

Song Variability Analysis

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R Markdown

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Here we are going to analyze patterns of variability in song structure among zebra finch populations. We are going to focus on an analysis of a summary statistic, PC1, derived from the log transformed raw K-L distances. We begin by testing for variation among two domesticated populations of *T. guttata castanotis* (TGC) [ECU and Chicago], one wild-derived population of TGC (Macquarie University) and one population of *T. guttata guttata* (TGG; ECU).

```
path<-("~/Documents/Projects/Domestication/OpenBiology/PopulationEffect"
dat=read.table(paste(path,"logKL_4pops_PCA.txt",sep="/"),sep = "\t",header=TRUE)
str(dat)
```

```

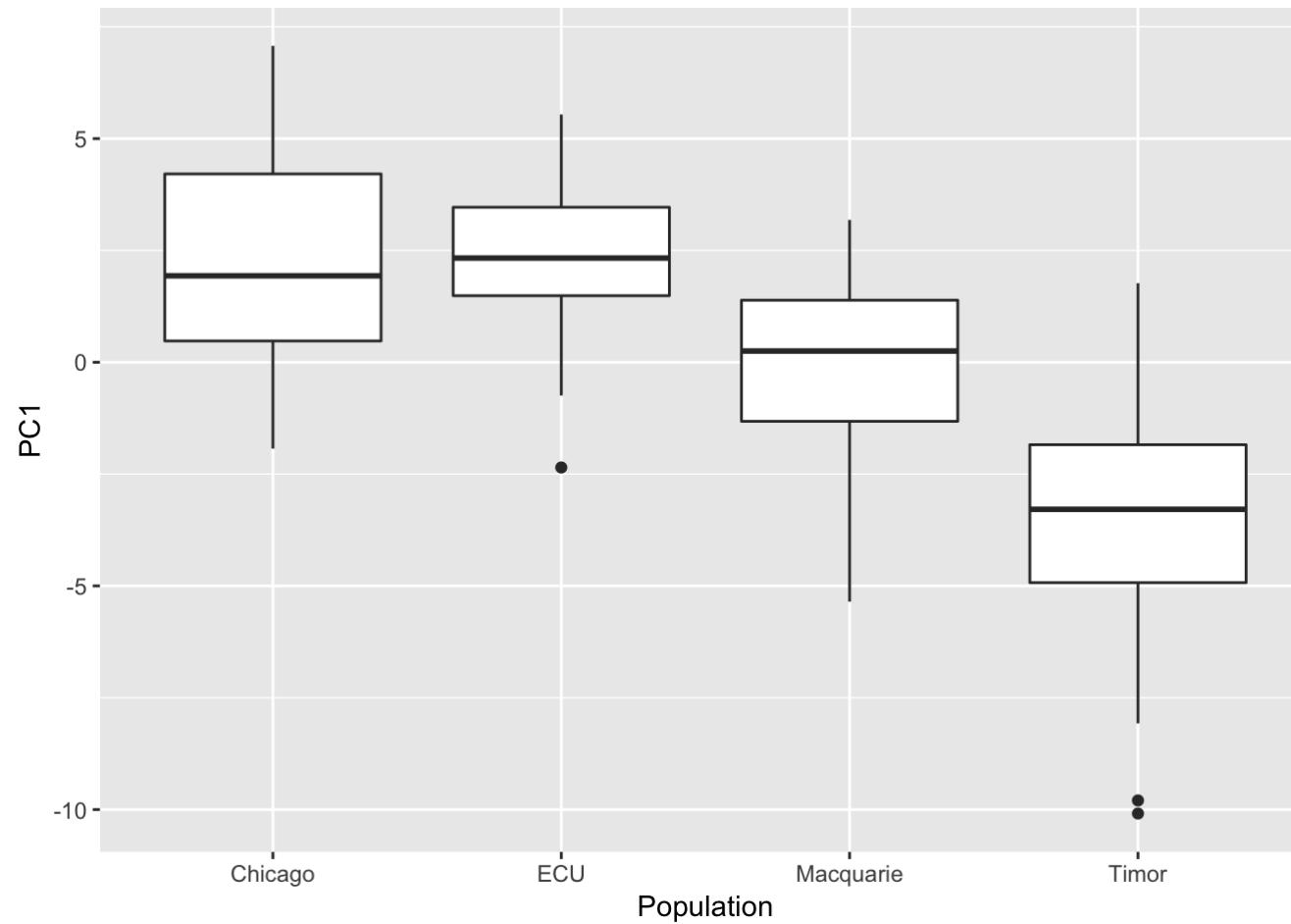
## 'data.frame': 278 obs. of 29 variables:
## $ X           : int 33 34 35 36 37 38 39 40 41 42 ...
## $ logamplitude : num 0.0731 0.3528 0.7059 0.3134 0.8614 ...
## $ logpitch    : num -0.0401 -0.1553 -0.1624 0.0384 0.0264 ...
## $ logFM        : num -0.05112 -0.04725 -0.10572 0.00904 0.20914 ...
## $ logAM2       : num -0.0501 0.0308 0.0118 0.0678 0.2659 ...
## $ logentropy   : num -0.1463 -0.039 -0.1209 0.0593 -0.0146 ...
## $ logpitchgoodness : num 0.1776 0.0858 -0.1597 0.1583 0.2722 ...
## $ logmeanfreq  : num 0.0247 -0.0111 -0.0465 0.1968 0.0784 ...
## $ logvpitch    : num -0.284 -0.644 -0.355 -0.212 -0.429 ...
## $ logvFM       : num -0.109 -0.305 -0.285 -0.15 -0.19 ...
## $ logventropy  : num -0.1256 -0.323 -0.2192 -0.1177 -0.0522 ...
## $ logvpitchgoodness: num 0.1502 -0.1482 -0.0221 0.1015 0.4843 ...
## $ vmeanfreq    : num -0.135 -0.119 -0.17 -0.112 -0.189 ...
## $ PC1          : num -4.42 -5.08 -4.86 -3.62 -2.74 ...
## $ PC1neg       : num 4.42 5.08 4.86 3.62 2.74 ...
## $ PC2          : num 0.473 -0.739 -1.706 -0.138 -1.931 ...
## $ PC3          : num -0.477 -0.42 0.486 -0.172 -0.842 ...
## $ PC4          : num -0.1379 -0.227 -0.0764 0.2167 -0.637 ...
## $ PC5          : num 0.839 0.0795 0.5432 0.5411 0.7339 ...
## $ PC6          : num -0.0179 -0.1383 -0.232 0.1277 0.2246 ...
## $ PC7          : num -0.4882 0.0517 -0.2967 0.05 -0.5216 ...
## $ PC8          : num -0.4904 0.2227 -0.0157 -0.1431 0.1708 ...
## $ PC9          : num -0.1904 -0.4548 -0.0159 -0.0331 0.2913 ...
## $ PC10         : num -0.346 -0.217 -0.359 -0.372 -0.802 ...
## $ PC11         : num 0.0963 -0.0212 0.1071 -0.0505 -0.2981 ...
## $ PC12         : num 0.000142 0.322964 -0.303615 0.046641 -0.440604 ...
## $ Population   : Factor w/ 4 levels "Chicago", "ECU", ...: 4 4 4 4 4 4 4 4 4 ...
## $ Template     : Factor w/ 35 levels "filtered_BL65_tclust", ...: 1 1 1 1 1 1 1 1 1 2 ...
## $ Target       : Factor w/ 35 levels "filtered_BL65_tclust", ...: 2 3 4 22 23 24 25 26 27 1 ...

```

```
require(ggplot2)
```

```
## Loading required package: ggplot2
```

```
print(ggplot(dat, aes(x=Population, y=PC1)) + geom_boxplot())
```



Since KL-distance is a pairwise measurement between two individual birds, we need to deal with the non-independence inherent in the data. To do this, we will analyze the data using generalized linear mixed models. Each KL measurement includes a “template” and a “target” and we will treat each of these as nested random effects in the model.

We've opted to use bayesian mixed models to avoid singularity issues we were having with standard lmer models.

```
require(lme4)
## Loading required package: lme4
## Loading required package: Matrix
```

```
library(blme)

mPC1 <- blmer(PC1 ~ Population + (1 | Population/Template) + (1 | Population/Target), data = dat, REML = FALSE)
summary(mPC1)
```

```

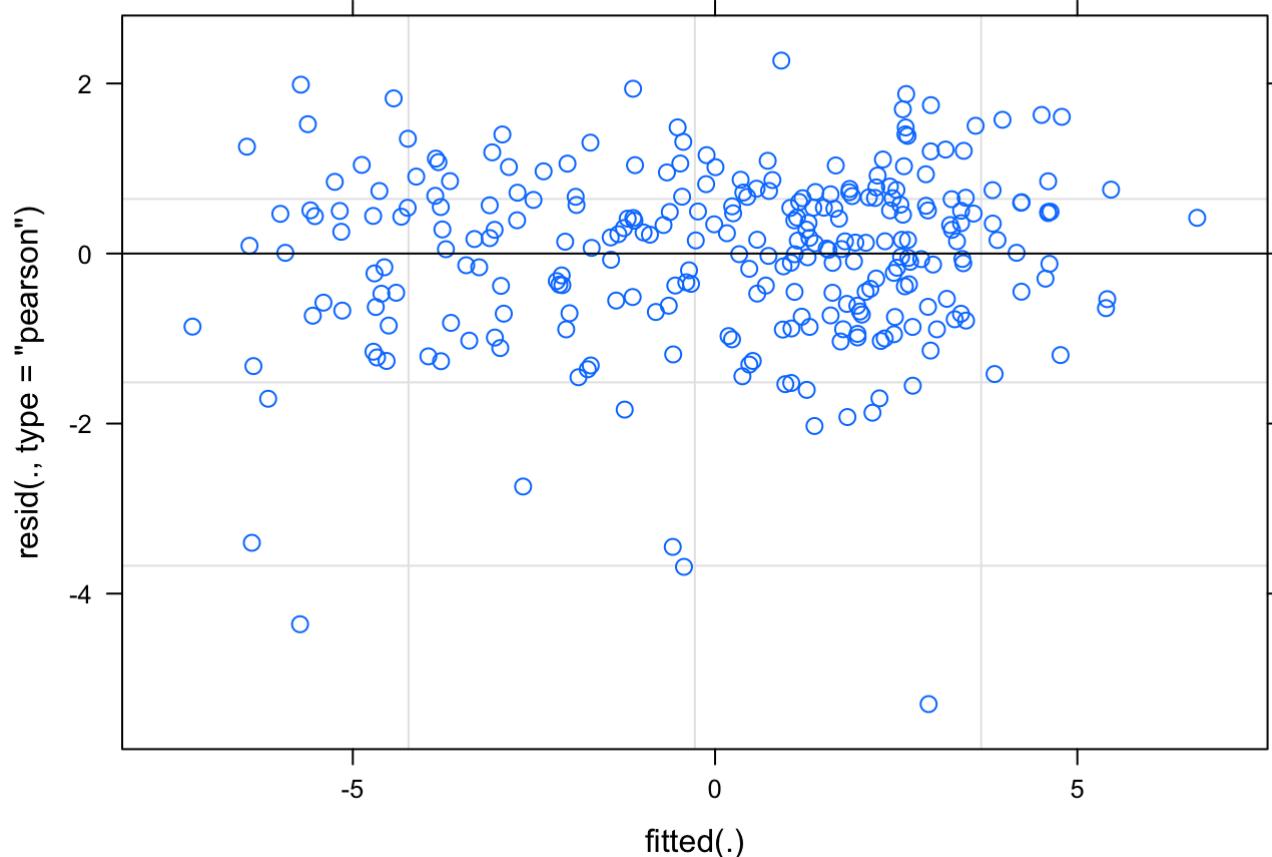
## Cov prior : Target:Population ~ wishart(df = 3.5, scale = Inf, posterior.scale = cov, common.scale = TRUE)
##           : Template:Population ~ wishart(df = 3.5, scale = Inf, posterior.scale = cov, common.scale = TRUE)
##           : Population ~ wishart(df = 3.5, scale = Inf, posterior.scale = cov, common.scale = TRUE)
## Prior dev : 1.0421
##
## Linear mixed model fit by maximum likelihood  ['blmerMod']
## Formula:
## PC1 ~ Population + (1 | Population/Template) + (1 | Population/Target)
##   Data: dat
##
##      AIC      BIC      logLik deviance df.resid
## 1050.3 1083.0   -516.2    1032.3       269
##
## Scaled residuals:
##    Min     1Q Median     3Q    Max
## -4.6529 -0.5387  0.1348  0.5755  1.9925
##
## Random effects:
## Groups            Name        Variance Std.Dev.
## Target:Population (Intercept) 1.5569   1.248
## Template:Population (Intercept) 2.0674   1.438
## Population         (Intercept) 0.6626   0.814
## Population.1       (Intercept) 0.6626   0.814
## Residual           1.2971   1.139
## Number of obs: 278, groups:
## Target:Population, 35; Template:Population, 35; Population, 4
##
## Fixed effects:
##             Estimate Std. Error t value
## (Intercept) 2.2516    1.3421  1.678
## PopulationECU 0.1097    1.8717  0.059
## PopulationMacquarie -2.4293   1.9171 -1.267
## PopulationTimor -5.5421   1.8717 -2.961
##
## Correlation of Fixed Effects:
##          (Intr) PplECU PpltnM
## PopulatnECU -0.717
## PopultnMcqr -0.700  0.502
## PopulatnTmr -0.717  0.514  0.502

```

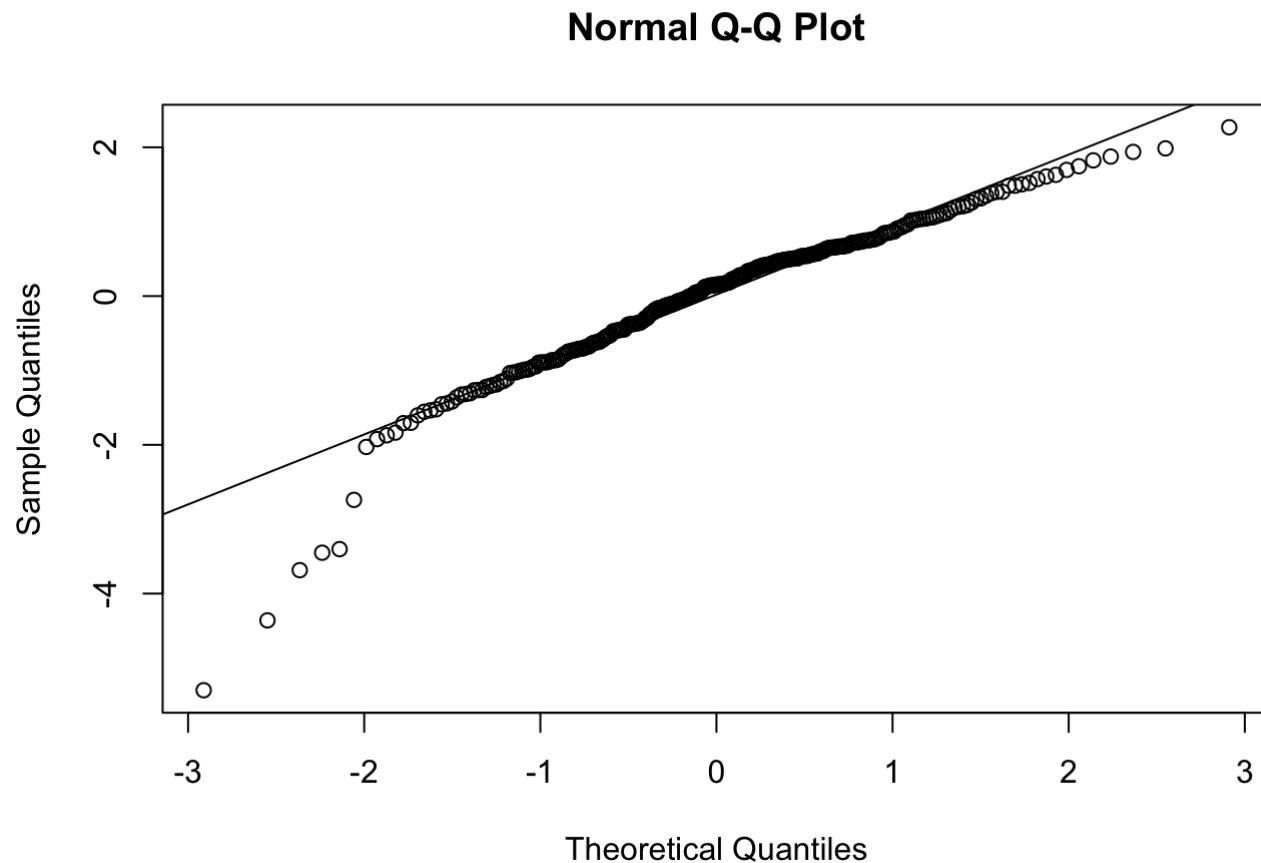
```
VarCorr(mPC1) # Check for singularity
```

```
## Groups           Name      Std.Dev.  
## Target.Population (Intercept) 1.24777  
## Template.Population (Intercept) 1.43786  
## Population        (Intercept) 0.81398  
## Population.1       (Intercept) 0.81398  
## Residual                         1.13889
```

```
plot(mPC1) # residuals plot
```



```
qqnorm(resid(mPC1)); qqline(resid(mPC1)) #qqplot
```



```
#check blmer
```

