# Monday, Mar 27

## Distributions for Over-dispersion

One way to model over-dispersion is to assume a model of the form

$$g[E(Y_i)] = \beta_0 + \beta_1 x_{i1} + \dots + \beta_k x_{ik} + \zeta_i,$$

where  $\zeta_i$  is an *unobserved* unit-specific random quantity that represents one or more unobserved explanatory variables that vary over units.

#### The Negative Binomial Distribution

Suppose that  $Y_i$  has a Poisson distribution conditional on  $\zeta_i$ , and  $e^{\zeta_i}$  has a gamma distribution such that  $E(e^{\gamma_i}) = 1$  and  $Var(e^{\gamma_i}) = \alpha > 0$ . The marginal distribution of  $Y_i$  is then a negative binomial distribution, with mean structure

$$g[E(Y_i)] = \eta_i,$$

and variance structure

$$Var(Y_i) = E(Y_i) + \alpha E(Y_i)^2 \ge E(Y_i).$$

The Poisson distribution is a special case where  $\alpha = 0$ . This variance structure does not have the form

$$Var(Y_i) = \phi V[E(Y_i)]$$

unless  $\alpha$  is known (which it normally is not), so this model is not a traditional GLM. But we can make inferences using maximum likelihood.

**Example:** Consider our model for the trawl fishing data. Here we will consider a negative binomial regression model.

```
library(COUNT)
data(fishing)

library(MASS) # for the glm.nb function (note there is no family argument)

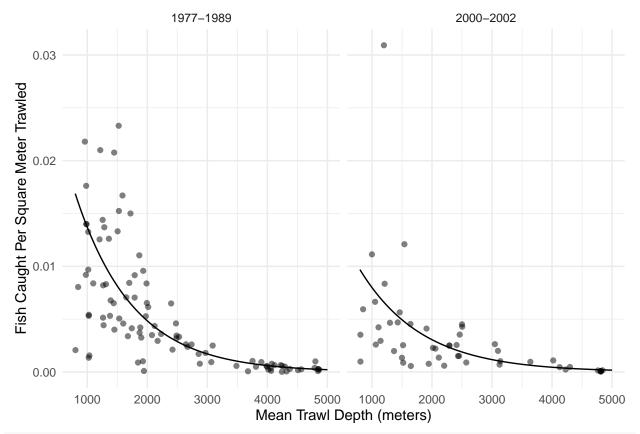
m <- glm.nb(totabund ~ period * meandepth + offset(log(sweptarea)),
    link = log, data = fishing)

d <- expand.grid(sweptarea = 1, period = levels(fishing$period),
    meandepth = seq(800, 5000, length = 100))

d$yhat <- predict(m, newdata = d, type = "response")

p <- ggplot(fishing, aes(x = meandepth, y = totabund/sweptarea)) +
    geom_point(alpha = 0.5) + facet_wrap(~ period) + theme_minimal() +
    labs(x = "Mean Trawl Depth (meters)",
        y = "Fish Caught Per Square Meter Trawled") +
    geom_line(aes(y = yhat), data = d)

plot(p)</pre>
```



summary(m) # note that what glm.nb calls theta equals 1/alpha

```
Call:
```

glm.nb(formula = totabund ~ period \* meandepth + offset(log(sweptarea)),
 data = fishing, link = log, init.theta = 1.961162176)

#### Deviance Residuals:

Min 1Q Median 3Q Max -3.366 -0.830 -0.208 0.365 2.893

#### Coefficients:

Estimate Std. Error z value Pr(>|z|)(Intercept) -3.25e+00 1.59e-01 -20.40 <2e-16 \*\*\* period2000-2002 -6.19e-01 0.023 \* 2.73e-01 -2.27-1.04e-03 5.92e-05 -17.58 <2e-16 \*\*\* meandepth period2000-2002:meandepth 7.95e-05 1.01e-04 0.79 0.432

---

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

(Dispersion parameter for Negative Binomial(1.961) family taken to be 1)

Null deviance: 471.79 on 146 degrees of freedom Residual deviance: 159.31 on 143 degrees of freedom

