R package glmm

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1 Bacteria example

The MASS package contains the command glmmPQL and the bacteria data-set. The manual describes the data set as follows: "Tests of the presence of the bacteria H. influenzae in children with otitis media in the Northern Territory of Australia."

The data were fit using glmmPQL, glmm with $m = 10^3$, glmm with $m = 10^4$, glmm with $m = 10^5$. (The data did need to be reformatted, which took only a couple minutes). The parameter estimates are summarized in the following table. More model details can be seen in the output that follows the table.

	Intercept	$\operatorname{trtdrug}$	$\operatorname{trtdrug} +$	I(week > 2) TRUE	ν
glmmPQL	3.41	-1.25	75	-1.61	1.99
glmm $m = 10^3$	3.02	-1.20	89	-1.44	.81
glmm $m = 10^4$	3.65	-1.36	92	-1.66	2.02
glmm $m = 10^5$	3.49	-1.65	90	-1.46	.90

It's safe to say that the bacteria glmm results with a paltry Monte Carlo sample size of $m = 10^3$ are not reliable. We conclude this because the estimates change quite a bit when $m = 10^4$. The results using $m = 10^4$ are very similar to the glmmPQL results.

The glmmPQL results:

```
> bac.pql<-glmmPQL(y ~ trt + I(week > 2), random = ~ 1 | ID,
+ family = binomial, data = bacteria)
> summary(bac.pql)
Linear mixed-effects model fit by maximum likelihood
Data: bacteria
AIC BIC logLik
NA NA NA
Random effects:
```

Variance function:

StdDev:

Formula: ~1 | ID

(Intercept) Residual

1.410637 0.7800511

```
Structure: fixed weights
 Formula: ~invwt
Fixed effects: y ~ trt + I(week > 2)
                    Value Std.Error DF t-value p-value
(Intercept)
                 3.412014 0.5185033 169 6.580506 0.0000
trtdrug
               -1.247355 0.6440635 47 -1.936696 0.0588
               -0.754327 0.6453978 47 -1.168779 0.2484
trtdrug+
I(week > 2)TRUE -1.607257 0.3583379 169 -4.485311 0.0000
 Correlation:
                (Intr) trtdrg trtdr+
trtdrug
                -0.598
                -0.571 0.460
trtdrug+
I(week > 2)TRUE -0.537 0.047 -0.001
Standardized Within-Group Residuals:
       Min
                   Q1
                             Med
                                         QЗ
                                                   Max
-5.1985361 0.1572336 0.3513075 0.4949482 1.7448845
Number of Observations: 220
Number of Groups: 50
   The bacteria glmm results with a Monte Carlo sample size of m = 10^3:
> set.seed(1234)
> bac.glmm1<-glmm(y2~trt+I(week > 2),list(~0+ID),
family=bernoulli.glmm, data=bacteria, m=10^3, varcomps.names=c("ID"))
> summary(bac.glmm1)
glmm(fixed = y2 ~ trt + I(week > 2), random = list(~0 + ID), varcomps.names = c("ID"),
data = bacteria, family.glmm = bernoulli.glmm,
                                                  m = 10^3
Fixed Effects:
               Estimate Std. Error z value Pr(>|z|)
(Intercept)
                3.0199
                             0.4667 6.471 9.75e-11 ***
trtdrug
                 -1.1989
                             0.4473 -2.680 0.007357 **
                            0.4742 -1.873 0.061021 .
trtdrug+
                 -0.8883
                            0.4258 -3.384 0.000716 ***
I(week > 2)TRUE -1.4407
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
```