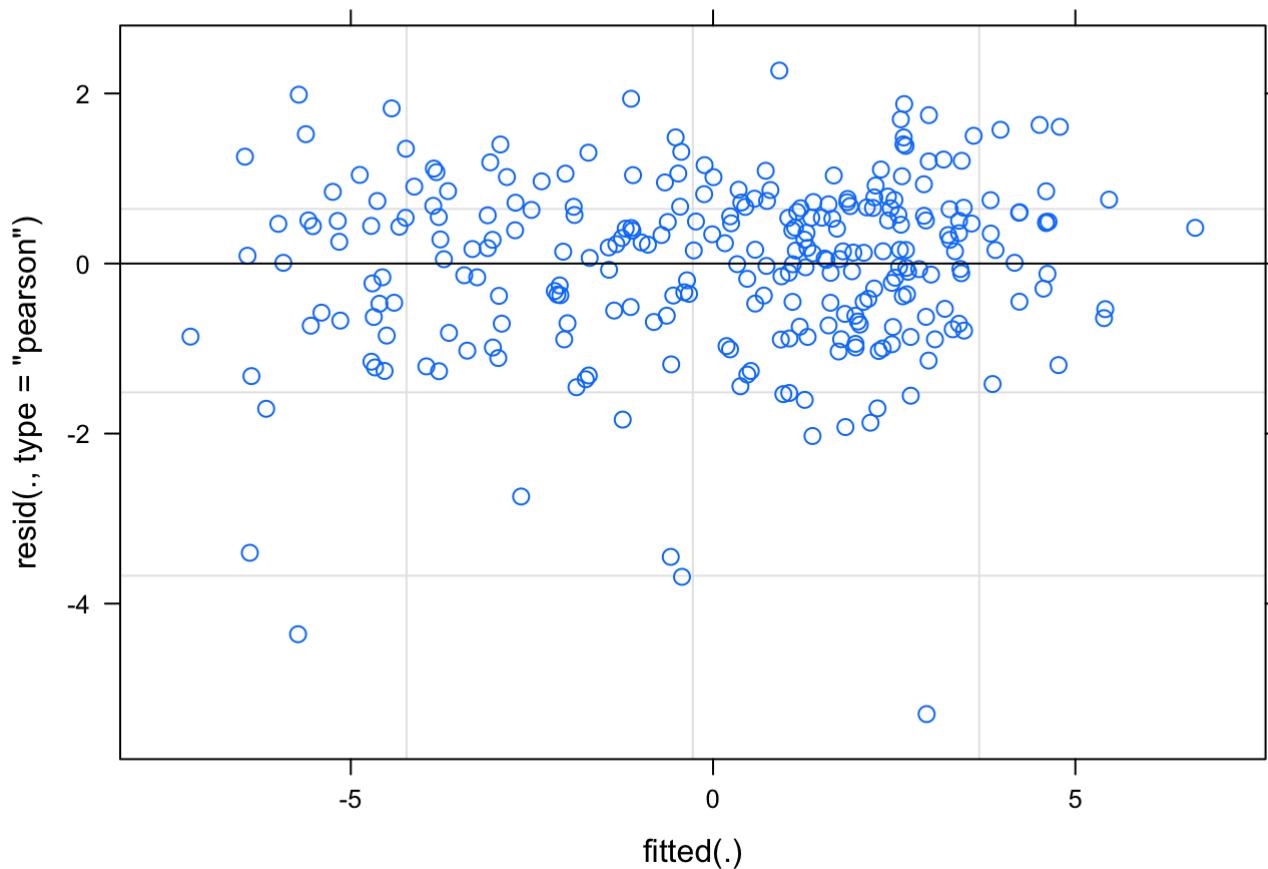
We can see above that we don't have an O estimates on our error terms. The residuals plot also looks pretty good as does the q-q plot. The lower tail isn't perfect, but overall this is not too bad.

So now let us fit a null model, removing Population as a fixed factor.

```
mNull <- blmer(PC1 ~ 1 + (1 | Population/Template) + (1 | Population/Target), data = dat, REML = FALSE)
VarCorr(mPC1) # Check for singularity
```

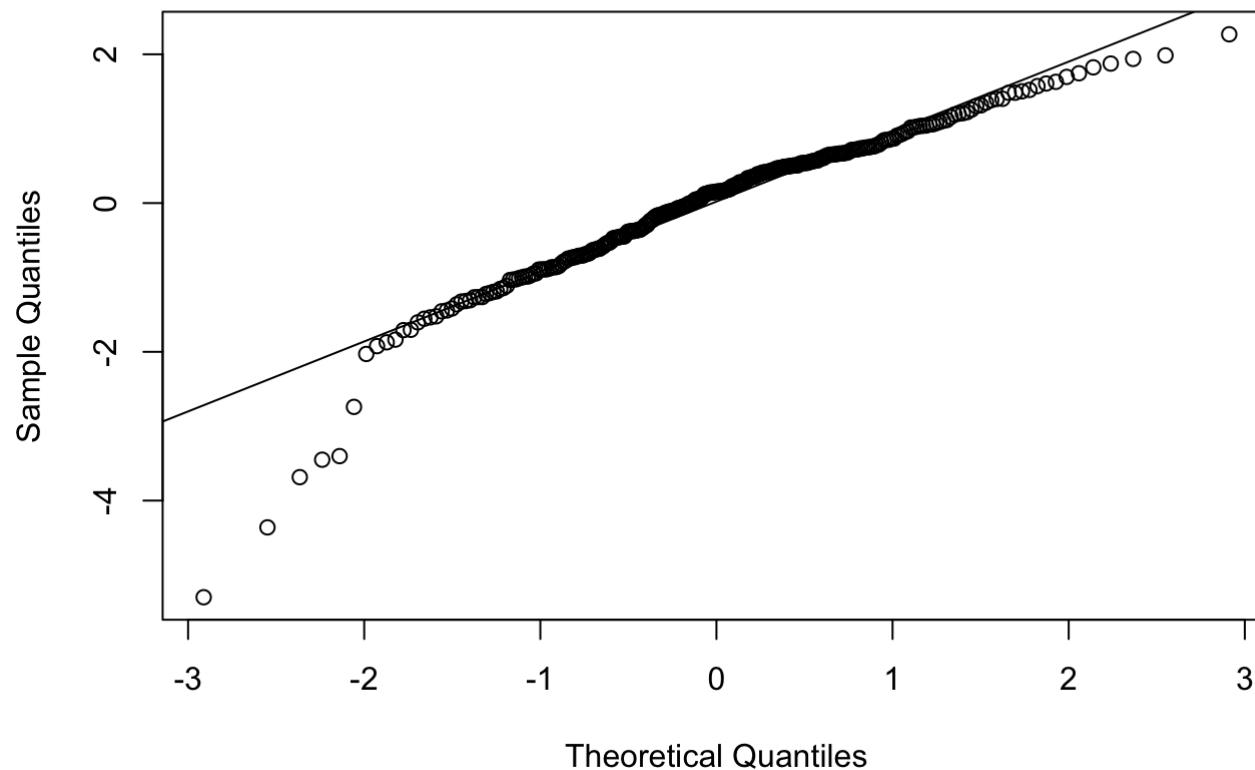
```
## Groups           Name      Std.Dev.  
## Target.Population (Intercept) 1.24777  
## Template.Population (Intercept) 1.43786  
## Population       (Intercept) 0.81398  
## Population.1     (Intercept) 0.81398  
## Residual          1.13889
```

```
plot(mPC1) # residuals plot
```



```
qqnorm(resid(mPC1)); qqline(resid(mPC1)) #qqplot
```

Normal Q-Q Plot



Now lets compare the models using

ANOVA

```
anova(mPC1,mNull)
```

```
## Data: dat
## Models:
## mNull: PC1 ~ 1 + (1 | Population/Template) + (1 | Population/Target)
## mPC1: PC1 ~ Population + (1 | Population/Template) + (1 | Population/Target)
##        Df    AIC    BIC  logLik deviance Chisq Chi Df Pr(>Chisq)
## mNull   6 1055.5 1077.3 -521.76    1043.5
## mPC1    9 1050.3 1083.0 -516.16    1032.3 11.184      3    0.01077 *
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

So there is a significant population effect!!!

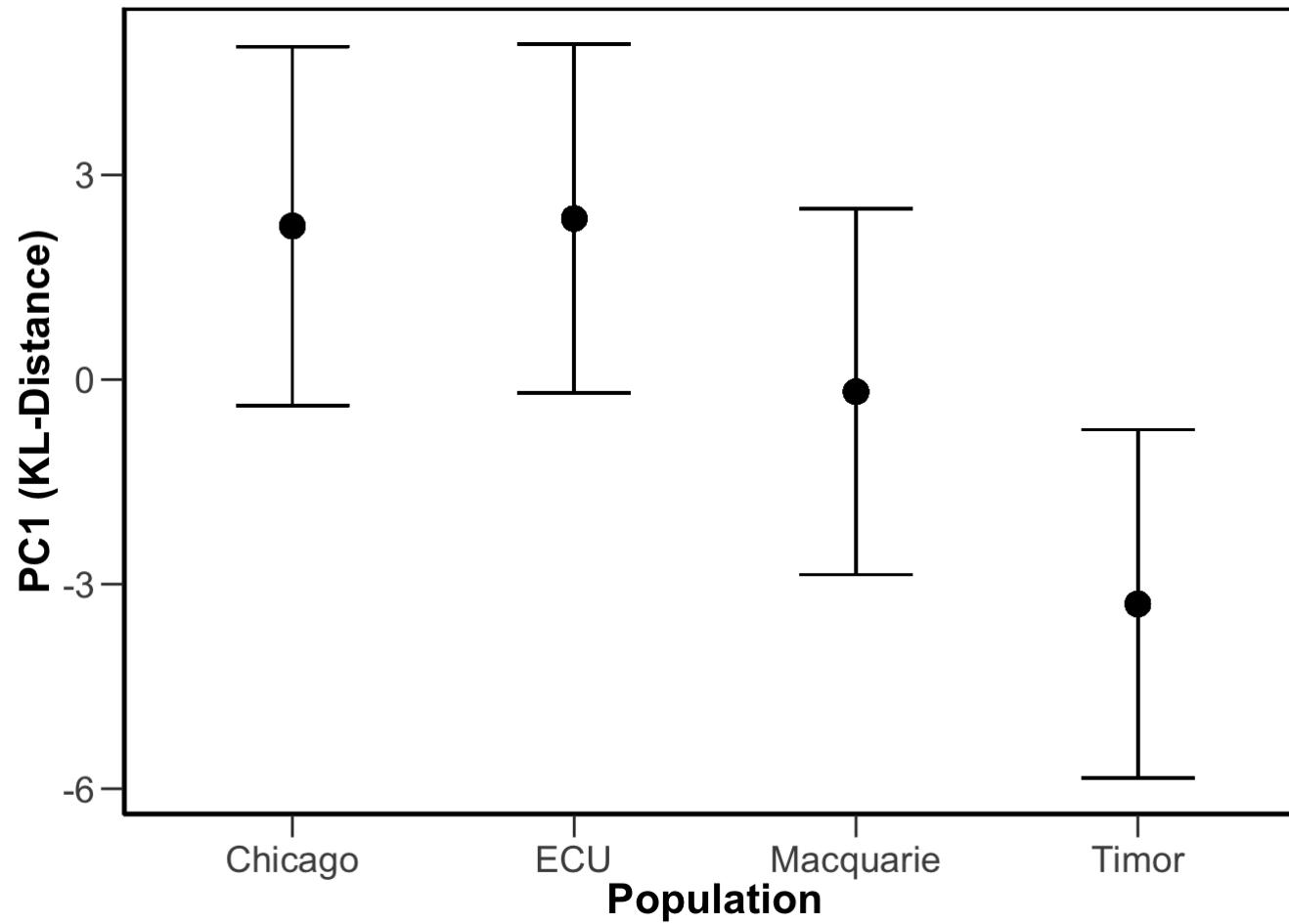
Lets load a cheesy little function for extracting the predictions and CIs for plotting.

```
easyPredCInorm <- function(model,newdata,alpha=0.05) {
  ## baseline prediction, on the linear predictor scale:
  pred0 <- predict(model,re.form=NA,newdata=newdata)
  ## fixed-effects model matrix for new data
  X <- model.matrix(formula(model,fixed.only=TRUE)[-2],
                     newdata)
  beta <- fixef(model) ## fixed-effects coefficients
  V <- vcov(model)      ## variance-covariance matrix of beta
  pred.se <- sqrt(diag(X %*% V %*% t(X))) ## std errors of predictions
  ## identity link
  linkinv <- I
  ## construct 95% Normal CIs on the log scale and
  ## transform back to the response (probability) scale:
  crit <- -qnorm(alpha/2)
  linkinv(cbind(lwr=pred0-crit*pred.se,
                upr=pred0+crit*pred.se))
}
```

Now we can extract model predicted means and confidence intervals and draw a pretty picture.

```
dat$predicted=(predict(mPC1,re.form=NA))
pframe <- data.frame(Population=dat$Population)
dat$LCI=as.vector((easyPredCInorm(mPC1,pframe))[,1])
dat$UCI=as.vector((easyPredCInorm(mPC1,pframe))[,2])

ggplot(dat,aes(x=Population,y=predicted))+geom_point(size=4)+geom_errorbar(aes(ymin=LCI,ymax=UCI),
  width=0.4,
  size=0.5)+
  ylab("PC1 (KL-Distance)")+xlab("Population")+theme_bw() +
  theme(axis.title=element_text(vjust=1,size=16,face="bold"), axis.text=element_text(size=14),
  axis.text.x = element_text(vjust=0.65, hjust=0.5, size=14), panel.border =
  element_rect(colour = "black",size=1.25)) + theme(axis.ticks.length=unit(0.3,"cm"))+
  theme(panel.grid.major = element_blank(), panel.grid.minor =element_blank(), axis.line =
  element_line(colour = "black"))
```



```
#ggsave( "PC1-Rplotv_4Pop_Nested.pdf", plot=last_plot(), device=NULL, path=NULL, scale=1, width=NA, height=NA, dpi =300, limitsize=TRUE)
```

Having demonstrated an overall effect of Population on song variability. We now want to explore this in more detail. As far as we can tell, post hoc tests are not supported for blme, but readers surely will want to know if there is significant variation within TGC, namely between domesticated and wild. So we drop the Timors from the dataset and run this again.

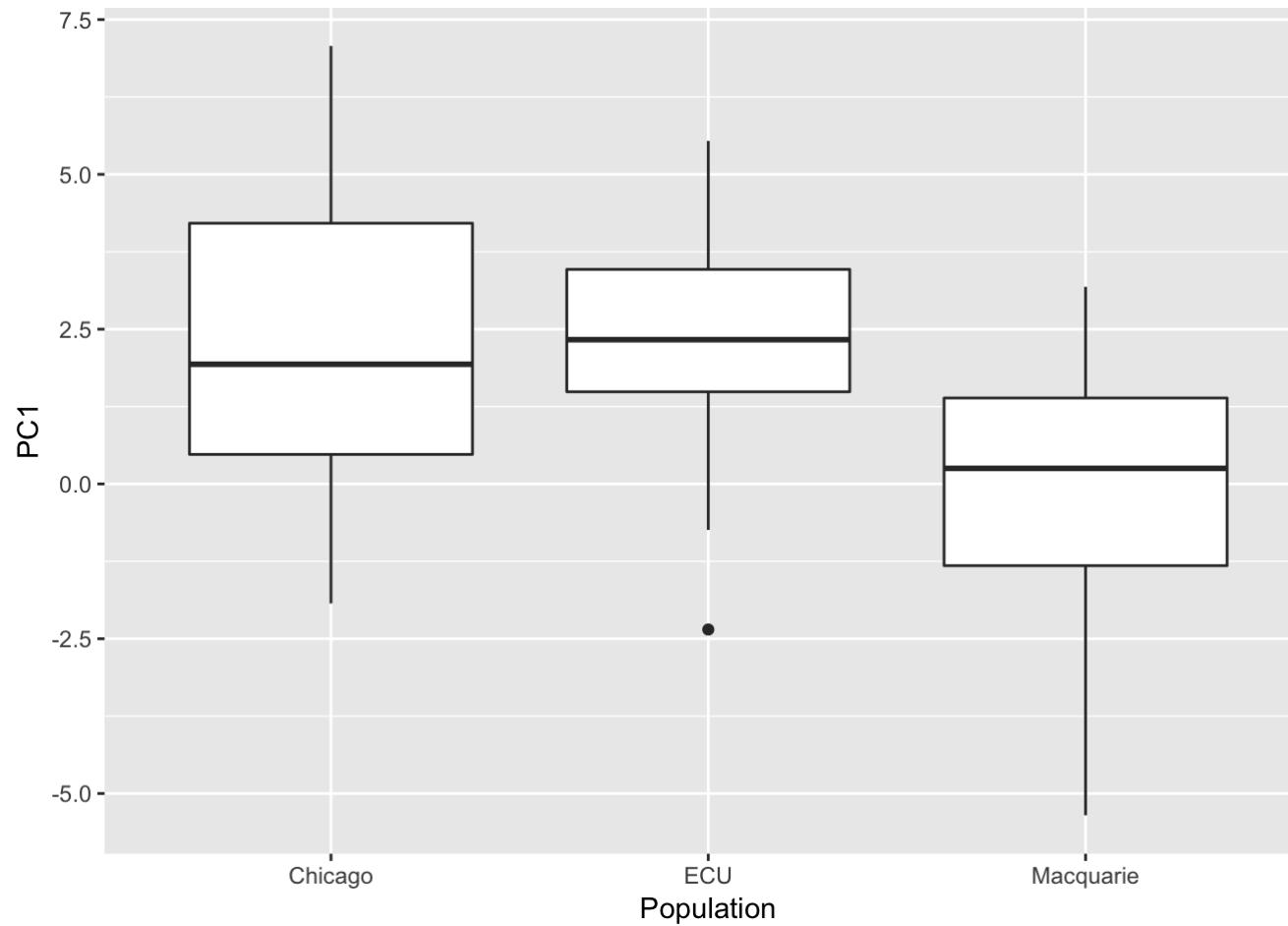
```
dat2=read.table(paste(path,"logKL_3pops_PCA.txt",sep="/"),sep = "\t",header=TRUE)
str(dat2)
```

