## Priors\_production

karine Durand
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## production des priors SI

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mon_script
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library(stats)
library("KScorrect", lib.loc="~/R/x86_64-pc-linux-gnu-library/3.3")
#####partie locus
#variables locus
\#-L=taille\ du\ gene
\#-t=theta
\#-r=rho
#-delta=taille du track recombinant
#boucle de 1000000 iterations(1000000 tirage 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 for the state of the
L<-scan("/home/kadurand/partage_windows/Xylella/analyses_genomiques/ABC/msms/lenght")#bound_taille du g
t<-runif(997,0, 0.004) #bound_theta=[0-0.0003]bornes vrai pour 13pauca_multiplex augmenter la borne sup
r<-runif(997,0,0.004)#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 (3) = Ts, N1, N2,
    ##Param demo (5) = Ts N1, N2, M12, M21
    Ts<-runif(1,0,20)#bound=[1,100,1E+7] Ts/4N0
    N1 < -runif(1,0,30) \#bound = [100,1E+6] X=N1/N0
    N2 < -runif(1,0,30) \#bound = [100,1E+6]
    Na < -runif(1,0,60) \#Bound = [100,1E+6]
#print( N1, N2, Ts)
m_demo=matrix(c(Ts,N1,N2,Na),ncol=4)
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_SI_ms_13/SI"</pre>
write.table(locus,file= paste(path,i, sep="-"),col.names=FALSE,row.names =FALSE)
```