AIC: 1763

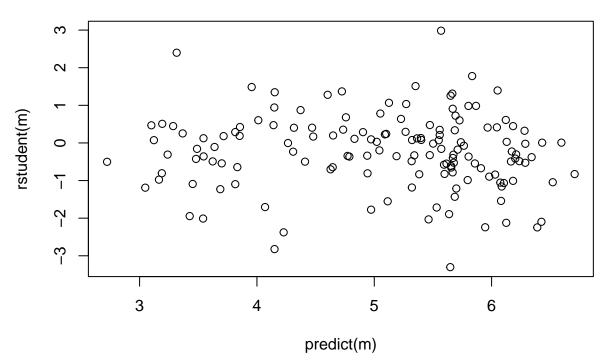
Number of Fisher Scoring iterations: 1

Theta: 1.961 Std. Err.: 0.219

2 x log-likelihood: -1752.713

```
plot(predict(m), rstudent(m), main = "Residual Plot")
```

# **Residual Plot**



Interestingly inferences based on the negative binomial model are very similar to those obtained using quasilikelihood assuming the variance structure  $V(Y_i) = \phi E(Y_i)^2$ . Here are the parameter estimates, standard errors, and confidence intervals.

```
m.negbn <- glm.nb(totabund ~ period * meandepth + offset(log(sweptarea)),
    link = log, data = fishing)
m.quasi <- glm(totabund ~ period * meandepth + offset(log(sweptarea)),
    family = quasi(link = "log", variance = "mu^2"), data = fishing)
cbind(summary(m.negbn)$coefficients, confint(m.negbn))</pre>
```

```
Estimate Std. Error z value Pr(>|z|) 2.5 % 97.5 % (Intercept) -3.249e+00 1.592e-01 -20.4044 1.529e-92 -3.5603979 -2.9274102 period2000-2002 -6.187e-01 2.726e-01 -2.2695 2.324e-02 -1.1815413 -0.0436165 meandepth -1.041e-03 5.923e-05 -17.5844 3.245e-69 -0.0011578 -0.0009213 period2000-2002:meandepth 7.955e-05 1.013e-04 0.7855 4.322e-01 -0.0001333 0.0002952 cbind(summary(m.quasi)$coefficients, confint(m.quasi))
```

Here are the estimates of the rate ratios for period at several different depths.

```
library(trtools)
contrast(m.negbn,
 a = list(meandepth = c(1000, 2000, 3000, 4000, 5000), period = "2000-2002", sweptarea = 1),
  b = list(meandepth = c(1000, 2000, 3000, 4000, 5000), period = "1977-1989", sweptarea = 1),
 cnames = c("1000m","2000m","3000m","4000m","5000m"), tf = exp)
      estimate lower upper
1000m
        0.5832 0.4017 0.8468
2000m
        0.6315 0.4868 0.8193
3000m
        0.6838 0.5184 0.9020
4000m 0.7404 0.4928 1.1125
5000m
        0.8017 0.4494 1.4301
contrast(m.quasi,
 a = list(meandepth = c(1000, 2000, 3000, 4000, 5000), period = "2000-2002", sweptarea = 1),
 b = list(meandepth = c(1000, 2000, 3000, 4000, 5000), period = "1977-1989", sweptarea = 1),
 cnames = c("1000m","2000m","3000m","4000m","5000m"), tf = exp)
      estimate lower upper
1000m
        0.5878 0.4046 0.8540
2000m
        0.6321 0.4869 0.8206
3000m
        0.6798 0.5173 0.8935
        0.7311 0.4905 1.0897
4000m
5000m
        0.7863 0.4458 1.3867
Here are the tests (likelihood ratio and F) for the "effect" of period. The null model assumes that expected
abundance per unit area trawled is the same each period at a given depth. Put another way, the null model
assumes that the rate ratio for period is one for all depths.
m.negbn.null <- glm.nb(totabund ~ meandepth + offset(log(sweptarea)),</pre>
 link = log, data = fishing)
anova(m.negbn.null, m.negbn)
Likelihood ratio tests of Negative Binomial Models
Response: totabund
                                         Model theta Resid. df
                                                                   2 x log-lik.
                                                                                           df LR stat.
                                                                                   Test.
           meandepth + offset(log(sweptarea)) 1.832
                                                            145
                                                                           -1764
2 period * meandepth + offset(log(sweptarea)) 1.961
                                                            143
                                                                          -1753 1 vs 2
                                                                                            2
                                                                                                  11.11
  Pr(Chi)
2 0.003872
m.quasi.null <- glm(totabund ~ meandepth + offset(log(sweptarea)),</pre>
  family = quasi(link = "log", variance = "mu^2"), data = fishing)
anova(m.quasi.null, m.quasi, test = "F")
Analysis of Deviance Table
Model 1: totabund ~ meandepth + offset(log(sweptarea))
Model 2: totabund ~ period * meandepth + offset(log(sweptarea))
  Resid. Df Resid. Dev Df Deviance
                                       F Pr(>F)
        145
                  90.5
1
2
        143
                  84.5 2
                               5.94 5.74 0.004 **
Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

Note: When using anova for a negative binomial model (estimated using the glm.nb function) we omit the test = "LRT" option which we use for generalized linear models. Somewhat confusingly, the anova function will do a likelihood ratio test for a glm.nb object, but will throw an error if we try to change the test type (even if we ask for a likelihood ratio test).

## Heteroscedastic Consistent (Robust) Standard Errors

An alternative is to accept that the specified variance structure is incorrect and estimate standard errors in a way that provides *consistent* estimates despite the misspecification of the variance structure.<sup>1</sup>

Note: I needed to specify the data set as trtools::rotifer below as there is a data set of the same name in another package that was loaded earlier. It's actually the same data but in a different format from the data frame in the trtools package.

Example: Consider the logistic regression model for the rotifer data from the trtools package.