```

## 'data.frame': 188 obs. of 30 variables:
## $ X           : int 123 124 125 126 127 128 129 130 131 132 ...
## $ logamplitude : num 1.04 0.943 0.9 1.061 0.951 ...
## $ logpitch     : num 0.534 0.682 0.766 0.723 0.911 ...
## $ logFM        : num 0.607 0.719 0.949 0.772 0.826 ...
## $ logAM2       : num 0.567 0.676 0.773 0.79 0.694 ...
## $ logentropy    : num 0.542 0.594 0.724 0.746 0.558 ...
## $ logpitchgoodness: num 0.575 0.694 0.891 0.775 0.939 ...
## $ logmeanfreq   : num 0.601 0.705 0.723 0.763 0.855 ...
## $ logvpitch     : num 0.383 0.542 0.547 0.67 0.142 ...
## $ logvFM        : num 0.511 0.608 0.728 0.756 0.538 ...
## $ logventropy    : num 0.453 0.61 0.642 0.694 0.54 ...
## $ logvpitchgoodness: num 0.587 0.632 0.726 0.749 0.772 ...
## $ vmeanfreq     : num 0.511 0.559 0.614 0.666 0.468 ...
## $ PC1          : num 2.23 3.2 4.17 4.26 3.55 ...
## $ PC1neg       : num -2.23 -3.2 -4.17 -4.26 -3.55 ...
## $ PC2          : num -1.147 -0.566 -0.271 -0.684 -0.721 ...
## $ PC3          : num 0.112 0.0935 -0.4043 0.1929 -1.2095 ...
## $ PC4          : num 0.0871 0.177 -0.202 0.0909 0.1591 ...
## $ PC5          : num 0.289 0.184 0.173 0.271 -0.344 ...
## $ PC6          : num -0.0182 0.1712 0.2571 -0.019 1.0204 ...
## $ PC7          : num -0.0914 -0.2547 -0.1412 -0.1078 -0.1745 ...
## $ PC8          : num -0.04286 -0.00937 0.23907 0.09437 -0.13775 ...
## $ PC9          : num 0.00132 0.15736 0.14971 0.14637 0.25303 ...
## $ PC10         : num -0.000771 0.074141 0.377118 -0.054397 -0.124958 ...
## $ PC11         : num 0.16405 0.05001 -0.03662 0.15893 -0.00725 ...
## $ PC12         : num -0.1052 0.016 0.1205 0.0165 0.1192 ...
## $ Population    : Factor w/ 3 levels "Chicago", "ECU", ...: 2 2 2 2 2 2 2 2 2 ...
## $ Domesticated  : Factor w/ 2 levels "Domesticated", ...: 1 1 1 1 1 1 1 1 1 ...
## $ Template      : Factor w/ 25 levels "filtered_NewRunDP99_tclust", ...: 1 1 1 1 1 1 1 1 1 2 ...
## $ Target        : Factor w/ 25 levels "filtered_NewRunDP99_tclust", ...: 2 3 4 5 6 7 8 9 10 1 ...

```

```
print(ggplot(dat2,aes(x=Population,y=PC1))+geom_boxplot())
```

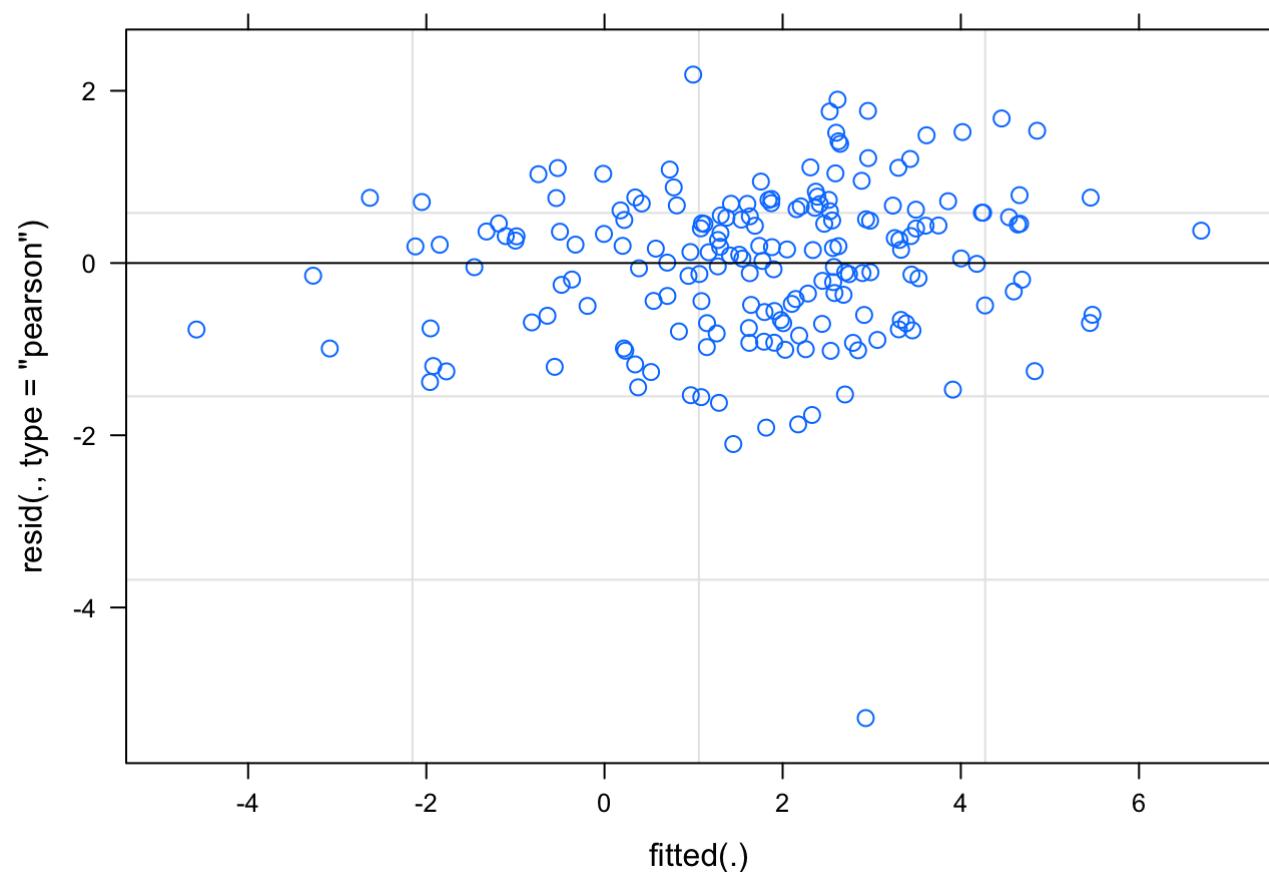


lets build some models

```
mPC1.3pop <- blmer(PC1 ~ Domesticated + (1 | Population/Template) + (1 | Population/Target), data = dat2, REML = F
ELSE)
VarCorr(mPC1.3pop) # Check for singularity
```

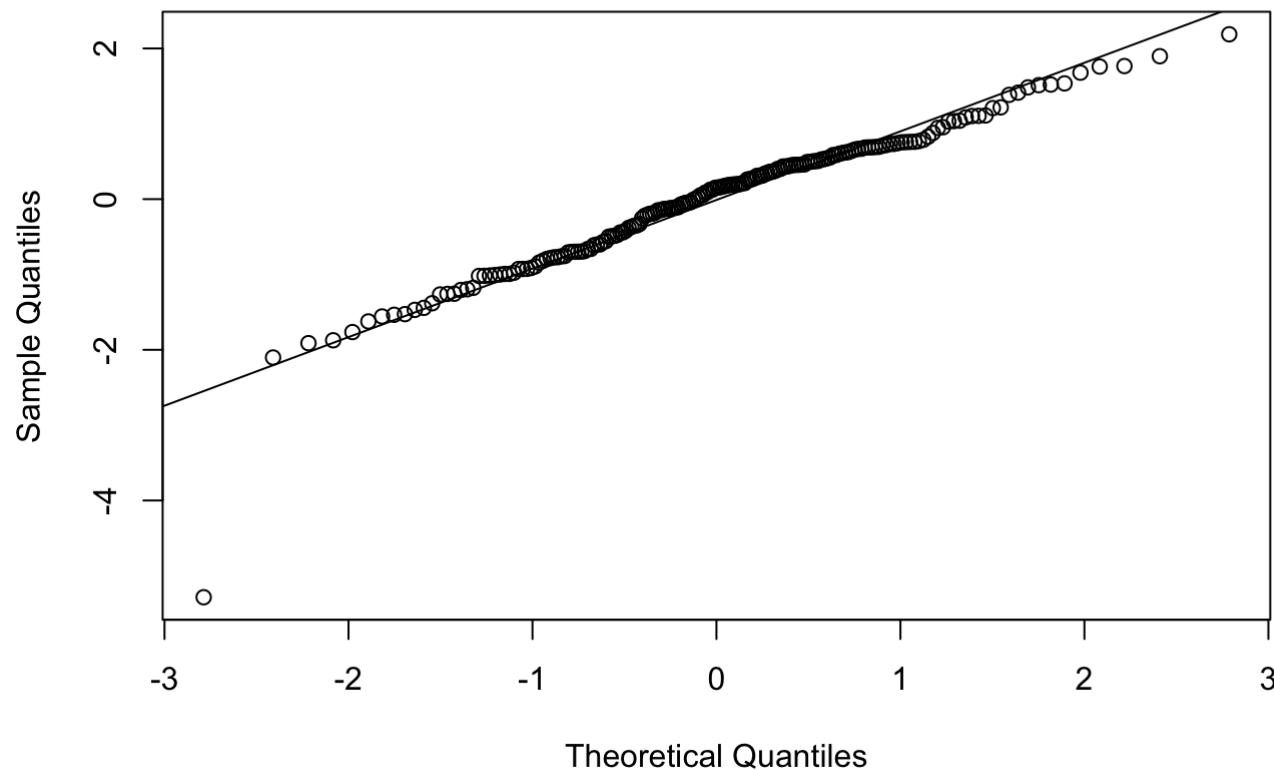
## Groups	Name	Std.Dev.
## Target.Population	(Intercept)	1.4479
## Template.Population	(Intercept)	1.1191
## Population	(Intercept)	29.6123
## Population.1	(Intercept)	29.6175
## Residual		1.0187

```
plot(mPC1.3pop) # residuals plot
```



```
qqnorm(resid(mPC1.3pop)); qqline(resid(mPC1.3pop)) #qqplot
```

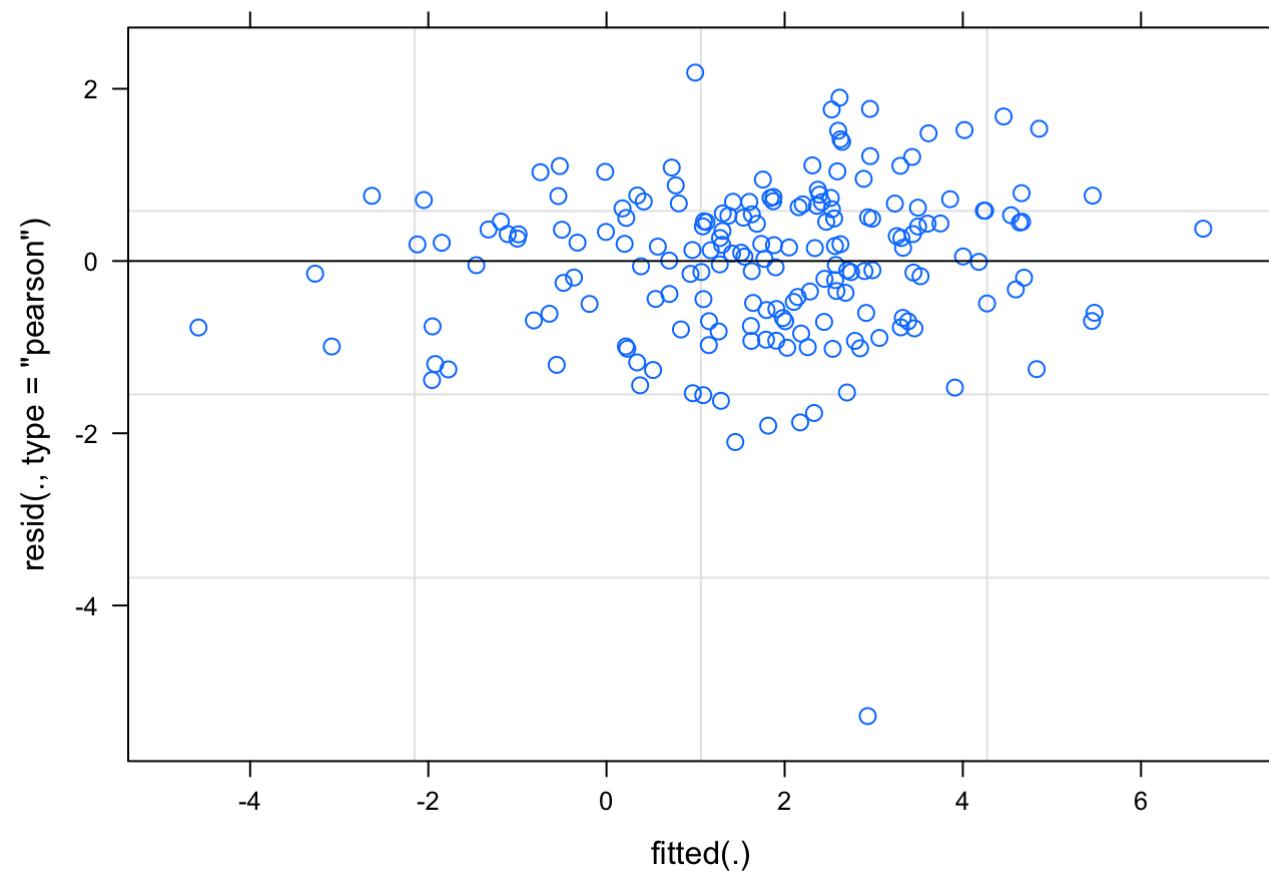
Normal Q-Q Plot



```
mPC1.3popNull <- blmer(PC1 ~ 1 + (1 | Population/Template) + (1 | Population/Target), data = dat2, REML = FALSE)
VarCorr(mPC1.3popNull) # Check for singularity
```

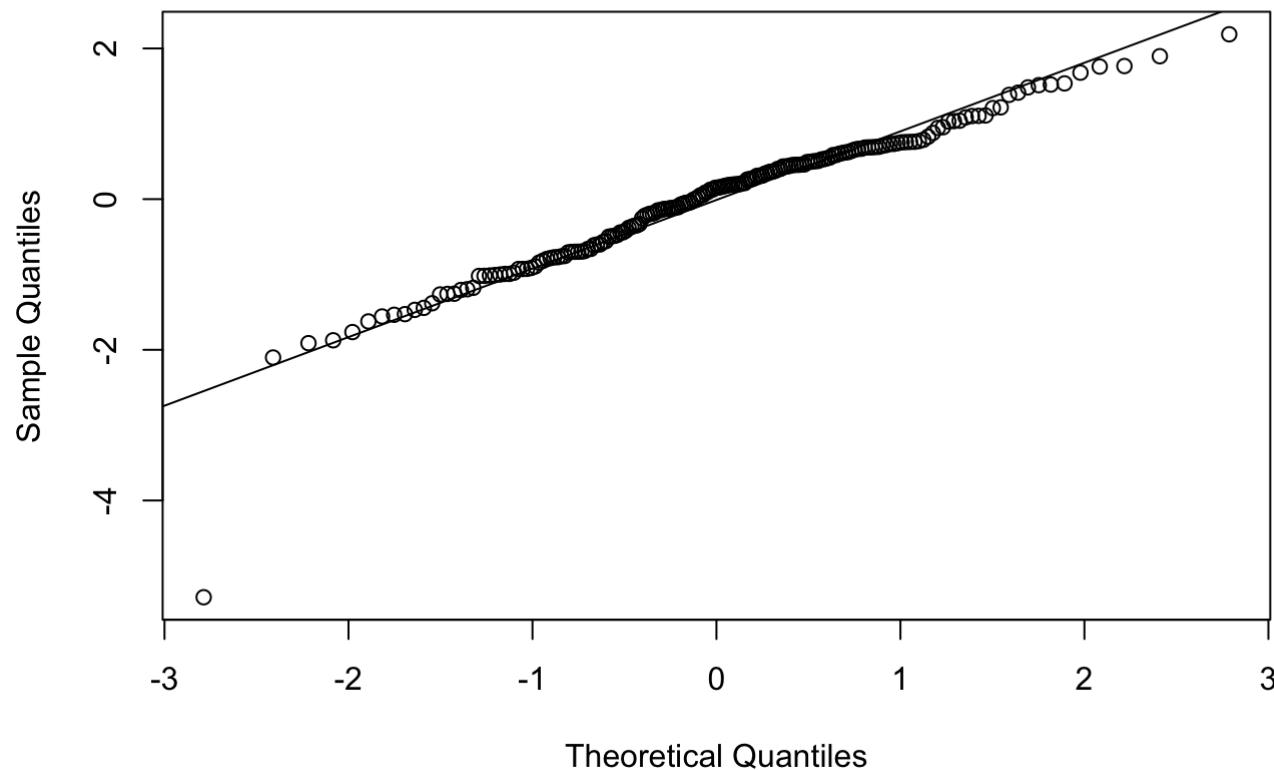
```
## Groups           Name      Std.Dev.
## Target.Population (Intercept) 1.4479
## Template.Population (Intercept) 1.1191
## Population        (Intercept) 45.4907
## Population.1       (Intercept) 45.4931
## Residual           1.0187
```

```
plot(mPC1.3popNull) # residuals plot
```



```
qqnorm(resid(mPC1.3popNull)); qqline(resid(mPC1.3popNull)) #qqplot
```

Normal Q-Q Plot



Note that the estimated errors are

quite large here, likely due to the small number of levels within the “Population” variable in the random factors.