Variance Components for Random Effects (P-values are one-tailed):

```
Estimate Std. Error z value Pr(>|z|)/2
    0.8175
                0.1635 4.998
                                 2.89e-07 ***
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1
The results from fitting the bacteria dataset with glmm and m = 10^4:
> set.seed(1234)
> bac.glmm2<-glmm(y2~trt+I(week > 2),list(~0+ID), family=bernoulli.glmm, data=bacteria, m=10^4, varcomps
> summary(bac.glmm2)
Call:
glmm(fixed = y2 ~ trt + I(week > 2), random = list(~0 + ID),
    varcomps.names = c("ID"), data = bacteria, family.glmm = bernoulli.glmm,
    m = 10^4)
Fixed Effects:
               Estimate Std. Error z value Pr(>|z|)
(Intercept)
                 3.6514
                             0.6327 5.771 7.9e-09 ***
                             0.7461 -1.829 0.067427 .
trtdrug
                 -1.3645
trtdrug+
                 -0.9186
                             0.7619 -1.206 0.227915
I(week > 2)TRUE -1.6660
                             0.4645 -3.587 0.000334 ***
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
Variance Components for Random Effects (P-values are one-tailed):
   Estimate Std. Error z value Pr(>|z|)/2
ID
     2.0244
               0.7515 2.694
                                  0.00353 **
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1
   Results of glmm with m = 10^5:
> set.seed(1234)
> bac.glmm3<-glmm(y2~trt+I(week > 2),list(~0+ID), family=bernoulli.glmm, data=bacteria, m=10^5, varcomps
> summary(bac.glmm3)
Call:
glmm(fixed = y2 ~ trt + I(week > 2), random = list(~0 + ID),
```

varcomps.names = c("ID"), data = bacteria, family.glmm = bernoulli.glmm,

```
m = 10^5)
```

```
Fixed Effects:
                Estimate Std. Error z value Pr(>|z|)
(Intercept)
                  3.4888
                              0.4861
                                       7.178 7.08e-13 ***
trtdrug
                 -1.6526
                              0.5232
                                      -3.159 0.001586 **
trtdrug+
                 -0.8968
                              0.5160
                                     -1.738 0.082196 .
I(week > 2)TRUE
                -1.4614
                              0.4334
                                      -3.372 0.000746 ***
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
Variance Components for Random Effects (P-values are one-tailed):
   Estimate Std. Error z value Pr(>|z|)/2
     0.8999
                0.2471
                                  0.000136 ***
ID
                         3.641
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
```

2 Herd CBPP Example

The cbpp dataset is located in the lme4 package. The lme4 package describes it thusly: "Contagious bovine pleuropneumonia (CBPP) is a major disease of cattle in Africa, caused by a mycoplasma. This dataset describes the serological incidence of CBPP in zebu cattle during a follow-up survey implemented in 15 commercial herds located in the Boji district of Ethiopia. The goal of the survey was to study the within-herd spread of CBPP in newly infected herds. Blood samples were quarterly collected from all animals of these herds to determine their CBPP status. These data were used to compute the serological incidence of CBPP (new cases occurring during a given time period). Some data are missing (lost to follow-up)."

First, I fit the data using glmer in lme4. Then I fit the data using glmm with $m = 10^4$. This took 16.75 minutes on my netbook. The point estimates and the standard errors were very similar between the two methods of model-fitting. (The data did need to be reformatted, which took only a couple minutes)

First, the results from glmer:

```
> summary(gm1)
Generalized linear mixed model fit by maximum likelihood (Laplace
   Approximation) [glmerMod]
Family: binomial ( logit )
Formula: cbind(incidence, size - incidence) ~ period + (1 | herd)
   Data: cbpp

AIC    BIC   logLik deviance df.resid
   194.1   204.2   -92.0   184.1   51
```

```
Scaled residuals:
```

```
Min 1Q Median 3Q Max
-2.3816 -0.7889 -0.2026 0.5142 2.8791
```

Random effects:

Groups Name Variance Std.Dev. herd (Intercept) 0.4123 0.6421 Number of obs: 56, groups: herd, 15

Fixed effects:

Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1

Correlation of Fixed Effects:

(Intr) perid2 perid3

period2 -0.363

period3 -0.340 0.280

period4 -0.260 0.213 0.198

Next, the results using glmm with $m = 10^4$:

> summary(herd.glmm1)