```
m <- glm(cbind(y, total - y) ~ species + density + species:density,
    family = binomial, data = trtools::rotifer)</pre>
```

Here are the parameter estimates and standard errors, with and without using the robust standard error estimates.

```
library(sandwich) # for the vcovHC function
library(lmtest) # for coeftest and coefci functions
cbind(summary(m)$coefficients, confint(m))
```

```
Estimate Std. Error z value
                                                Pr(>|z|)
                                                             2.5 %
                                                                     97.5 %
(Intercept)
                  -114.352
                               4.034 -28.3454 9.534e-177 -122.420 -106.598
speciespm
                     4.629
                                6.598
                                       0.7016 4.830e-01
                                                            -8.464
                                                                     17.431
density
                   108.746
                               3.857 28.1910 7.535e-175 101.332 116.460
speciespm:density
                   -3.077
                                6.329 -0.4862 6.268e-01
                                                          -15.354
                                                                      9.487
cbind(coeftest(m, vcov = vcovHC), coefci(m, vcov = vcovHC))
```

```
Estimate Std. Error z value Pr(>|z|)
                                                         2.5 % 97.5 %
(Intercept)
                  -114.352
                                18.31 -6.2449 4.240e-10 -150.24 -78.46
speciespm
                     4.629
                                29.91 0.1548 8.770e-01 -53.99 63.25
density
                   108.746
                               17.50 6.2137 5.175e-10
                                                         74.44 143.05
speciespm:density
                                28.82 -0.1068 9.150e-01
                                                        -59.57 53.42
                   -3.077
```

An alternative to using coeftest and coefci is lincon(m, fcov = vcovHC). Now compare our inferences for the odds ratios for the effect of a 0.01 increase in density.

```
contrast(m,
  a = list(density = 0.02, species = c("kc","pm")),
  b = list(density = 0.01, species = c("kc","pm")),
  cnames = c("kc","pm"), tf = exp)
```

```
estimate lower upper
kc    2.967 2.751 3.200
pm    2.877 2.607 3.174

contrast(m,
    a = list(density = 0.02, species = c("kc","pm")),
    b = list(density = 0.01, species = c("kc","pm")),
    cnames = c("kc","pm"), tf = exp, fcov = vcovHC)
```

<sup>&</sup>lt;sup>1</sup>Consistency is a rather technical condition, but roughly speaking a *consistent estimator* is one such that its sampling distribution becomes increasingly concentrated around the value being estimated as n increases.

```
estimate lower upper
kc 2.967 2.105 4.181
pm 2.877 1.836 4.507
```

For comparison consider also the results when using quasi-likelihood.