```
anova(mPC1.3popNull, mPC1.3pop)
```

```
## Data: dat2
## Models:
## mPC1.3popNull: PC1 ~ 1 + (1 | Population/Template) + (1 | Population/Target)
## mPC1.3pop: PC1 ~ Domesticated + (1 | Population/Template) + (1 | Population/Target)
##              Df    AIC    BIC  logLik deviance Chisq Chi Df Pr(>Chisq)
## mPC1.3popNull 6 700.96 720.38 -344.48     688.96
## mPC1.3pop     7 700.38 723.04 -343.19     686.38 2.5761      1     0.1085
```

Ok, so a model including “domesticated” as a fixed effect is not better than the null model.

Lastly, we also want to directly compare the two subspecies. This is not strictly a post-hoc test in this case. Here we are able to incorporate additional data. For each of the two subspecies, TGG and TGC we have a small sample of additional data from the UIUC. This data allows to determine whether the low variability observed above has been stable over time/across populations.

```
path<-("~/Documents/Projects/Domestication/OpenBiology/Subspecies"
dat3=read.table(paste(path,"logKL_TvA_PCA.txt",sep="/"),sep = "\t",header=TRUE)
str(dat3)
```

```

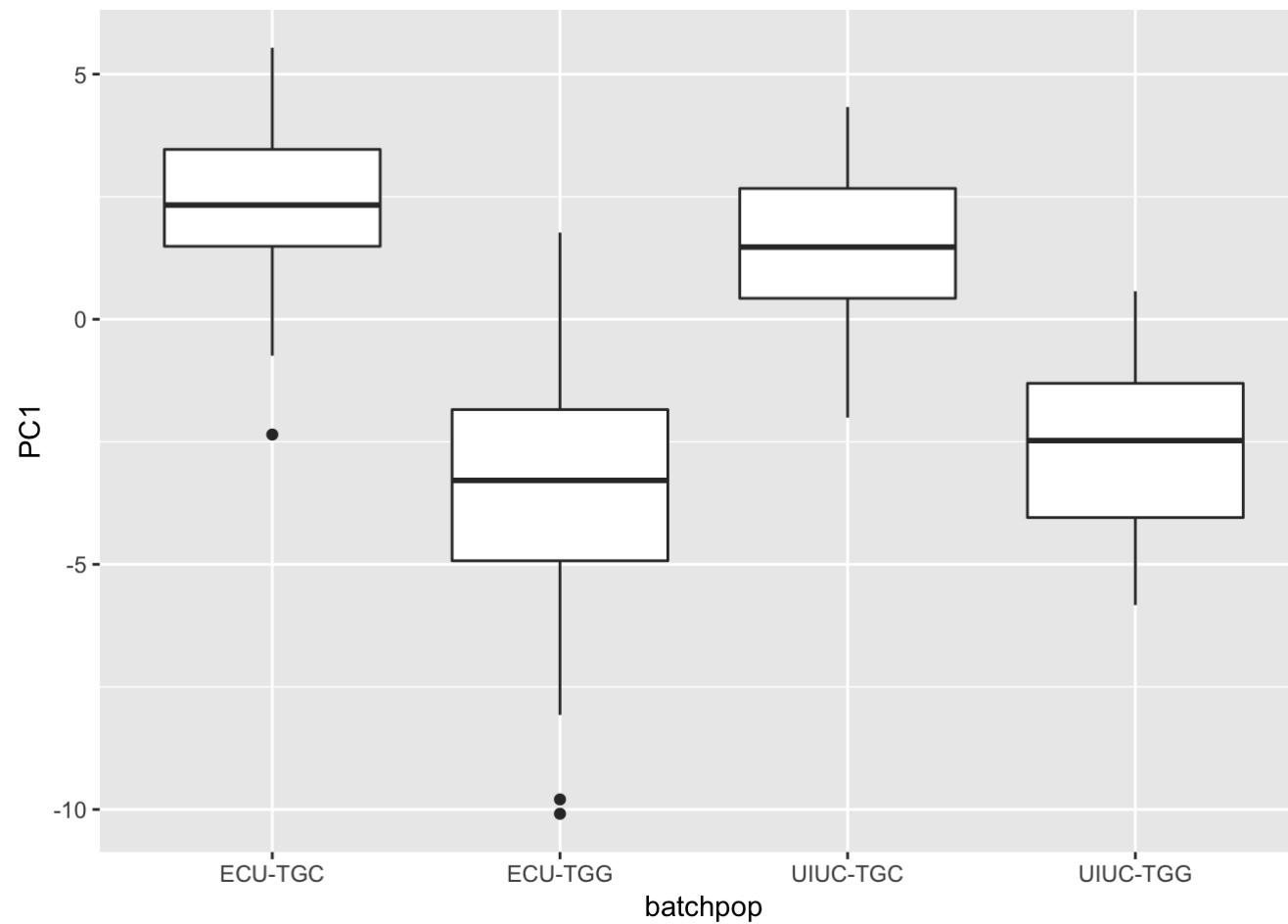
## 'data.frame': 212 obs. of 31 variables:
## $ X           : int 1 2 3 4 5 6 7 8 9 10 ...
## $ logamplitude : num 0.593 0.652 0.478 0.294 0.478 ...
## $ logpitch     : num 0.713 0.721 0.497 0.179 0.421 ...
## $ logFM        : num 0.611 0.725 0.448 0.23 0.373 ...
## $ logAM2       : num 0.625 0.719 0.615 0.398 0.447 ...
## $ logentropy    : num 0.617 0.724 0.456 0.247 0.361 ...
## $ logpitchgoodness: num 0.721 0.665 0.594 0.33 0.585 ...
## $ logmeanfreq   : num 0.621 0.691 0.447 0.407 0.491 ...
## $ logvpitch     : num 0.5507 0.6487 0.4843 0.0129 0.3282 ...
## $ logvFM        : num 0.5171 0.6269 0.4334 0.0189 0.4684 ...
## $ logventropy    : num 0.492 0.676 0.5417 -0.0239 0.4716 ...
## $ logvpitchgoodness: num 0.607 0.552 0.463 0.171 0.521 ...
## $ vmeanfreq      : num 0.471 0.546 0.408 -0.174 0.306 ...
## $ PC1          : num 2.489 3.208 1.314 -2.006 0.758 ...
## $ PC1neg        : num -2.489 -3.208 -1.314 2.006 -0.758 ...
## $ PC2          : num 0.448 0.479 0.612 0.309 0.454 ...
## $ PC3          : num 0.0136 0.368 0.3479 -0.3448 -0.1312 ...
## $ PC4          : num 0.029 0.1011 -0.1367 0.4306 0.0731 ...
## $ PC5          : num 0.382 0.0571 0.3869 0.5023 0.5184 ...
## $ PC6          : num 0.5253 0.1652 0.0143 0.3325 -0.1171 ...
## $ PC7          : num -0.017 -0.0171 -0.4781 0.1454 -0.4093 ...
## $ PC8          : num 0.0176 0.1835 0.0938 0.6944 -0.3397 ...
## $ PC9          : num 0.0264 0.2188 0.02 0.5259 0.1631 ...
## $ PC10         : num -0.0851 0.1441 -0.2972 -0.3736 -0.3687 ...
## $ PC11         : num 0.0508 -0.067 -0.0722 -0.1573 0.129 ...
## $ PC12         : num 0.1568 0.0736 0.2731 0.2577 0.1433 ...
## $ batch         : Factor w/ 2 levels "ECU","UIUC": 2 2 2 2 2 2 2 2 2 ...
## $ Population    : Factor w/ 2 levels "TGC","TGG": 1 1 1 1 1 1 1 1 1 ...
## $ batchpop      : Factor w/ 4 levels "ECU-TGC","ECU-TGG",...: 3 3 3 3 3 3 3 3 3 ...
## $ Template      : Factor w/ 29 levels "filtered_BL65_tclust",...: 15 15 15 16 16 16 17 17 17 18 ...
## $ Target        : Factor w/ 29 levels "filtered_BL65_tclust",...: 16 17 18 15 17 18 15 16 18 15 ...

```

```

require(ggplot2)
print(ggplot(dat3,aes(x=batchpop,y=PC1))+geom_boxplot())

```



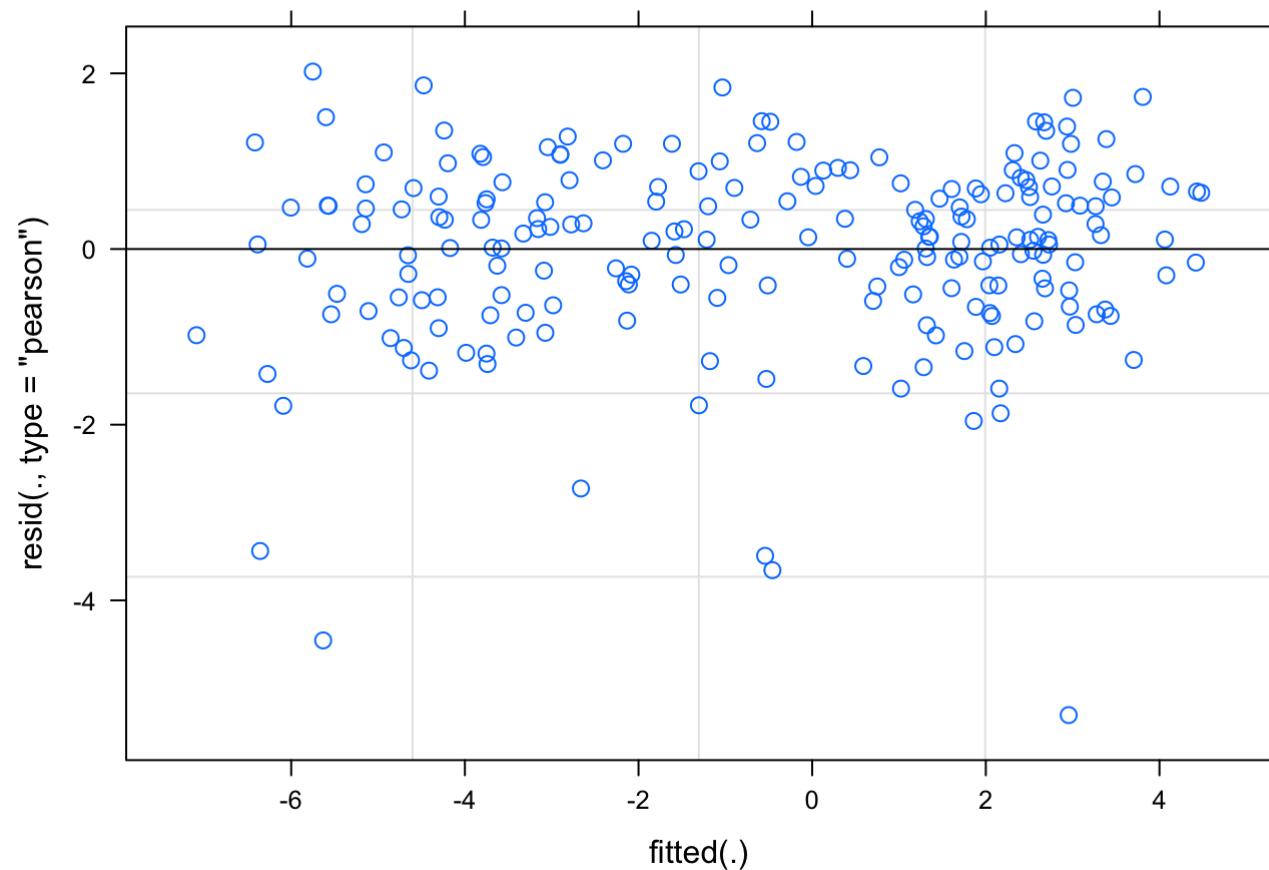
Superficially, we can clearly see that

TGG is low variability in both populations. Lets test.

```
mPC1.TvA<- blmer(PC1 ~ batchpop + (1 | batchpop/Template) + (1 | batchpop/Target), data = dat3, REML = FALSE)
VarCorr(mPC1.TvA) # Check for singularity
```

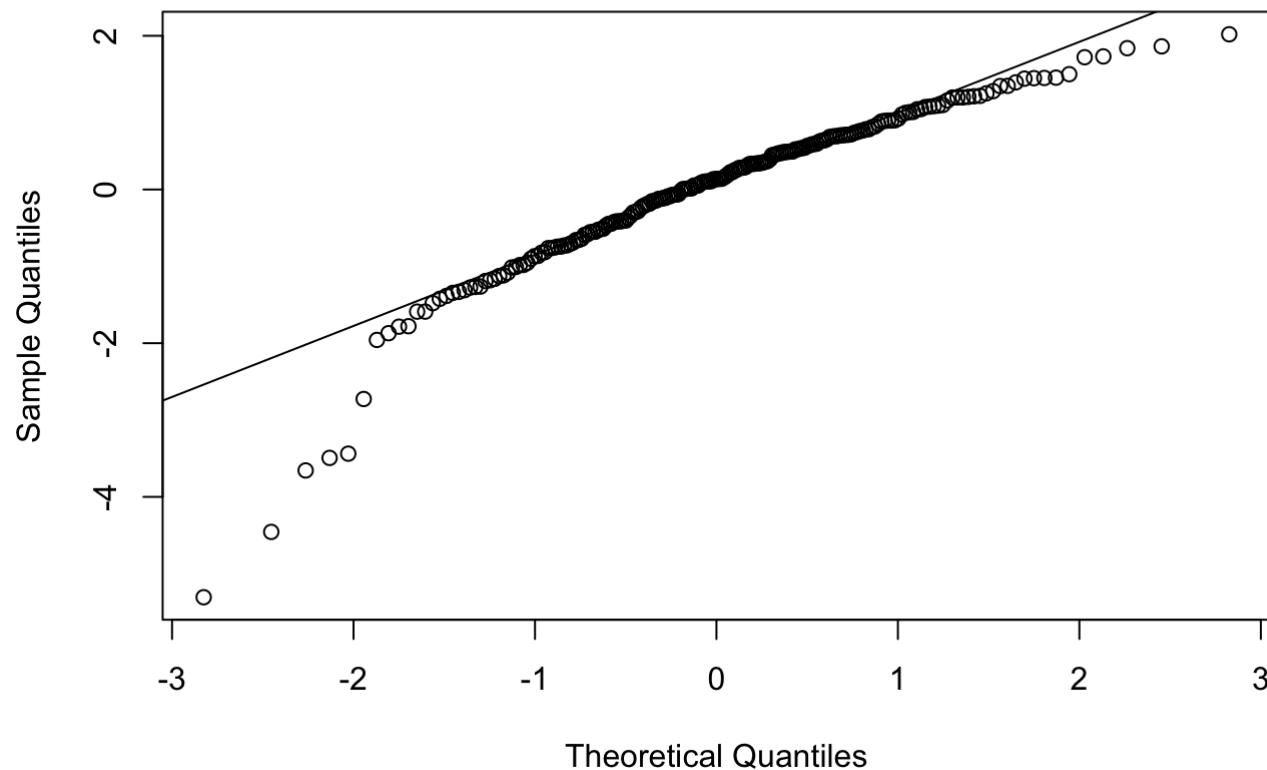
```
## Groups           Name      Std.Dev.
## Target.batchpop (Intercept) 0.83666
## Template.batchpop (Intercept) 1.50896
## batchpop        (Intercept) 0.87592
## batchpop.1       (Intercept) 0.87592
## Residual          1.17491
```

