

Friday, Mar 22

## Over-dispersion

Over-dispersion can occur for generalized linear models that assume a Poisson or binomial distribution for the response variable.

When we specify a distribution in a generalized linear model, what we are actually specifying is the *variance structure*

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

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

*Over-dispersion* is when

$$\text{Var}(Y_i) > \phi V[E(Y_i)],$$

and *underdispersion* is when

$$\text{Var}(Y_i) < \phi V[E(Y_i)].$$

Over-dispersion is fairly common in practice, but under-dispersion is relatively rare.

## Over-dispersion in Poisson Regression

If  $Y_i$  has a *Poisson* distribution, then

$$\text{Var}(Y_i) = E(Y_i),$$

so that it is implicitly assumed that  $\phi = 1$  and  $V(z) = z$ . Over-dispersion occurs if

$$\text{Var}(Y_i) > E(Y_i).$$

## Over-dispersion in Binomial Regression

If  $C_i$  has a *binomial* distribution, and  $Y_i = C_i/m_i$ , then

$$\text{Var}(Y_i) = E(Y_i)[1 - E(Y_i)]/m_i,$$

so that it is implicitly assumed that  $\phi = 1$  and  $V(z) = z(1 - z)/m_i$ . over-dispersion occurs if

$$\text{Var}(Y_i) > E(Y_i)[1 - E(Y_i)]/m_i.$$

In general, failing to account for over-dispersion (or a misspecification of the variance structure in general) may yield incorrect standard errors (usually too small in the case of over-dispersion), leading to incorrect test statistics and confidence intervals.

## Causes of Over-dispersion

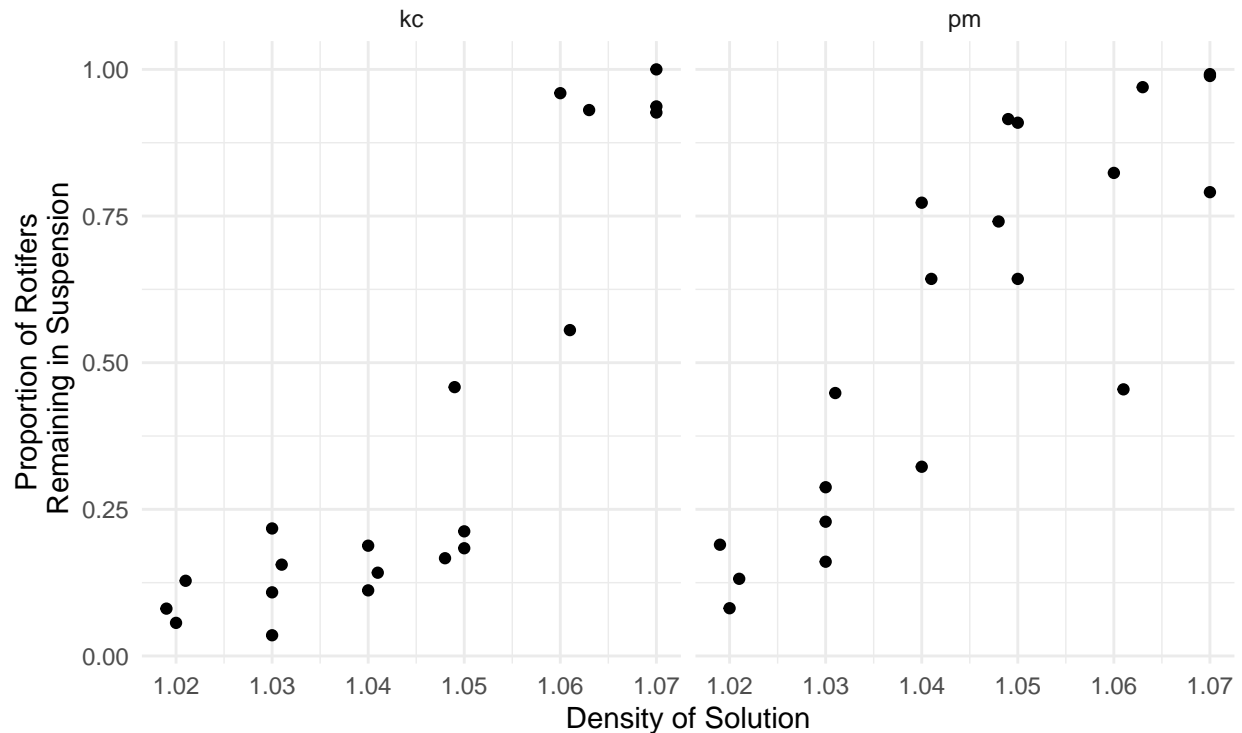
1. Wrong assumed distribution for the response variable.
2. Unobserved explanatory variables that vary over observations.