```
m <- glm(cbind(y, total - y) ~ species + density + species:density,
    family = quasibinomial, data = trtools::rotifer)
cbind(summary(m)$coefficients, confint(m))</pre>
```

```
Estimate Std. Error t value Pr(>|t|)
                                                           2.5 % 97.5 %
(Intercept)
                  -114.352
                                14.95 -7.6472 4.736e-09 -146.02 -87.01
speciespm
                     4.629
                                24.46 0.1893 8.509e-01
                                                          -46.15 51.31
density
                   108.746
                                14.30 7.6056 5.358e-09
                                                           82.60 139.02
speciespm:density
                    -3.077
                                23.46 -0.1312 8.964e-01
                                                          -47.81
                                                                  45.70
contrast(m,
  a = list(density = 0.02, species = c("kc", "pm")),
  b = list(density = 0.01, species = c("kc", "pm")),
  cnames = c("kc","pm"), tf = exp)
```

```
estimate lower upper
kc 2.967 2.220 3.965
pm 2.877 1.973 4.195
```

Recall that heteroscedastic consistent standard errors are best used with generous sample sizes. For modest sample sizes (such as this experiment) quasi-likelihood is probably better.

### Generalized Linear Models Revisited

Recall that a generalized linear model (GLM) has the form

$$g[E(Y_i)] = \underbrace{\beta_0 + \beta_1 x_{i1} + \beta_1 x_{i2} + \dots + \beta_k x_{ik}}_{\eta_i},$$

where g is the link function and  $\eta_i$  is the linear predictor or systematic component. This is the mean structure of the model.

The variance structure of a GLM is

$$Var(Y_i) = \phi V[E(Y_i)],$$

where  $\phi$  is a dispersion parameter and V is the variance function.

If we define  $h = g^{-1}$  so that  $E(Y_i) = h(\eta_i)$  we can write a GLM concisely as

$$E(Y_i) = h(\eta_i) \tag{1}$$

$$Var(Y_i) = \phi V[h(\eta_i)] \tag{2}$$

to define the mean structure and a variance structure for  $Y_i$ , respectively, by specifying the mean and variance of  $Y_i$  to be functions of  $x_{i1}, x_{i2}, \ldots, x_{ik}$ .

The specification of a generalized linear model therefore requires three components.

- 1. The systematic component  $\eta_i = \beta_0 + \beta_1 x_{i1} + \beta_1 x_{i2} + \cdots + \beta_k x_{ik}$ .
- 2. The link function g for the mean structure  $g[E(Y_i)] = \eta_i$ .
- 3. The distribution of the response variable  $Y_i$ , which implies the variance structure  $Var(Y_i) = \phi V[E(Y_i)]$ , or we can specify the variance structure directly.

Four common distributions from the exponential family of distributions (normal/Gaussian, Poisson, gamma, and inverse-Gaussian) imply variance structures of the form

$$Var(Y_i) = \phi E(Y_i)^p$$

The values of p are p = 0 (normal/Gaussian), p = 1 (Poisson if  $\phi = 1$ ), p = 2 (gamma), and p = 3 (inverse-Gaussian). Also note that when using quasi-likelihood we can use other values of p via the tweedie function from the **statmod** package.

## GLMs for Gamma-Distributed Response Variables

If  $Y_i$  has a gamma distribution then  $Y_i$  is a positive and continuous random variable, and  $Var(Y_i) = \phi E(Y_i)^2$ . Such models are sometimes suitable for response variables that are bounded below by zero and right-skewed. Common link functions include the log and inverse functions. With a log link function we have a mean structure like that for Poisson regression where

$$\log E(Y_i) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \dots + \beta_k x_{ik},$$

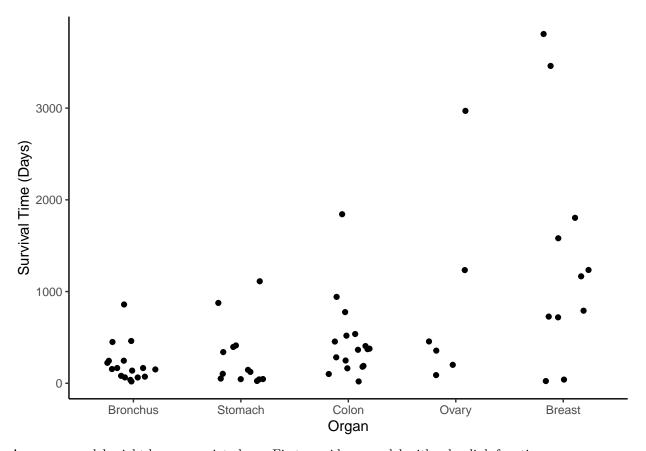
or

$$E(Y_i) = \exp(\beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \dots + \beta_k x_{ik}),$$

so the effects of explanatory variables and contrasts can be interpreted by applying the exponential function  $e^x$  and interpreting the effects as multiplicative factors or percent increase/decrease or percent larger/smaller.

**Example**: Consider again the cancer survival time data.