```
plot(mPC1.TvA) # residuals plot
```



```
qqnorm(resid(mPC1.TvA)); qqline(resid(mPC1.TvA)) #qqplot
```

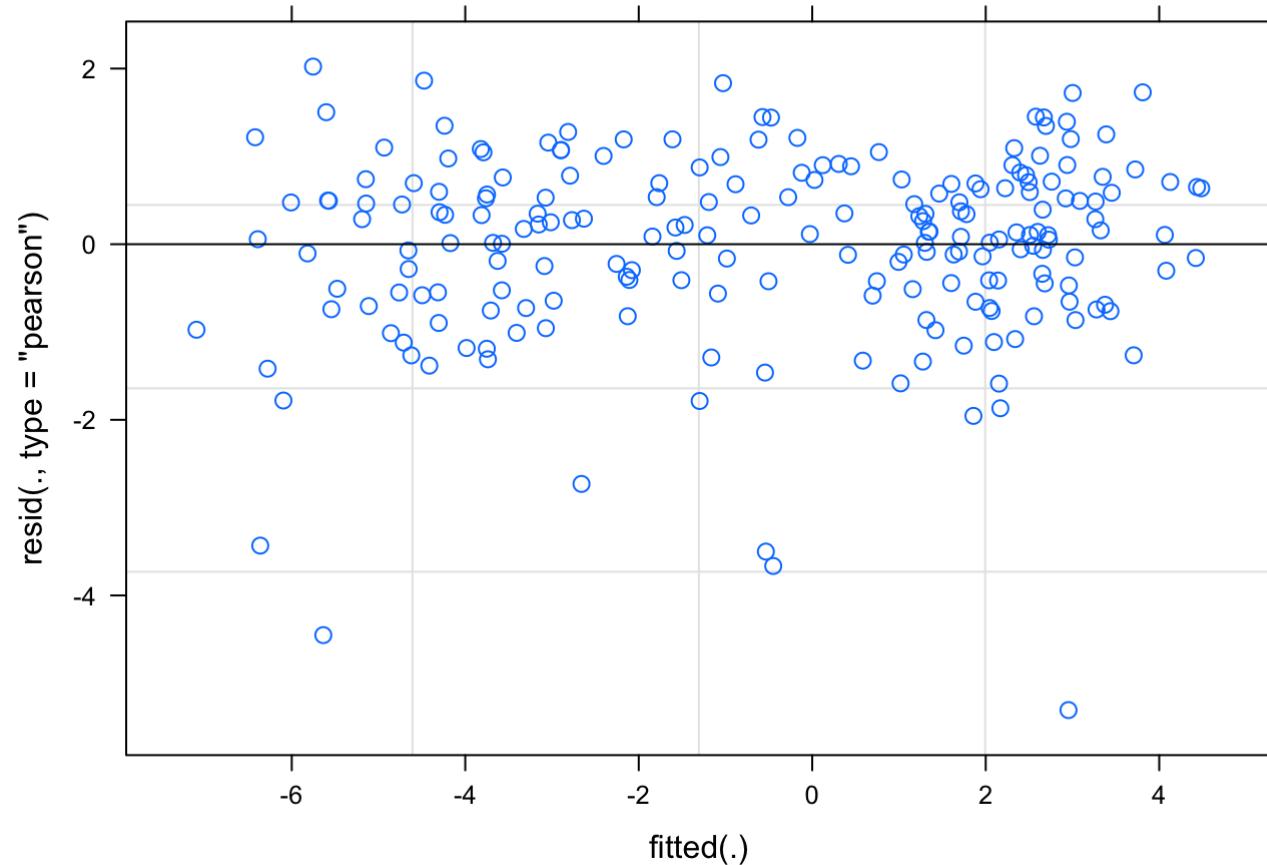
Normal Q-Q Plot



```
mPC1.TvA.null <- blmer(PC1 ~ 1 + (1 | batchpop/Template) + (1 | batchpop/Target), data = dat3, REML = FALSE)
VarCorr(mPC1.TvA.null) # Check for singularity
```

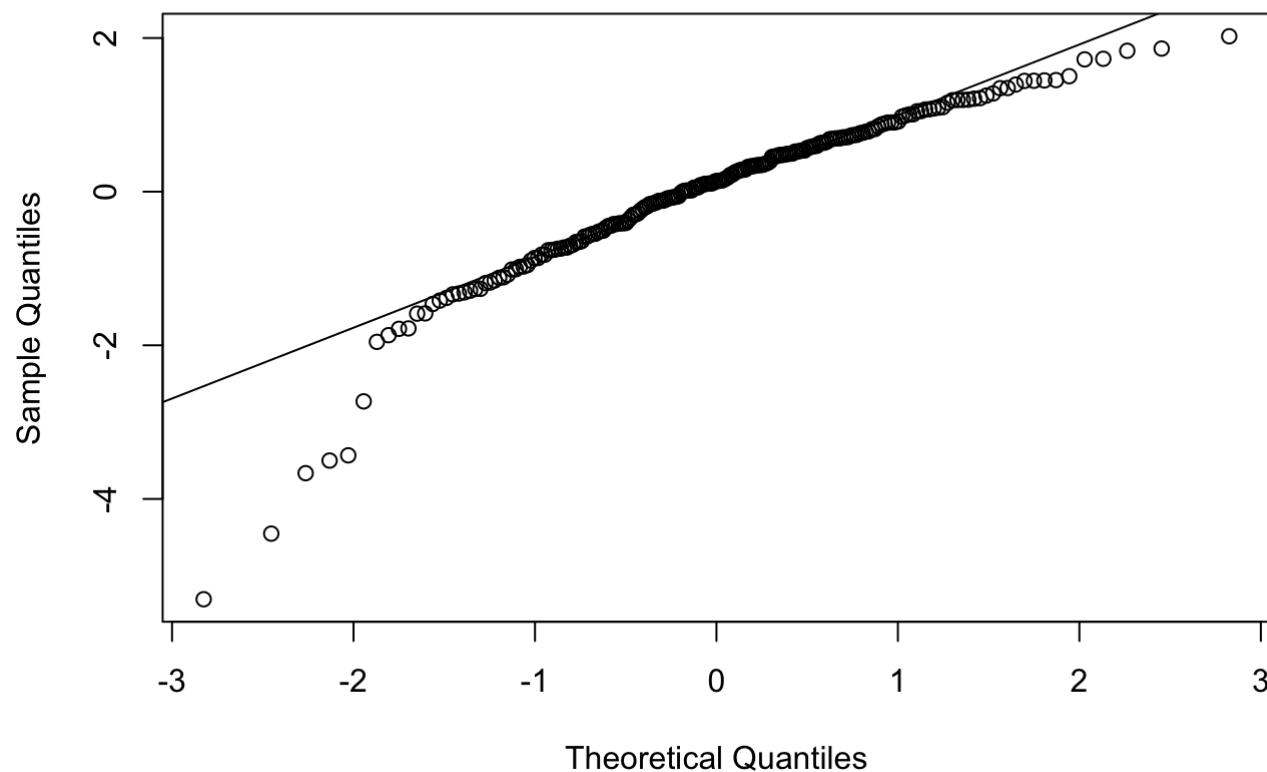
```
## Groups           Name      Std.Dev.
## Target.batchpop (Intercept) 0.84285
## Template.batchpop (Intercept) 1.53347
## batchpop        (Intercept) 3.53784
## batchpop.1       (Intercept) 3.53784
## Residual          1.17481
```

```
plot(mPC1.TvA.null) # residuals plot
```



```
qqnorm(resid(mPC1.TvA.null)); qqline(resid(mPC1.TvA.null)) #qqplot
```

Normal Q-Q Plot

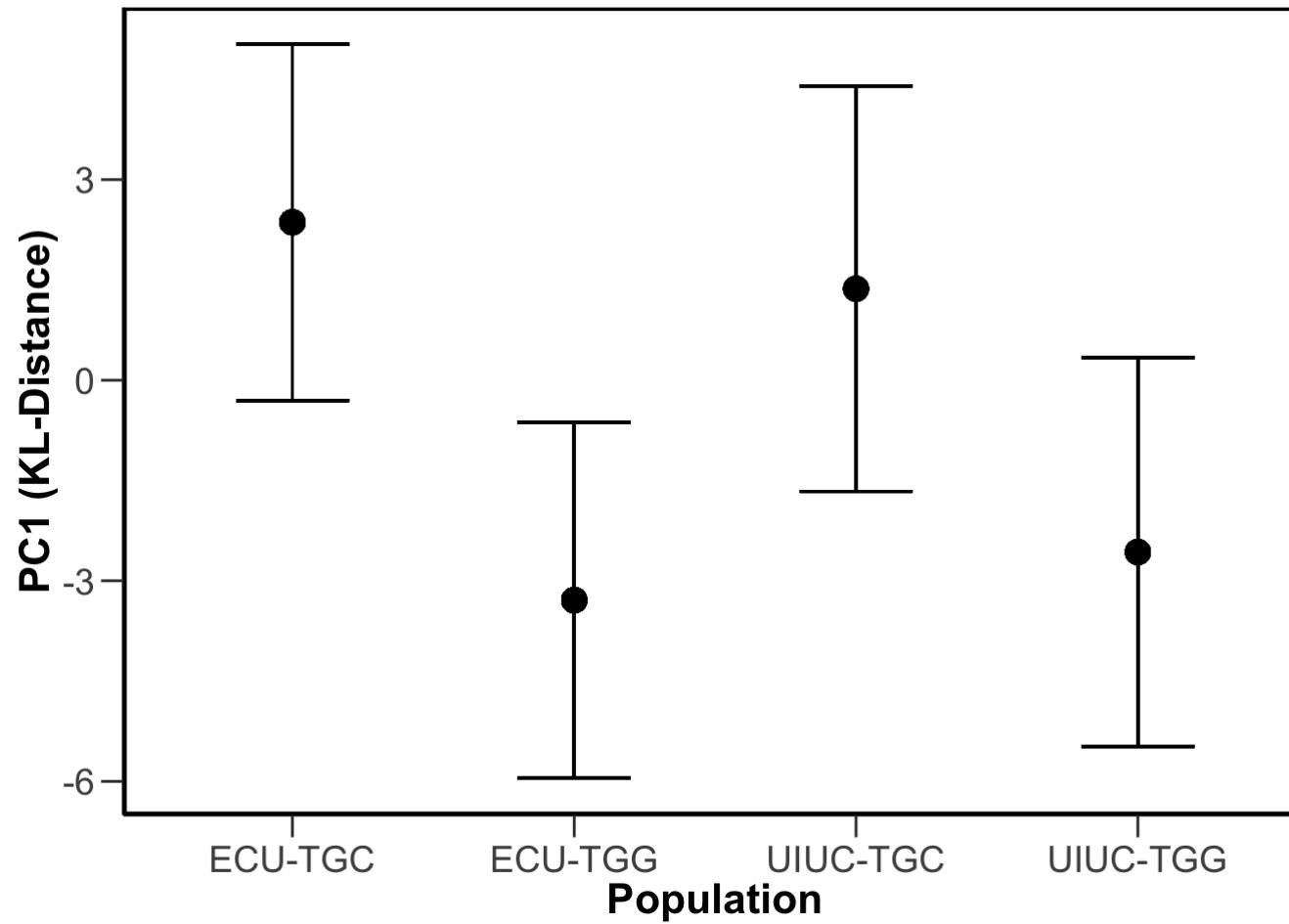


```
anova(mPC1.TvA,mPC1.TvA.null)
```

```
## Data: dat3
## Models:
## mPC1.TvA.null: PC1 ~ 1 + (1 | batchpop/Template) + (1 | batchpop/Target)
## mPC1.TvA: PC1 ~ batchpop + (1 | batchpop/Template) + (1 | batchpop/Target)
##             Df      AIC      BIC logLik deviance Chisq Chi Df Pr(>Chisq)
## mPC1.TvA.null 6  809.92  830.06 -398.96    797.92
## mPC1.TvA       9  804.85  835.06 -393.43    786.85 11.068      3   0.01136 *
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
dat3$predicted=(predict(mPC1.TvA,re.form=NA))
pframe <- data.frame(batchpop=dat3$batchpop)
dat3$LCI=as.vector((easyPredCInorm(mPC1.TvA,pframe))[,1])
dat3$UCI=as.vector((easyPredCInorm(mPC1.TvA,pframe))[,2])

ggplot(dat3,aes(x=batchpop,y=predicted))+geom_point(size=4)+geom_errorbar(aes(ymin=LCI,ymax=UCI),
width=0.4,
size=0.5)+
ylab("PC1 (KL-Distance)")+xlab("Population")+theme_bw() +
theme(axis.title=element_text(vjust=1,size=16,face="bold"), axis.text=element_text(size=14),
axis.text.x = element_text(vjust=0.65, hjust=0.5, size=14), panel.border =
element_rect(colour = "black",size=1.25)) + theme(axis.ticks.length=unit(0.3,"cm"))+
theme(panel.grid.major = element_blank(), panel.grid.minor =element_blank(), axis.line =
element_line(colour = "black"))
```



```
#ggsave("PC1-RplotTVA_4pop.pdf", plot=last_plot(), device=NULL, path=NULL, scale=1, width=NA, height=NA, dpi=300, limitsize=TRUE)
```

Now, the last model we wish to run deals with comparing TGC and TGG finches that were experimentally tutored by Bengalese finches. We first examine KL distances to tutors.

```
path<-("~/Documents/Projects/Domestication"
KLtoTutors=read.table(paste(path, "logKL_forPCAandMMA_Xfost.txt", sep="/"), sep = "\t", header=TRUE)
str(KLtoTutors)
```

```

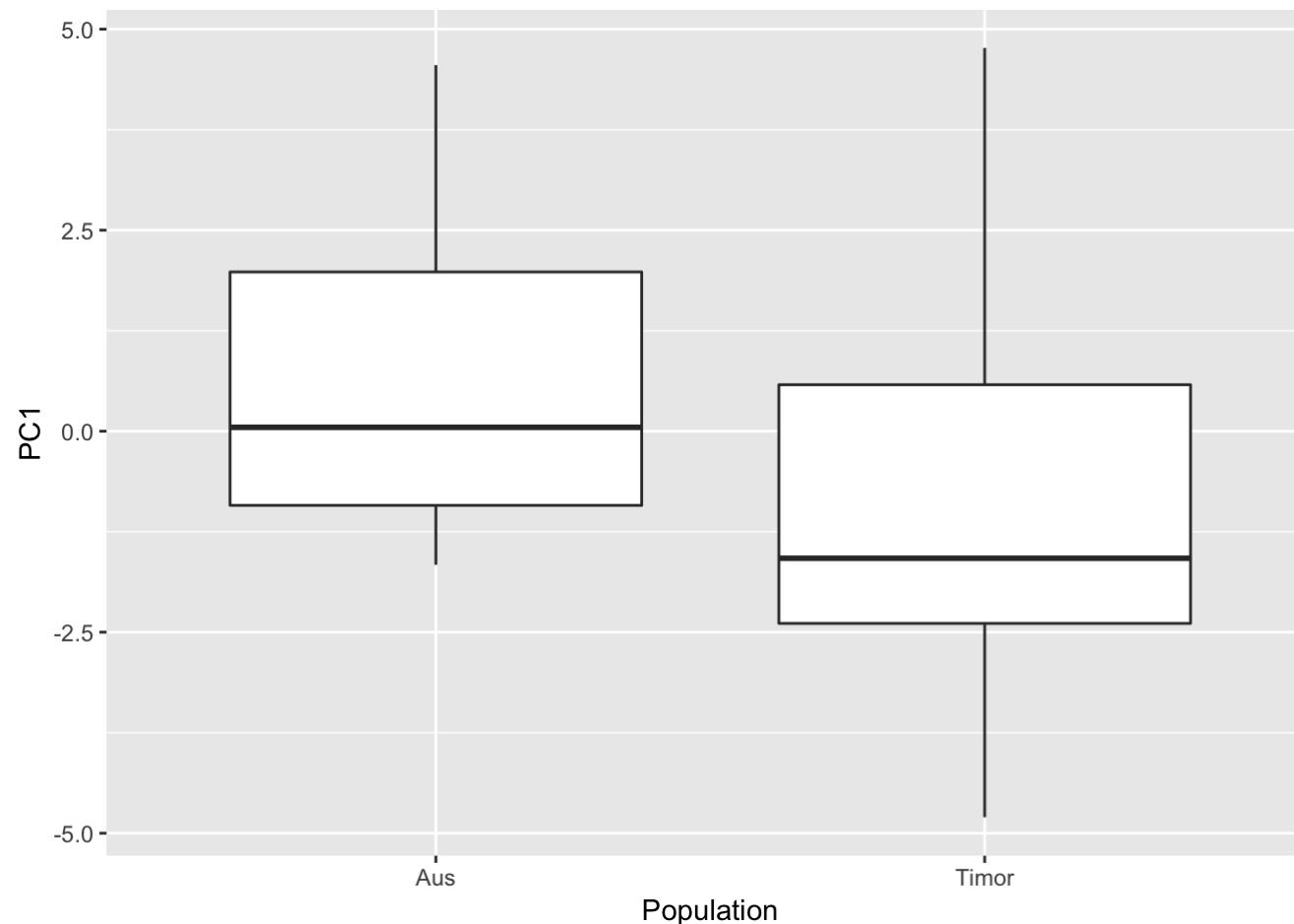
## 'data.frame':   18 obs. of  27 variables:
## $ logamplitude    : num  0.432 0.721 0.469 0.633 0.967 ...
## $ logpitch        : num  0.399 0.517 0.274 0.322 0.923 ...
## $ logFM           : num  0.55 0.279 0.524 0.337 0.585 ...
## $ logAM2          : num  0.5575 0.0545 0.3093 0.0957 0.6293 ...
## $ logentropy       : num  0.49 0.742 0.657 0.447 0.861 ...
## $ logpitchgoodness: num  0.669 0.49 0.276 0.42 0.725 ...
## $ logmeanfreq     : num  0.623 0.698 0.356 0.671 0.504 ...
## $ logvpitch       : num  -0.0527 -0.3973 -0.2404 -0.5944 0.2876 ...
## $ logvFM          : num  0.326 0.212 0.126 0.191 0.761 ...
## $ logventropy     : num  0.1304 0.143 0.0728 0.0296 0.932 ...
## $ logvpitchgoodness: num  0.232 0.232 -0.157 -0.103 0.368 ...
## $ vmeanfreq       : num  0.0758 0.142 -0.0503 -0.0874 0.4318 ...
## $ PC1             : num  1.268 0.663 -0.566 -0.894 4.553 ...
## $ PC2             : num  -0.947 -1.255 -0.733 -1.513 0.714 ...
## $ PC3             : num  0.79884 -1.21753 -0.00629 -0.32591 0.78802 ...
## $ PC4             : num  0.466 0.269 0.671 0.406 0.196 ...
## $ PC5             : num  -0.95 0.794 0.175 -0.363 1.261 ...
## $ PC6             : num  0.22301 0.00671 0.2758 0.09311 0.23199 ...
## $ PC7             : num  0.3141 0.5155 -0.9192 -0.0643 -0.2569 ...
## $ PC8             : num  0.299 -0.189 0.879 -0.607 -0.558 ...
## $ PC9             : num  -0.6274 0.0216 0.2184 0.2184 -0.1268 ...
## $ PC10            : num  0.1677 0.1854 -0.2392 0.0494 -0.0106 ...
## $ PC11            : num  0.2671 0.2916 0.2222 0.1099 -0.0342 ...
## $ Population      : Factor w/ 2 levels "Aus","Timor": 1 1 1 1 1 1 1 1 1 ...
## $ Stage           : Factor w/ 2 levels "egg","P30": 1 2 1 1 2 2 2 2 2 ...
## $ Tutor           : Factor w/ 5 levels "B_BR90","B_DB10",...: 3 3 3 3 2 2 1 4 4 5 ...
## $ Target.File     : Factor w/ 18 levels "filtered_BfADP56.xls",...: 3 6 9 10 4 8 5 1 2 7 ...

```