Call:

```
glmm(fixed = Y ~ period, random = list(~0 + herd), varcomps.names = c("herd"),
    data = herddat, family.glmm = bernoulli.glmm, m = 10^4)
```

Fixed Effects:

Estimate Std. Error z value Pr(>|z|)

Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1

```
Variance Components for Random Effects (P-values are one-tailed):
     Estimate Std. Error z value Pr(>|z|)/2
herd
       0.4354
                  0.2488
                            1.75
                                     0.0401 *
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
    Salamander
3
> set.seed(1234)
> sal.glmm3<-glmm(Mate~Cross,random=list(~0+Female,~0+Male),varcomps.names=c("F","M"),data=salamander,fa
> summary(sal.glmm3)
Call:
glmm(fixed = Mate ~ Cross, random = list(~0 + Female, ~0 + Male),
    varcomps.names = c("F", "M"), data = salamander, family.glmm = bernoulli.glmm,
    m = 10^4)
Fixed Effects:
            Estimate Std. Error z value Pr(>|z|)
(Intercept)
              1.1123
                         0.2593 4.290 1.79e-05 ***
CrossRW
             -0.5545
                        0.3558 -1.558
                                           0.119
CrossWR.
             -3.1701
                       0.4029 -7.867 3.62e-15 ***
CrossWW
             -0.2122
                         0.3681 -0.577
                                           0.564
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1
Variance Components for Random Effects (P-values are one-tailed):
  Estimate Std. Error z value Pr(>|z|)/2
   1.2087
               0.2534
                        4.769
                                9.24e-07 ***
    0.9485
               0.1809
                        5.244
                              7.86e-08 ***
М
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1
   Here are the results with m = 10^4 but I forgot to set the seed.
> summary(sal.glmm2)
Call:
glmm(fixed = Mate ~ Cross, random = list(~0 + Female, ~0 + Male),
    varcomps.names = c("F", "M"), data = salamander, family.glmm = bernoulli.glmm,
    m = 10^4)
```

Fixed Effects:

```
Estimate Std. Error z value Pr(>|z|)
(Intercept) 0.93879
                        0.26963
                                  3.482 0.000498 ***
CrossRW
            -0.76157
                        0.37376 -2.038 0.041593 *
CrossWR
            -2.72491
                        0.40736 -6.689 2.24e-11 ***
            0.02256
                        0.37902
                                  0.060 0.952541
CrossWW
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
Variance Components for Random Effects (P-values are one-tailed):
  Estimate Std. Error z value Pr(>|z|)/2
    1.5991
               0.2965
                        5.394
                              3.45e-08 ***
  0.9908
               0.1885
                        5.255
                              7.41e-08 ***
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
```

4 Reformatting the Datsets

4.1 Bacteria Reformatting

The issue with the bacteria dataset is the response was y/n rather than 1/0. I created a new response that changed the y to 1 and the n to 0.

bacteria\$y2<-as.integer(bacteria\$y)-1

4.2 CBPP Reformatting

The cbpp dataset was created for binomial but my package was written for Bernoulli responses. In other words, my package needs a row for each success or failure. I did this in the following way:

```
cbpp$nonincidence<-cbpp$size-cbpp$incidence #number of "failures"
herddat<-matrix(data=NA,nrow=842,ncol=3)
colnames(herddat)<-c("Y","period","herd")
rowid<-1
for(i in 1:nrow(cbpp)){
  #make a row for each one of the incidences
  ntimes<-cbpp[i,2]
  if(ntimes>0){
  for(j in 1:ntimes){
   herddat[rowid,]<-c(1,cbpp[i,4],cbpp[i,1])
  rowid<-rowid+1
}</pre>
```

```
#make a row for each of the nonincidences
ntimes<-cbpp[i,5]
if(ntimes>0){
for(j in 1:ntimes){
herddat[rowid,]<-c(0,cbpp[i,4],cbpp[i,1])
rowid<-rowid+1
}
}
herddat<-as.data.frame(herddat)
herddat$herd<-as.factor(herddat$herd)
herddat$period<-as.factor(herddat$period)</pre>
```