```
library(Stat2Data)
data(CancerSurvival)
CancerSurvival$0rgan <- with(CancerSurvival, reorder(Organ, Survival, mean))
p <- ggplot(CancerSurvival, aes(x = Organ, y = Survival)) +
   geom_jitter(height = 0, width = 0.25) +
   labs(y = "Survival Time (Days)") + theme_classic()
plot(p)</pre>
```



A gamma model might be appropriate here. First consider a model with a log link function.

```
m <- glm(Survival ~ Organ, family = Gamma(link = log), data = CancerSurvival)
cbind(summary(m)$coefficients, confint(m))</pre>
```

```
Estimate Std. Error t value Pr(>|t|)
                                                     2.5 % 97.5 %
                         0.2504 21.3854 1.773e-29
(Intercept)
              5.3546
                                                  4.90101
OrganStomach
              0.3013
                         0.3804 0.7923 4.314e-01 -0.44036
                                                            1.068
OrganColon
              0.7709
                         0.3541 2.1772 3.348e-02 0.06991
OrganOvary
              1.4302
                         0.4902 2.9174 4.987e-03 0.52462
                                                            2.486
              1.8867
                         0.3995 4.7228 1.479e-05 1.11572 2.702
OrganBreast
```

We might compare the survival times to the type of cancer with lowest expected survival time.

```
contrast(m, tf = exp,
    a = list(Organ = c("Stomach", "Colon", "Ovary", "Breast")),
    b = list(Organ = "Bronchus"),
    cnames = paste(c("Stomach", "Colon", "Ovary", "Breast"), "/", "Bronchus", sep = ""))
```

```
      estimate
      lower
      upper

      Stomach/Bronchus
      1.352
      0.6314
      2.893

      Colon/Bronchus
      2.162
      1.0644
      4.391

      Ovary/Bronchus
      4.180
      1.5671
      11.147

      Breast/Bronchus
      6.597
      2.9663
      14.673
```

Now suppose we specify the same variance structure directly. Note that the results are identical.

```
m <- glm(Survival ~ Organ, family = quasi(link = log, variance = "mu^2"), data = CancerSurvival)
cbind(summary(m)$coefficients, confint(m))</pre>
```

```
Estimate Std. Error t value Pr(>|t|)
                                                       2.5 % 97.5 %
(Intercept)
               5.3546
                          0.2504 21.3854 1.773e-29 4.90101 5.889
OrganStomach
               0.3013
                          0.3804 0.7923 4.314e-01 -0.44036
                                                              1.068
OrganColon
               0.7709
                          0.3541
                                  2.1772 3.348e-02
                                                    0.06991
                                                              1.472
OrganOvary
               1.4302
                          0.4902
                                  2.9174 4.987e-03
                                                     0.52462
                                                              2.486
OrganBreast
               1.8867
                          0.3995
                                 4.7228 1.479e-05
                                                     1.11572
                                                              2.702
contrast(m, tf = exp,
  a = list(Organ = c("Stomach", "Colon", "Ovary", "Breast")),
  b = list(Organ = "Bronchus"),
  cnames = paste(c("Stomach", "Colon", "Ovary", "Breast"), "/", "Bronchus", sep = ""))
```

 stimate
 lower
 upper

 Stomach/Bronchus
 1.352
 0.6314
 2.893

 Colon/Bronchus
 2.162
 1.0644
 4.391

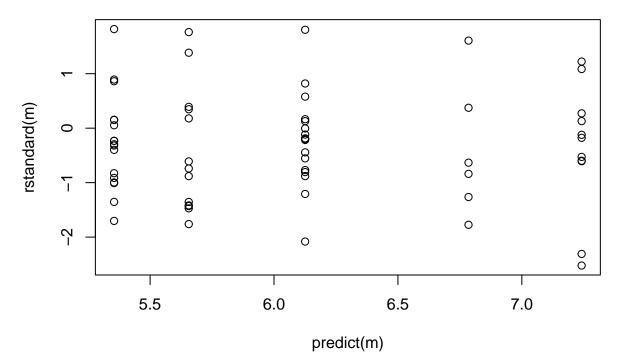
 Ovary/Bronchus
 4.180
 1.5671
 11.147

 Breast/Bronchus
 6.597
 2.9663
 14.673

Naturally we should check the residuals to see if the variance structure is reasonable.