```

require(ggplot2)
print(ggplot(KLtoTutors,aes(x=Population,y=PC1))+geom_boxplot())

```



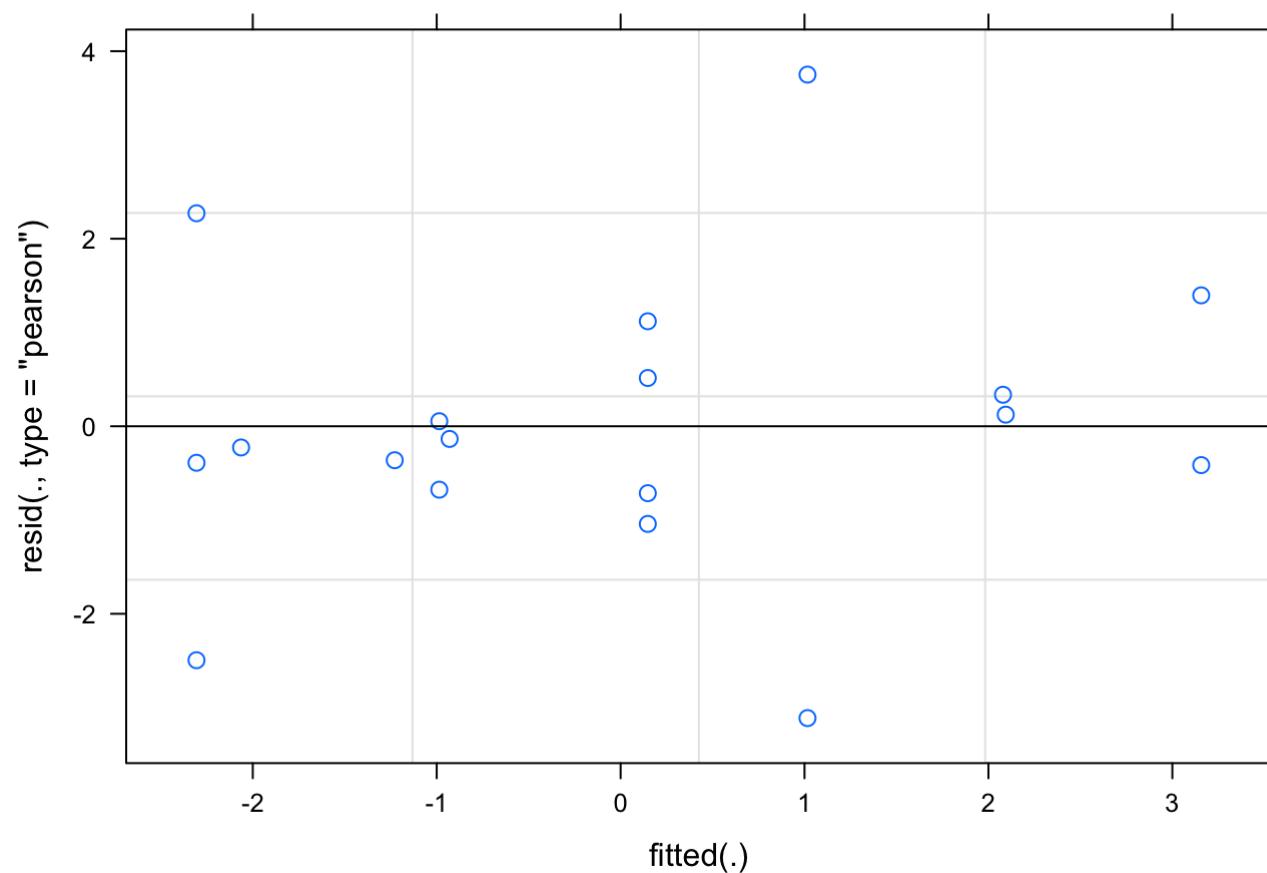
Ok, clearly not much exciting goin on

here... but we should probably do stats anyway. Here Template is the Bengalese Finch Tutor

```
library(blme)
mKLtoTutor <- blmer(PC1 ~ Population + (1 | Tutor), data = KLtoTutors, REML = FALSE)
VarCorr(mKLtoTutor) # Check for singularity
```

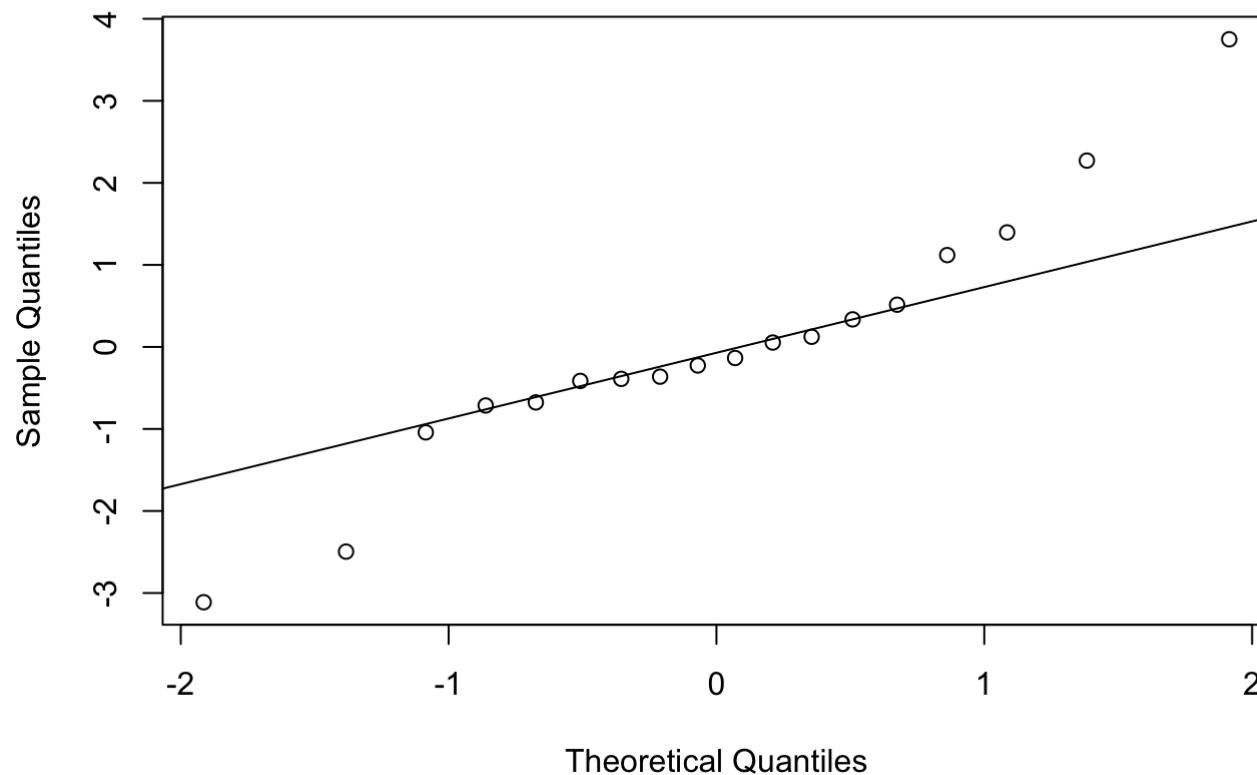
```
## Groups      Name       Std.Dev.
## Tutor      (Intercept) 2.283
## Residual           1.651
```

```
plot(mKLtoTutor) # residuals plot
```



```
qqnorm(resid(mKLtoTutor)); qqline(resid(mKLtoTutor)) #qqplot
```

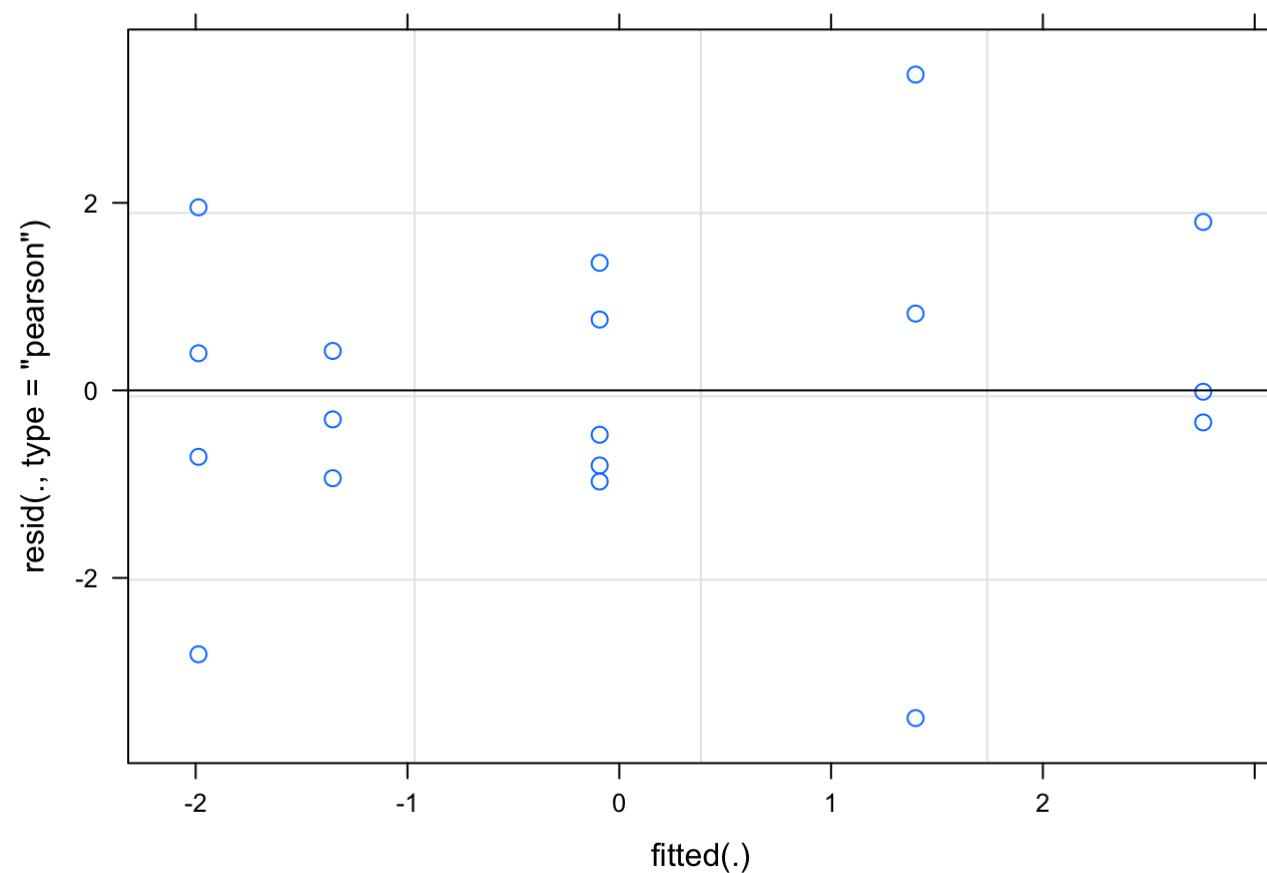
Normal Q-Q Plot



```
mKLtoTutorNull <- blmer(PC1 ~ 1 + (1 | Tutor), data = KLtoTutors, REML = FALSE)
VarCorr(mKLtoTutorNull) # Check for singularity
```

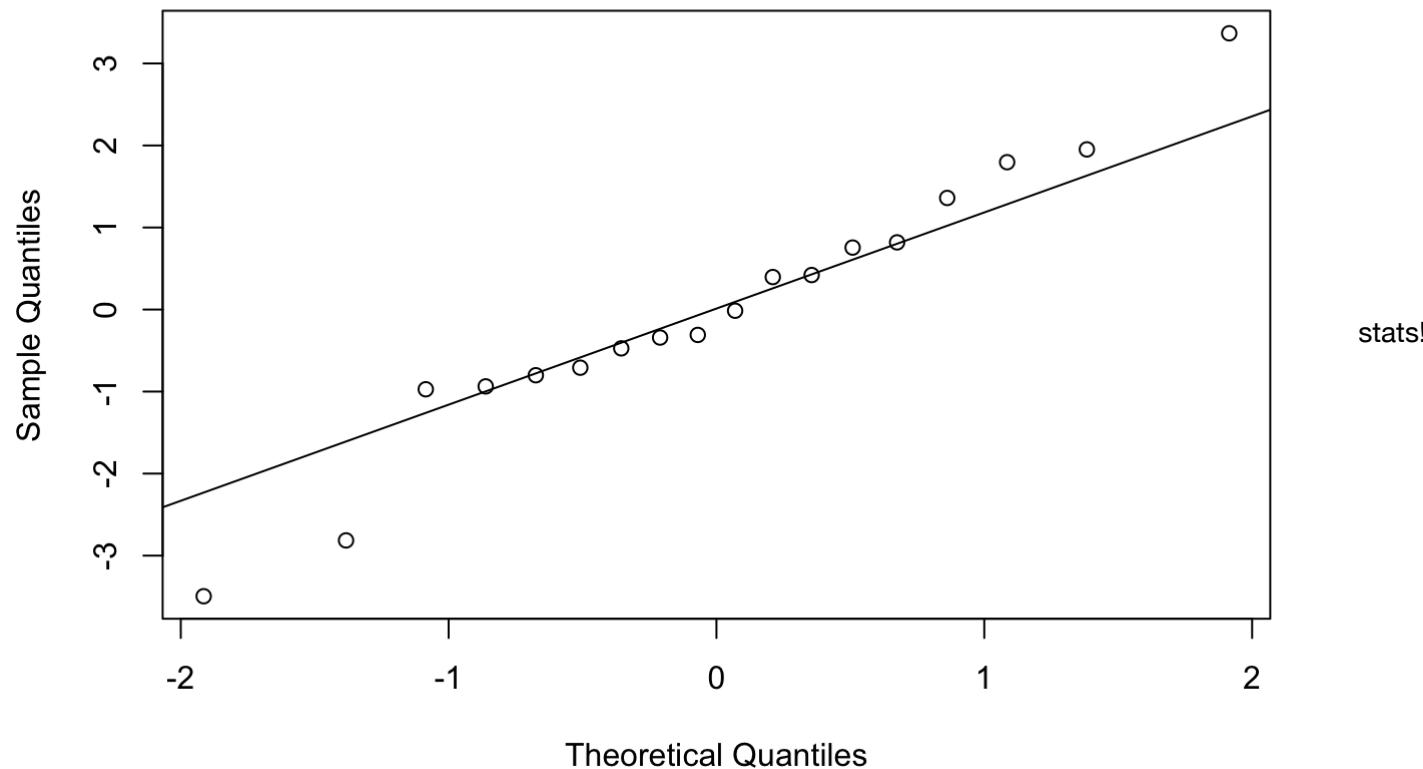
```
## Groups      Name       Std.Dev.
## Tutor      (Intercept) 2.3271
## Residual           1.7280
```