Note: A misspecified *mean structure* (e.g., failing to transform an explanatory variable or omitting a strong interaction) may appear as overdispersion.

**Example:** Consider the following data from an experiment that investigated the proportion of rotifers of two species remaining in suspension in different solution densities after being put into a centrifuge.

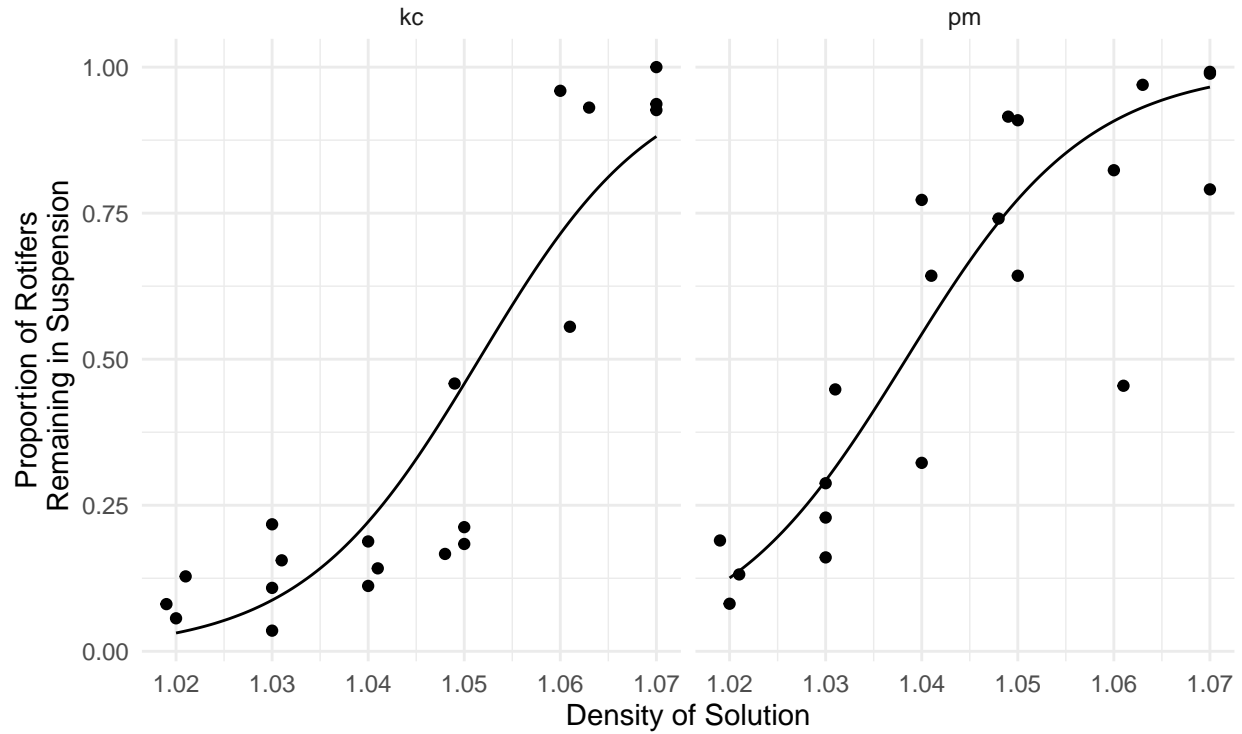
```
myrotifer <- trtools::rotifer
```

```
p <- ggplot(myrotifer, aes(x = density, y = y/total)) +  
  geom_point() + facet_wrap(~species) +  
  labs(y = "Proportion of Rotifers\nRemaining in Suspension",  
       x = "Density of Solution") + theme_minimal()  
plot(p)
```