```
plot(predict(m), rstandard(m), main = "Residual Plot")
```

## **Residual Plot**



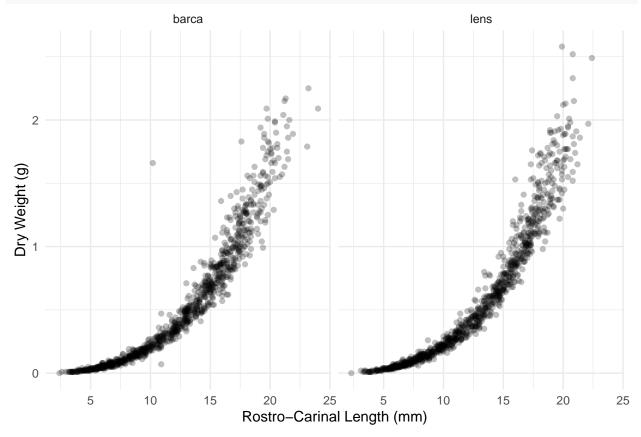
**Example:** Consider the following observations of dry weight (in grams) and rostro-carinal length (in mm) of a species of barnacles sampled from the inter-tidal zones near Punta Lens and Punta de la Barca along the Atlantic coast of Spain.

```
library(npregfast)
head(barnacle)
```

```
DW RC F
1 0.14 9.5 barca
2 0.00 2.4 barca
```

```
3 0.42 13.1 barca
4 0.01  3.7 barca
5 0.03  5.6 barca
6 1.56 18.6 barca

p <- ggplot(barnacle, aes(x = RC, y = DW))
p <- p + geom_point(alpha = 0.25) + facet_wrap(~ F) + theme_minimal()
p <- p + labs(x = "Rostro-Carinal Length (mm)", y = "Dry Weight (g)")
plot(p)</pre>
```



A common allometric regression model would have the form

$$E(Y_i) = ax_i^b$$

where  $Y_i$  is the dry weight for the *i*-th observation, and  $x_i$  is the rostro-carinal length for the *i*-th observation. We can also write this as

$$\log E(Y_i) = \log a + b \log x_i$$

or, equivalently,

$$E(Y_i) = \exp(\log a + b \log x_i)$$

or

$$E(Y_i) = \exp(\beta_0 + \beta_1 \log x_i)$$

where  $\beta_0 = \log a$  and  $\beta_1 = b$ . This is basically a log-linear model since we can write

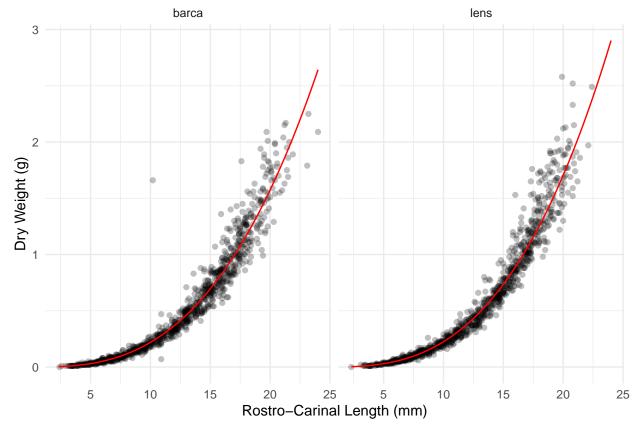
$$\log E(Y_i) = \beta_0 + \beta_1 \log x_i.$$

Because dry weight is continuous and positive, with the variability appearing to increase with the expected dry weight, we might specify a gamma distribution for dry weight.