```
plot(mKLtoTutorNull) # residuals plot
```



```
qqnorm(resid(mKLtoTutorNull)); qqline(resid(mKLtoTutorNull)) #qqplot
```

Normal Q-Q Plot



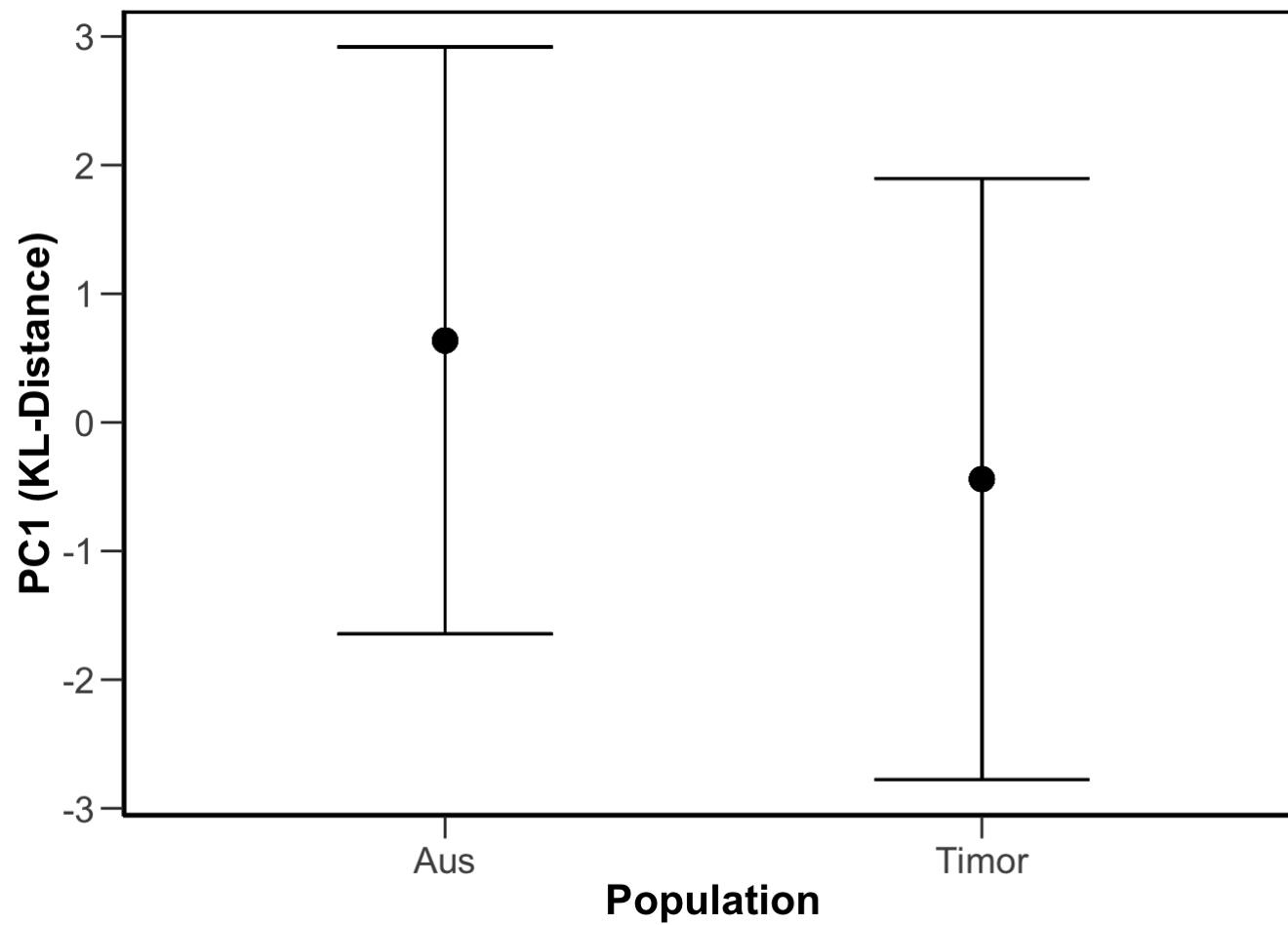
```
anova(mKLtoTutor, mKLtoTutorNull)
```

```
## Data: KLtoTutors
## Models:
## mKLtoTutorNull: PC1 ~ 1 + (1 | Tutor)
## mKLtoTutor: PC1 ~ Population + (1 | Tutor)
##          Df    AIC    BIC  logLik deviance Chisq Chi Df Pr(>Chisq)
## mKLtoTutorNull 3 86.780 89.451 -40.390    80.780
## mKLtoTutor     4 87.369 90.931 -39.685    79.369 1.411      1   0.2349
```

as predicted, no difference here. but lets plot anyway.

```
KLtoTutors$predicted=(predict(mKLtoTutor,re.form=NA))
pframe <- data.frame(Population=KLtoTutors$Population)
KLtoTutors$LCI=as.vector((easyPredCInorm(mKLtoTutor,pframe))[,1])
KLtoTutors$UCI=as.vector((easyPredCInorm(mKLtoTutor,pframe))[,2])

ggplot(KLtoTutors,aes(x=Population,y=predicted))+geom_point(size=4)+geom_errorbar(aes(ymin=LCI,ymax=UCI),
width=0.4,
size=0.5)+
ylab("PC1 (KL-Distance)")+xlab("Population") +theme_bw() +
theme(axis.title=element_text(vjust=1,size=16,face="bold"), axis.text=element_text(size=14),
axis.text.x = element_text(vjust=0.65, hjust=0.5, size=14), panel.border =
element_rect(colour = "black",size=1.25)) + theme(axis.ticks.length=unit(0.3,"cm"))+
theme(panel.grid.major = element_blank(), panel.grid.minor =element_blank(), axis.line =
element_line(colour = "black"))
```



```
#ggsave("logPC1-KLxFosterToTutors.pdf", plot=last_plot(), device=NULL, path=NULL, scale=1, width=NA, height=NA, dpi=300, limitsize=TRUE)
```

so the last thing we need to do is examine variability within the sets of birds that have been cross fostered.

```
path<-("~/Documents/Projects/Domestication/"
KLxfost=read.table(paste(path,"logXfostPairwiseWithinPops.txt",sep="/"),sep = "\t",header=TRUE)
str(KLxfost)
```

```

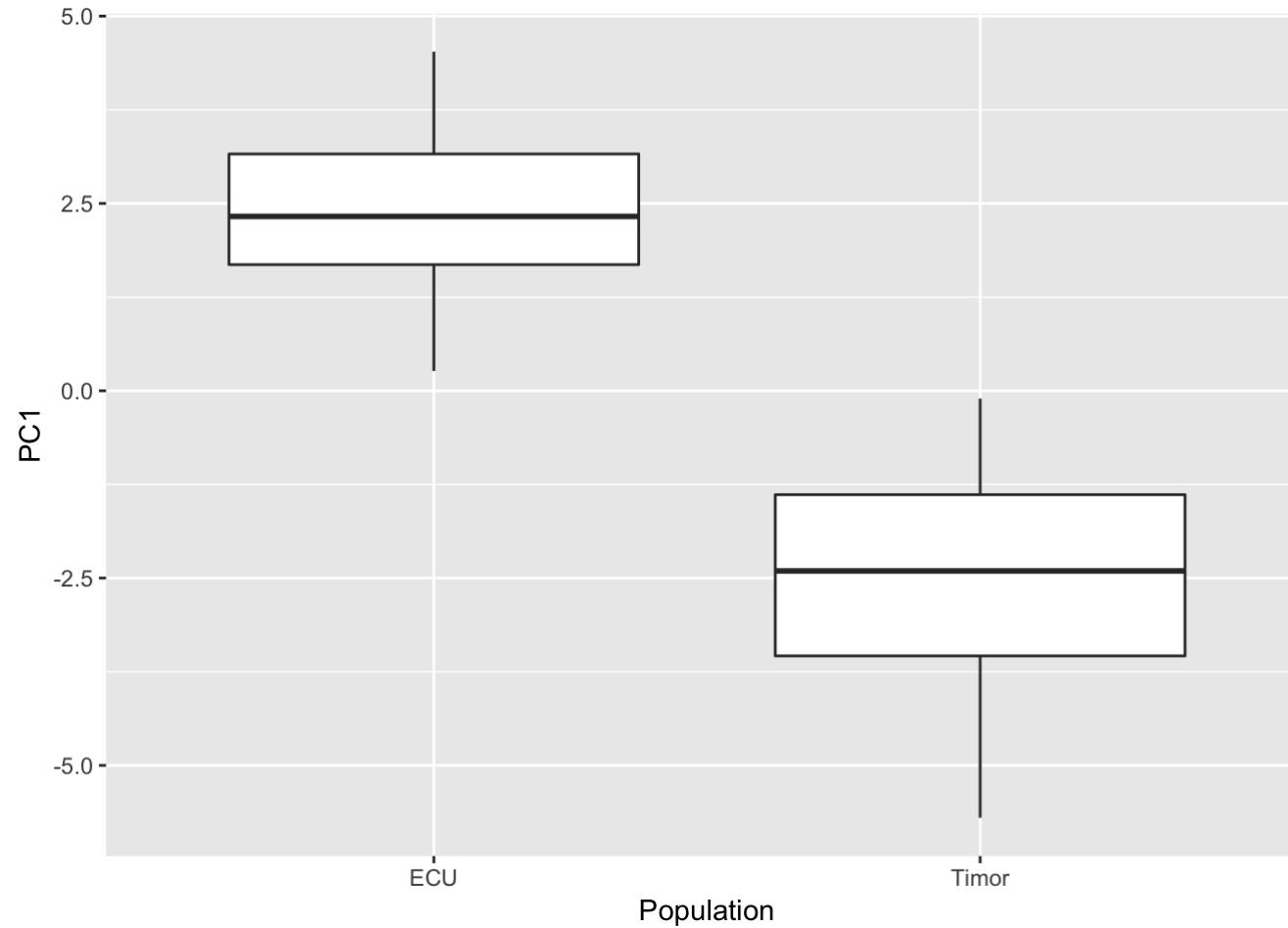
## 'data.frame': 20 obs. of 28 variables:
## $ X : int 1 2 3 4 5 6 7 8 9 10 ...
## $ logamplitude : num -0.157 0.324 0.382 0.15 0.351 ...
## $ logpitch : num 0.0853 0.0279 0.0741 0.1064 0.1239 ...
## $ logFM : num -0.15 0.32 0.117 0.108 0.565 ...
## $ logAM2 : num -0.115 -0.164 0.318 -0.289 0.2 ...
## $ logentropy : num -0.2459 -0.0596 0.0853 -0.1641 0.0292 ...
## $ logpitchgoodness : num -0.0742 0.3476 0.0979 -0.0919 0.3875 ...
## $ logmeanfreq : num -0.3064 -0.186 0.0188 -0.0168 0.2168 ...
## $ logvpitch : num -1.287 -0.846 -0.281 -0.799 -0.639 ...
## $ logvFM : num -0.5164 -0.2281 -0.1063 -0.2109 -0.0626 ...
## $ logventropy : num -0.1112 -0.01312 -0.07199 -0.00938 0.03991 ...
## $ logvpitchgoodness: num -0.5112 -0.0907 -0.1372 -0.2589 -0.0846 ...
## $ vmeanfreq : num -0.106 -0.392 -0.009 -0.274 0.315 ...
## $ PC1 : num -5.699 -3.149 -1.661 -4 -0.331 ...
## $ PC2 : num 0.606 1.507 -1.114 0.483 0.456 ...
## $ PC3 : num -0.598 1.398 -0.178 0.706 -1.238 ...
## $ PC4 : num 1.52 -1.284 -0.674 0.439 -1.433 ...
## $ PC5 : num 0.703 -0.474 -0.259 -0.648 1.176 ...
## $ PC6 : num 0.56 0.459 0.516 -0.567 -0.736 ...
## $ PC7 : num 0.357 0.109 0.598 0.191 -0.282 ...
## $ PC8 : num 0.4017 -0.2142 -0.0457 -0.4466 0.1006 ...
## $ PC9 : num 0.0812 -0.0111 0.5832 -0.2039 0.054 ...
## $ PC10 : num 0.0902 -0.1312 -0.0009 -0.317 -0.0206 ...
## $ PC11 : num -0.1156 -0.2461 0.2461 0.0657 -0.2288 ...
## $ PC12 : num -0.0774 0.1448 -0.1109 -0.1328 -0.0748 ...
## $ Population : Factor w/ 2 levels "ECU","Timor": 2 2 2 2 2 2 2 2 2 ...
## $ Template : Factor w/ 8 levels "filtered_BfAPU187.xls",...: 5 6 7 8 6 7 8 7 8 ...
## $ Target : Factor w/ 8 levels "filtered_BfADP56.xls",...: 5 5 5 5 6 6 7 7 8 ...

```

```

require(ggplot2)
print(ggplot(KLxfost,aes(x=Population,y=PC1))+geom_boxplot())

```



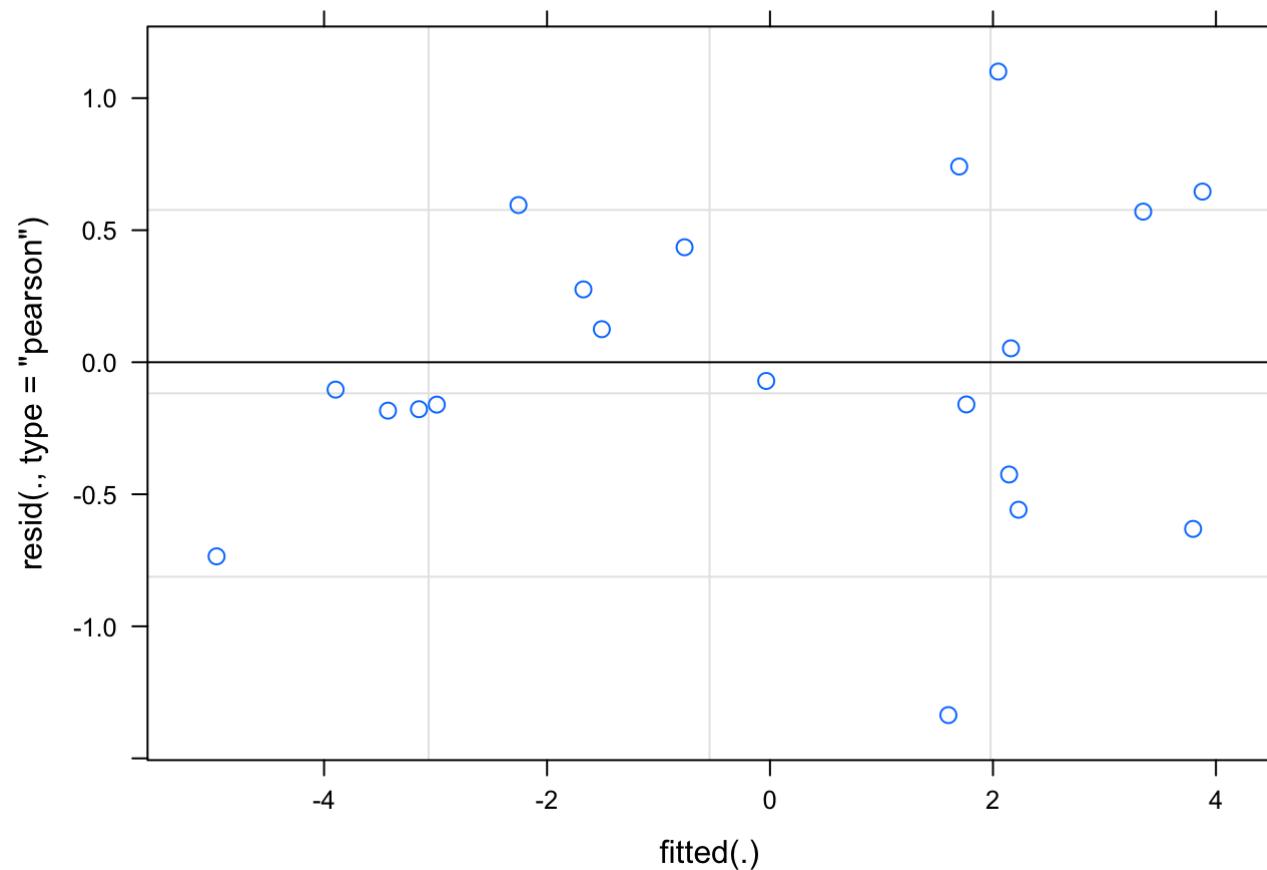
So here again we see a very similar

pattern as was described for the non cross fostered birds.