Logistic regression might be a reasonable model here.

```
m <- glm(cbind(y, total - y) ~ species * density,  
        family = binomial, data = myrotifer)  
  
d <- expand.grid(species = c("kc", "pm"), density = seq(1.02, 1.07, length = 100))  
d$yhat <- predict(m, newdata = d, type = "response")  
  
p <- p + geom_line(aes(y = yhat), data = d)  
plot(p)
```



Do these data exhibit over-dispersion for this model?

## Detection of Over-dispersion

Standardized residuals can be used to detect over-dispersion. There are several types for GLMs.

1. Pearson residuals. Pearson residuals are defined as

$$\frac{y_i - \hat{y}_i}{\sqrt{\widehat{\text{Var}}(Y_i)}}.$$

Dividing a Pearson residual by another term to account for the variance  $\hat{y}_i$  creates a *standardized* Pearson residual. These are obtained using `rstandard(m, type = "pearson")`.

2. Deviance residuals. The residual deviance can be decomposed into a per-observation contribution so that  $D = \sum_{i=1}^n d_i$ . Then the residual deviance is defined as

$$\text{sign}(y_i - \hat{y}_i) \sqrt{d_i},$$

where

$$\text{sign}(z) = \begin{cases} 1, & \text{if } z > 0, \\ 0, & \text{if } z = 0, \\ -1, & \text{if } z < 0. \end{cases}$$

Dividing a deviance residual by another term to account for the variance  $\hat{y}_i$  creates a *standardized* deviance residual. These are obtained using `rstandard(m, type = "deviance")`. A numerical approximation to these residuals obtained when omitting the observation can be obtained using `rstudent(m)`.