```
barnacle <- subset(barnacle, DW > 0) # remove observations of zero weight to avoid errors
m <- glm(DW ~ F + log(RC) + F:log(RC), family = Gamma(link = log), data = barnacle)
summary(m)$coefficients

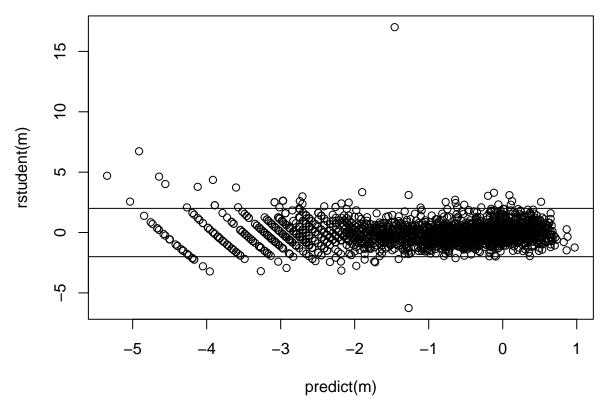
Estimate Std. Error t value Pr(>|t|)
```

```
(Intercept)
               -8.06129
                           0.03898 -206.786 0.0000000
Flens
               -0.15688
                           0.05724
                                      -2.741 0.0061837
log(RC)
                2.84234
                           0.01585 179.296 0.0000000
Flens:log(RC) 0.07884
                           0.02315
                                       3.405 0.0006744
d \leftarrow expand.grid(F = c("barca","lens"), RC = seq(2.3, 24, length = 100))
d$yhat <- predict(m, newdata = d, type = "response")</pre>
p <- p + geom_line(aes(y = yhat), color = "red", data = d)</pre>
plot(p)
```



```
# effect of a 20% increase in RC
contrast(m, tf = exp,
    a = list(F = c("barca","lens"), RC = 6),
    b = list(F = c("barca","lens"), RC = 5),
    cnames = c("barca","lens"))
```

# **Residual Plot**



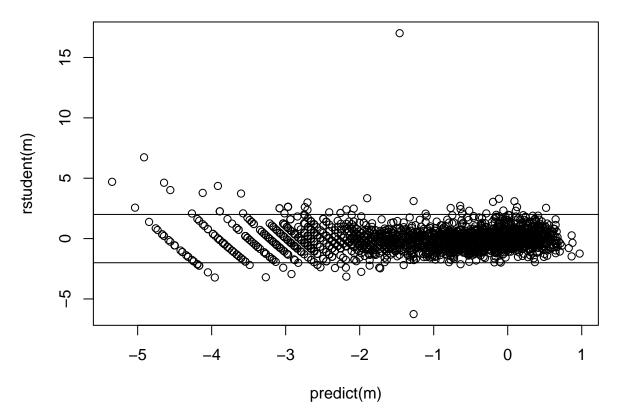
Note: Eliminating a couple of observations due to having a zero dry weight is not of much consequence here since there are so many observations. But if there were fewer observations this would not be a good idea. A better approach would be to just specify the same model using quasi. Note that using quasi with variance = "mu^2" is effectively equivalent to using family = gamma.

```
m <- glm(DW ~ F + log(RC) + F:log(RC), data = barnacle,
   family = quasi(link = "log", variance = "mu^2"))
summary(m)$coefficients</pre>
```

```
Estimate Std. Error
                                  t value Pr(>|t|)
(Intercept)
                          0.03898 -206.786 0.0000000
              -8.06129
Flens
              -0.15688
                          0.05724
                                     -2.741 0.0061837
log(RC)
                                    179.296 0.0000000
               2.84234
                          0.01585
Flens:log(RC)
               0.07884
                          0.02315
                                      3.405 0.0006744
```

```
# effect of a 20% increase in RC
contrast(m, tf = exp,
    a = list(F = c("barca", "lens"), RC = 6),
    b = list(F = c("barca", "lens"), RC = 5),
    cnames = c("barca","lens"))
      estimate lower upper
         1.679 1.670 1.689
barca
lens
         1.703 1.693 1.714
# comparing the two locations at different values of RC
contrast(m, tf = exp,
    a = list(F = "lens", RC = c(10,15,20)),
    b = list(F = "barca", RC = c(10,15,20)),
    cnames = c("10mm","15mm","20mm"))
     estimate lower upper
10mm
        1.025 1.004 1.046
        1.058 1.034 1.083
15mm
20mm
        1.083 1.048 1.118
Checking the residuals.
plot(predict(m), rstudent(m), main = "Residual Plot")
abline(-2,0)
abline(2,0)
```

# **Residual Plot**



Inverse-gaussian GLMs are similar. There the variance increases a bit faster with the expected response. To estimate such a model use family = inverse.gaussian. An equivalent model is to use quasi with

variance = mu^3.