000]
#print(L,t,r,delta)
m_locus=matrix(c(L,t,r,delta),ncol=4)
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(1,100,1E+5)#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)

path <- "/home//kadurand/partage_windows/Xylella/analyses_genomiques/ABC/msms/Priors_IM_10000/IM"
write.table(locus,file= paste(path,i, sep="-"),col.names=FALSE,row.names =FALSE)
}
```