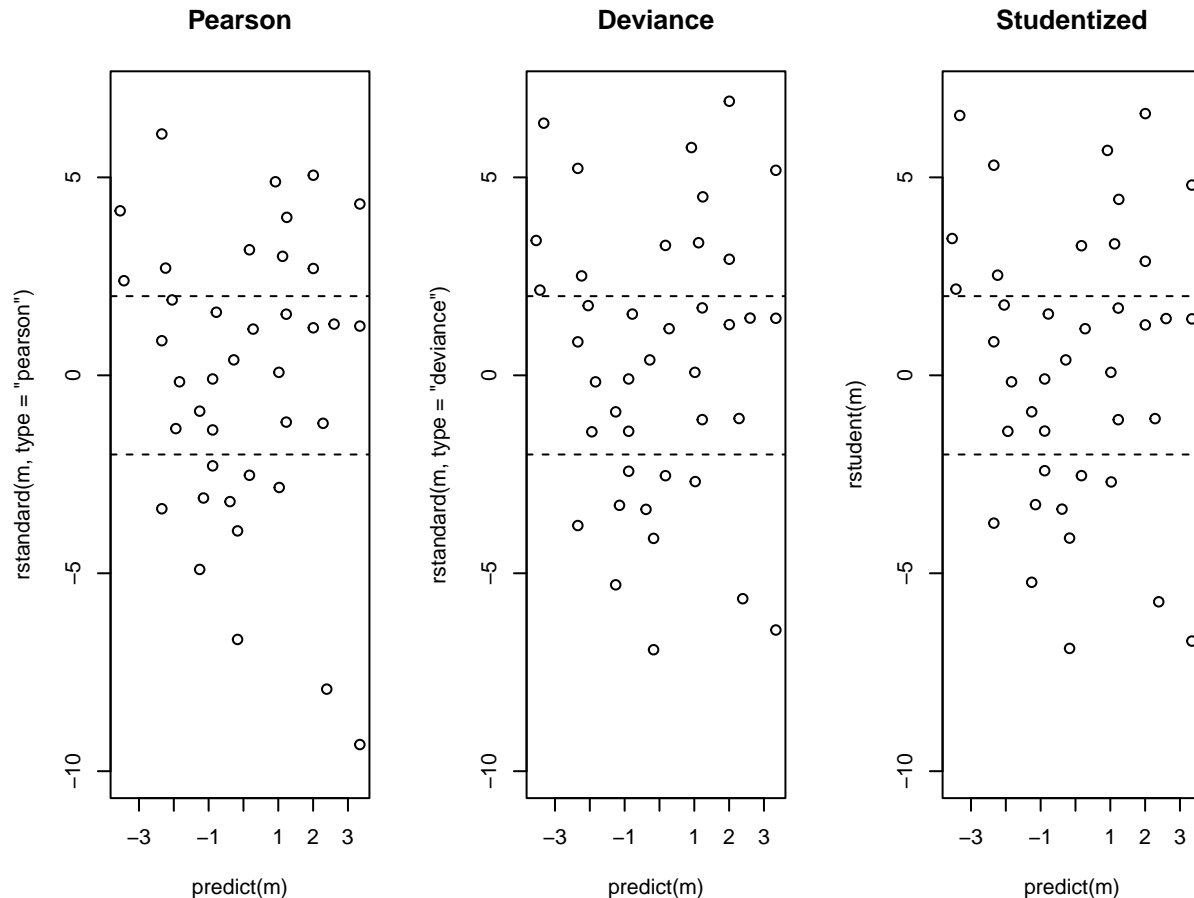
3. Studentized residuals. The function `rstudent` will produce *approximate* studentized residuals for GLMs.

Comment: If the model is correct the residuals *might* be approximately normally distributed with a mean of zero and standard deviation of one (i.e., “standard normal”), so an excess of values greater than two (in absolute value) may indicate over-dispersion or some other problem with the model. But with very *coarse*

data (e.g., very small counts in a Poisson regression model or proportions with small  $m_i$  in a logistic regression model), the distribution of these residuals is not approximately normal.

**Example:** Let's look at the residuals for the `rotifer` model.

```
par(mfcol = c(1,3))
plot(predict(m), rstandard(m, type = "pearson"), ylim = c(-10, 7), main = "Pearson")
abline(h = c(-2,2), lty = 2)
plot(predict(m), rstandard(m, type = "deviance"), ylim = c(-10, 7), main = "Deviance")
abline(h = c(-2,2), lty = 2)
plot(predict(m), rstudent(m), ylim = c(-10, 7), main = "Studentized")
abline(h = c(-2,2), lty = 2)
```



Is there an explanation of the over-dispersion?

Another metric is to compare the residual deviance to the residual degrees of freedom in a GLM with a response variable with either a Poisson or binomial distribution. If the model is (approximately) correct then the ratio of the residual deviance to the residual degrees of freedom is approximately one.