```
mPC1xFost <- blmer(PC1 ~ Population + (1 | Template) + (1 | Target) , data = KLxfost, REML = FALSE)
VarCorr(mPC1xFost) # Check for singularity
```

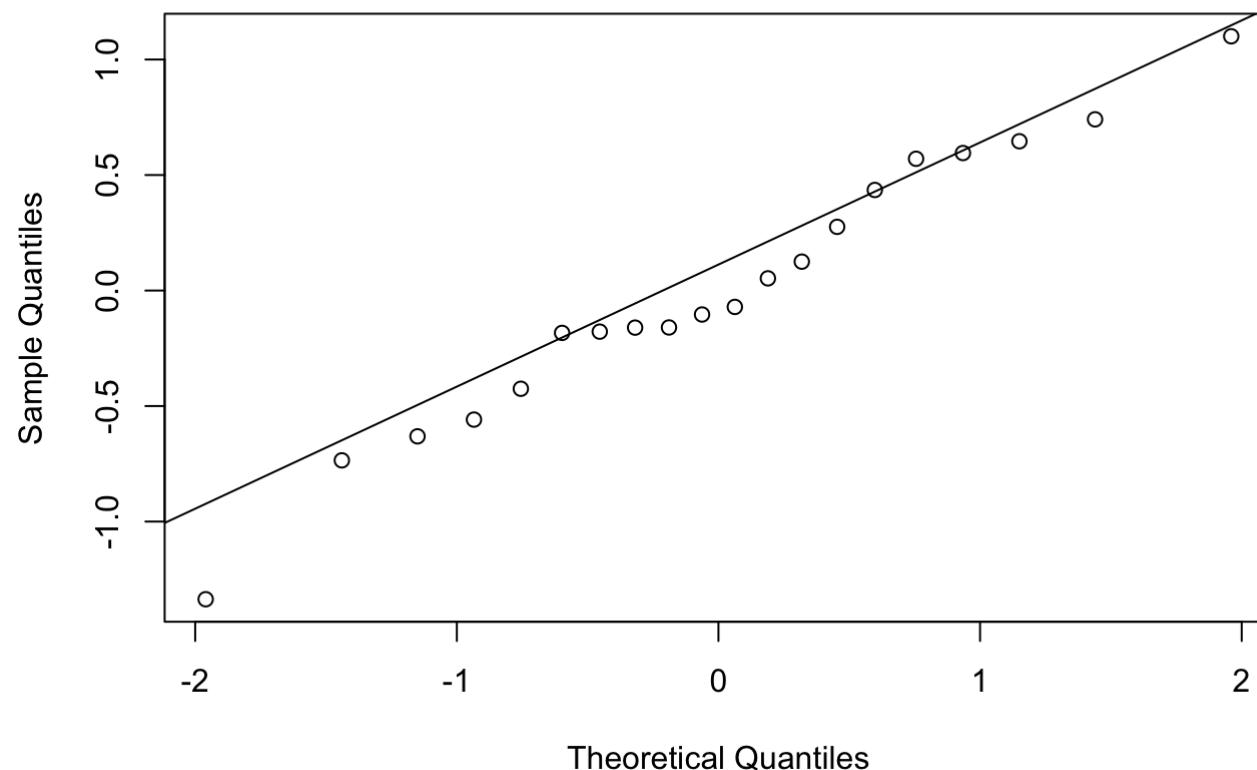
```
## Groups      Name       Std.Dev.
## Template (Intercept) 1.03509
## Target    (Intercept) 1.07980
## Residual             0.74016
```

```
plot(mPC1xFost) # residuals plot
```



```
qqnorm(resid(mPC1xFost)); qqline(resid(mPC1xFost)) #qqplot
```

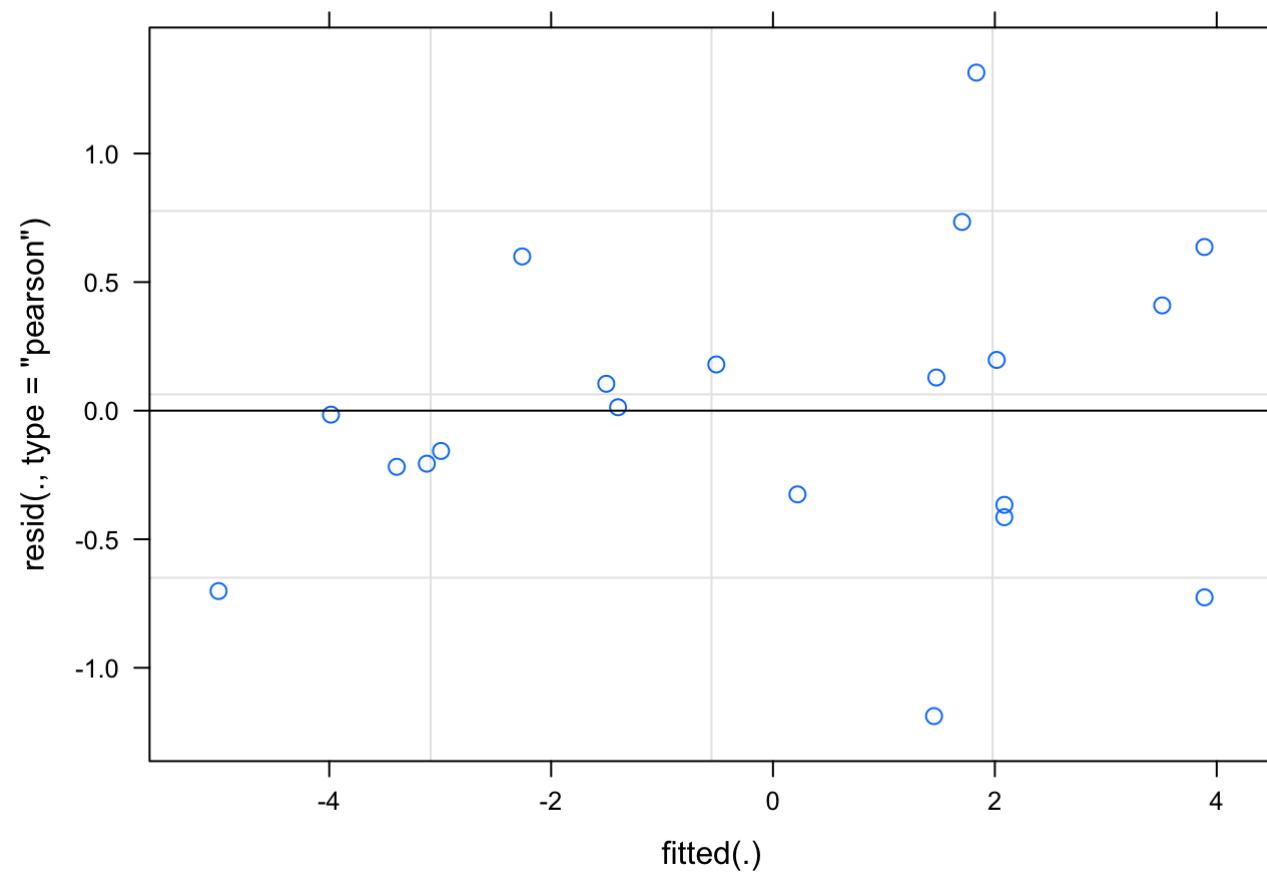
Normal Q-Q Plot



```
mPC1xFostNull <- blmer(PC1 ~ 1 + (1 | Template) + (1 | Target), data = KLxfost, REML = FALSE)
VarCorr(mPC1xFostNull) # Check for singularity
```

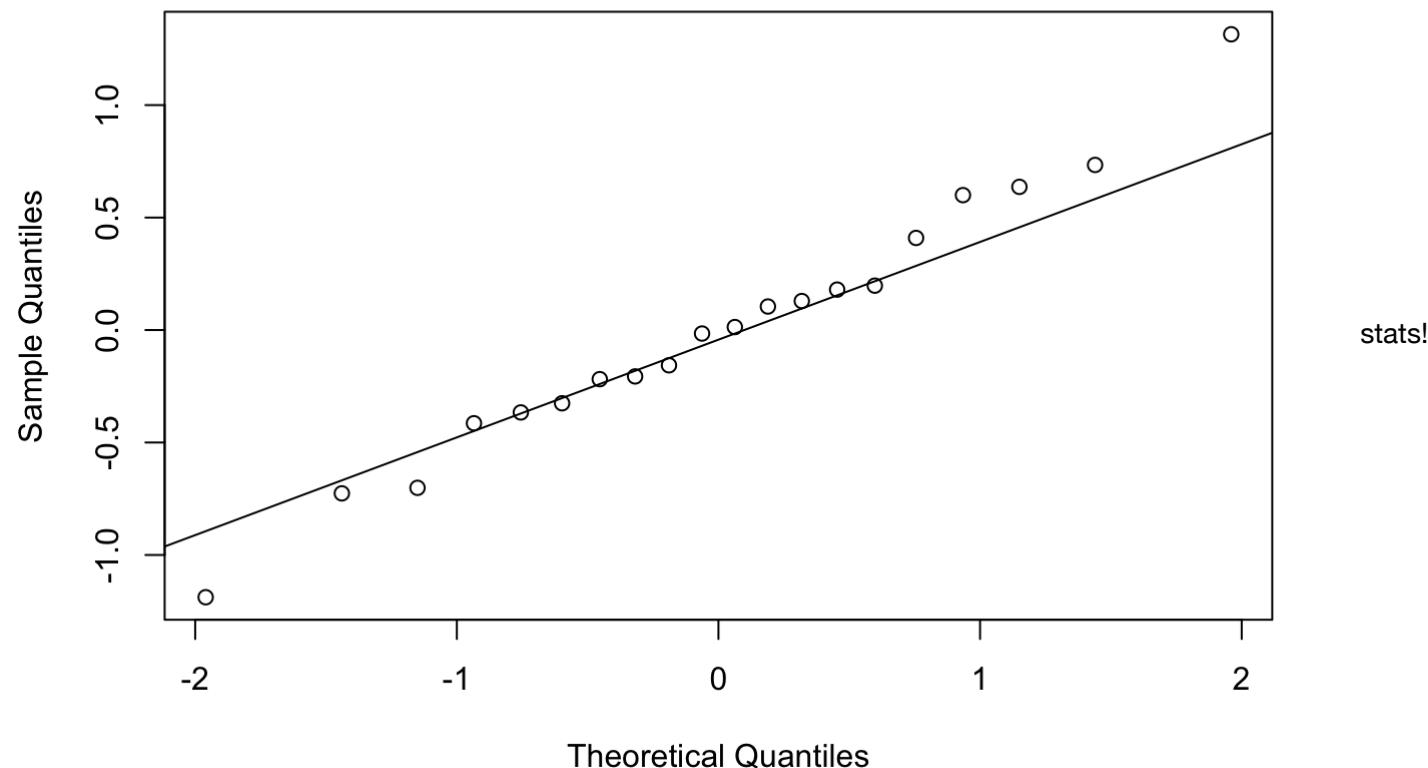
```
## Groups      Name       Std.Dev.
## Template (Intercept) 1.2867
## Target    (Intercept) 2.5255
## Residual             0.7735
```

```
plot(mPC1xFostNull) # residuals plot
```



```
qqnorm(resid(mPC1xFostNull)); qqline(resid(mPC1xFostNull)) #qqplot
```

Normal Q-Q Plot



stats!

```
anova(mPC1xFostNull, mPC1xFost)
```

```

## Data: KLxfost
## Models:
## mPC1xFostNull: PC1 ~ 1 + (1 | Template) + (1 | Target)
## mPC1xFost: PC1 ~ Population + (1 | Template) + (1 | Target)
##          Df      AIC     BIC   logLik deviance Chisq Chi Df Pr(>Chisq)
## mPC1xFostNull 4 91.535 95.518 -41.767    83.535
## mPC1xFost     5 79.379 84.358 -34.690    69.379 14.155       1 0.0001683
##
## mPC1xFostNull
## mPC1xFost      ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

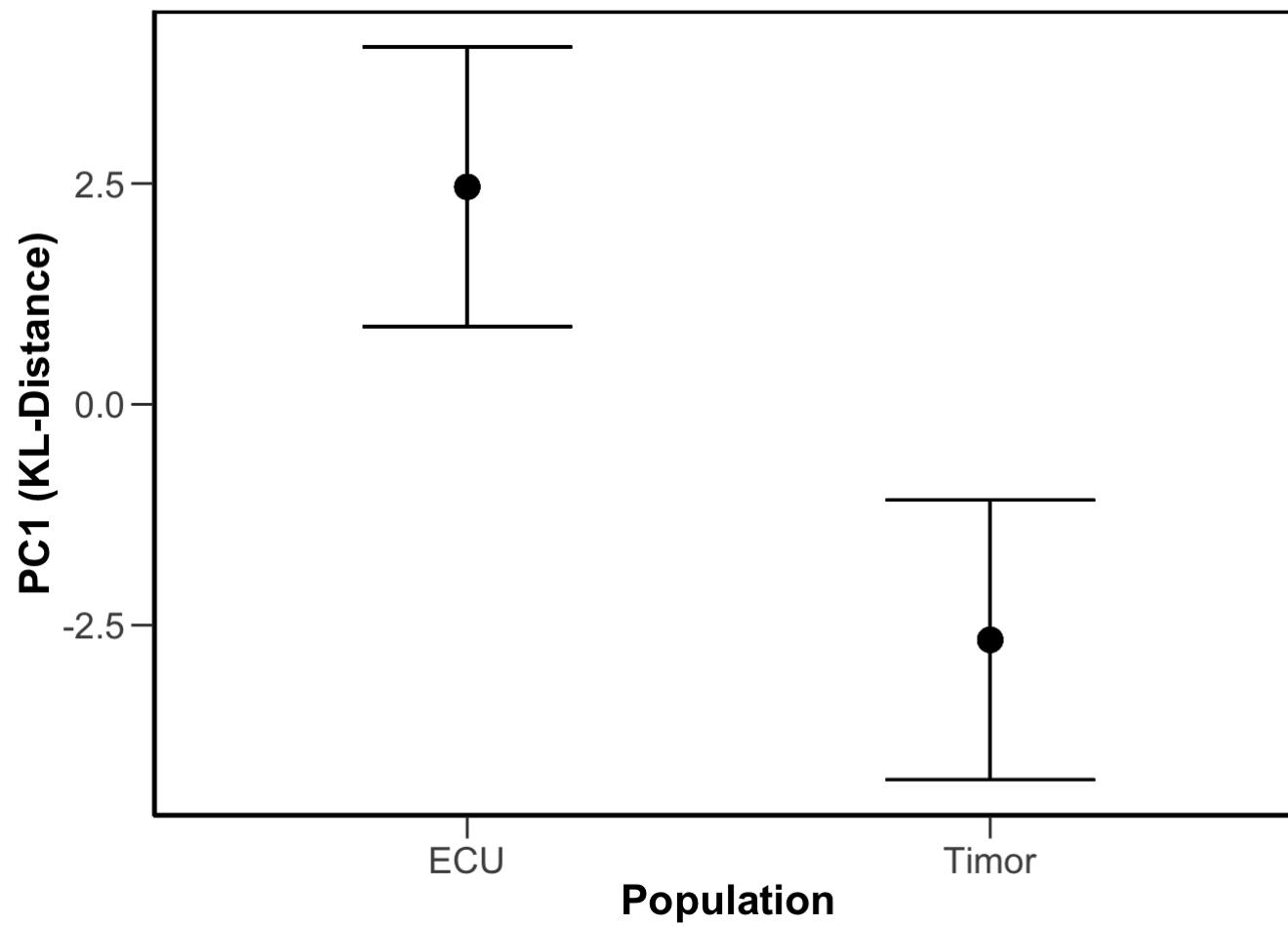
```

```

KLxfost$predicted=(predict(mPC1xFost,re.form=NA))
pframe <- data.frame(Population=KLxfost$Population)
KLxfost$LCI=as.vector((easyPredCInorm(mPC1xFost,pframe))[,1])
KLxfost$UCI=as.vector((easyPredCInorm(mPC1xFost,pframe))[,2])

ggplot(KLxfost,aes(x=Population,y=predicted))+geom_point(size=4)+geom_errorbar(aes(ymin=LCI,ymax=UCI),
width=0.4,
size=0.5)+
ylab("PC1 (KL-Distance)")+xlab("Population")+theme_bw() +
theme(axis.title=element_text(vjust=1,size=16,face="bold"), axis.text=element_text(size=14),
axis.text.x = element_text(vjust=0.65, hjust=0.5, size=14), panel.border =
element_rect(colour = "black",size=1.25)) + theme(axis.ticks.length=unit(0.3,"cm"))+
theme(panel.grid.major = element_blank(), panel.grid.minor =element_blank(), axis.line =
element_line(colour = "black"))

```



```
#ggsave( "PC1-KLxFosterWithin.pdf", plot=last_plot(), device=NULL, path=NULL, scale=1, width=NA, height=NA, dpi=300, limitsize=TRUE)
```