**Example:** Consider the residual deviance and residual degrees of freedom for the `rotifer` model.

```
summary(m)
```

Call:

```
glm(formula = cbind(y, total - y) ~ species * density, family = binomial,
     data = myrotifer)
```

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	-114.35	4.03	-28.35	<2e-16 ***
speciespm	4.63	6.60	0.70	0.48
density	108.75	3.86	28.19	<2e-16 ***
speciespm:density	-3.08	6.33	-0.49	0.63

---

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

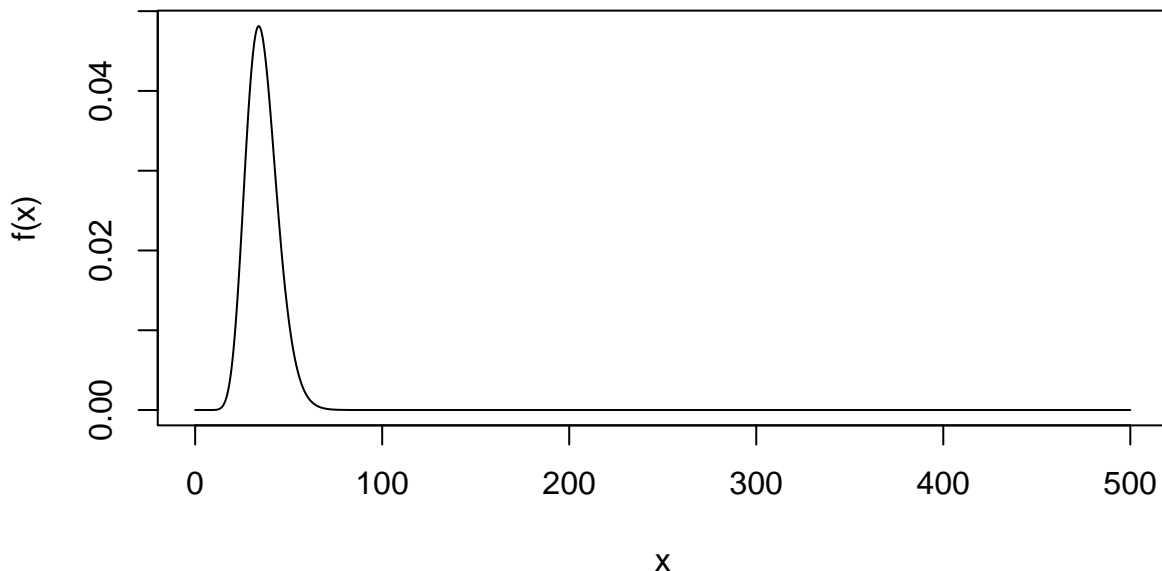
(Dispersion parameter for binomial family taken to be 1)

Null deviance: 3180.99 on 39 degrees of freedom  
Residual deviance: 434.02 on 36 degrees of freedom  
AIC: 596.6

Number of Fisher Scoring iterations: 5

If the model is correct and there is no over-dispersion, the residual deviance has approximate a  $\chi^2$  distribution with degrees of freedom equal to the residual degrees of freedom. We can use this as an informal test for over-dispersion.

```
f <- function(x) dchisq(x, 36)
curve(f, from = 0, to = 500, n = 1000)
```



```
1 - pchisq(434.02, df = 36)
```

```
[1] 0
```

Residuals are more informative, but the residual deviance is a quick way to check to see if over-dispersion may be an issue.

Note: For logistic regression, over-dispersion *cannot* be diagnosed in this way for *binary* data (and the residual deviance may not be reliable if the  $m_i$  are very small).

**Example:** Let's look again at the Poisson regression model for the trawling data.

```
library(COUNT)
data(fishing)
```

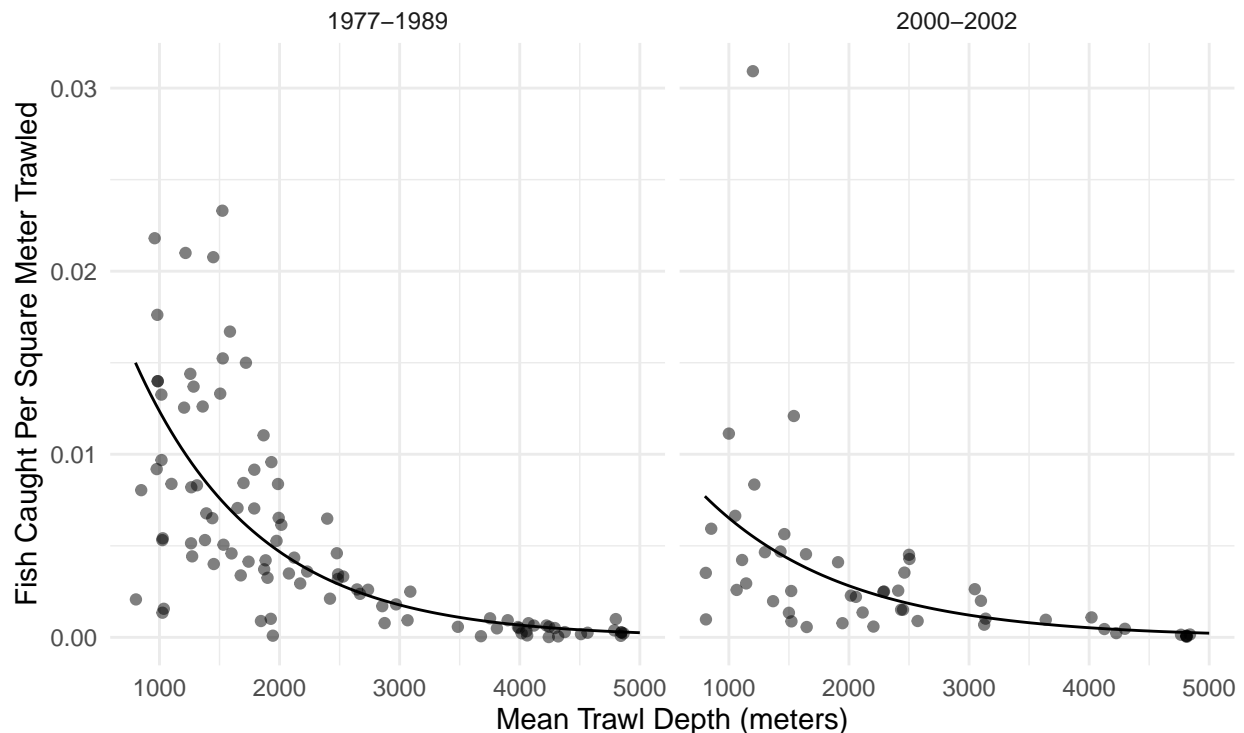
```

m <- glm(totabund ~ period * meandepth + offset(log(sweptarea)),
  family = poisson, 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)

```



Might there be over-dispersion here?

```
summary(m)
```

Call:

```
glm(formula = totabund ~ period * meandepth + offset(log(sweptarea)),
    family = poisson, data = fishing)
```

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	-3.42e+00	1.49e-02	-229.67	<2e-16 ***
period2000-2002	-7.71e-01	2.97e-02	-25.94	<2e-16 ***
meandepth	-9.71e-04	7.96e-06	-121.94	<2e-16 ***
period2000-2002:meandepth	1.32e-04	1.52e-05	8.65	<2e-16 ***

---

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

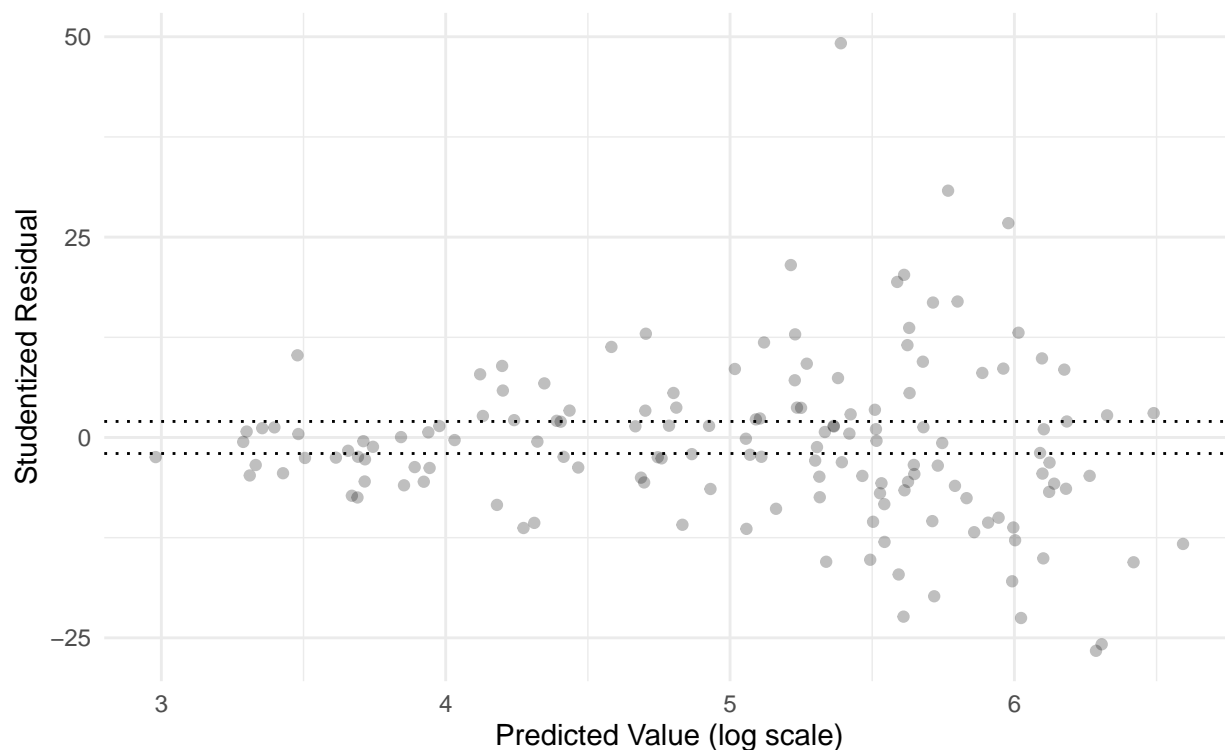
(Dispersion parameter for poisson family taken to be 1)

Null deviance: 46176 on 146 degrees of freedom  
Residual deviance: 14982 on 143 degrees of freedom  
AIC: 15962

Number of Fisher Scoring iterations: 5

```
fishing$eta <- predict(m)
fishing$res <- rstudent(m)

p <- ggplot(fishing, aes(x = eta, y = res)) + theme_minimal() +
  geom_point(alpha = 0.25) +
  labs(x = "Predicted Value (log scale)",
       y = "Studentized Residual") +
  geom_hline(yintercept = c(-2, 2), linetype = 3)
plot(p)
```



Over-dispersion is not the only issue here. The variance of the residuals is not constant.

## Solutions to Over-dispersion

There are several potential solutions to over-dispersion.

1. Quasi-likelihood. Specify a variance structure other than the one implied by a specified distribution.
2. Specify a different distribution (possibly outside the exponential family).
3. Use a robust estimator of the standard errors (i.e., heteroscedastic consistent standard errors).

## Quasi-Likelihood Solutions to Over-dispersion

The Poisson and binomial distributions assume the variance structures

$$\text{Var}(Y_i) = \phi E(Y_i) \quad \text{and} \quad \text{Var}(Y_i) = \phi E(Y_i)[1 - E(Y_i)]/m_i,$$

respectively, where the *dispersion parameter* is *fixed* at  $\phi = 1$ . One solution is to allow  $\phi$  to be an unknown parameter to “relax” the variance structure and allow the variance to be larger than it would be for a Poisson or binomial distribution. The dispersion parameter can be estimated. R uses

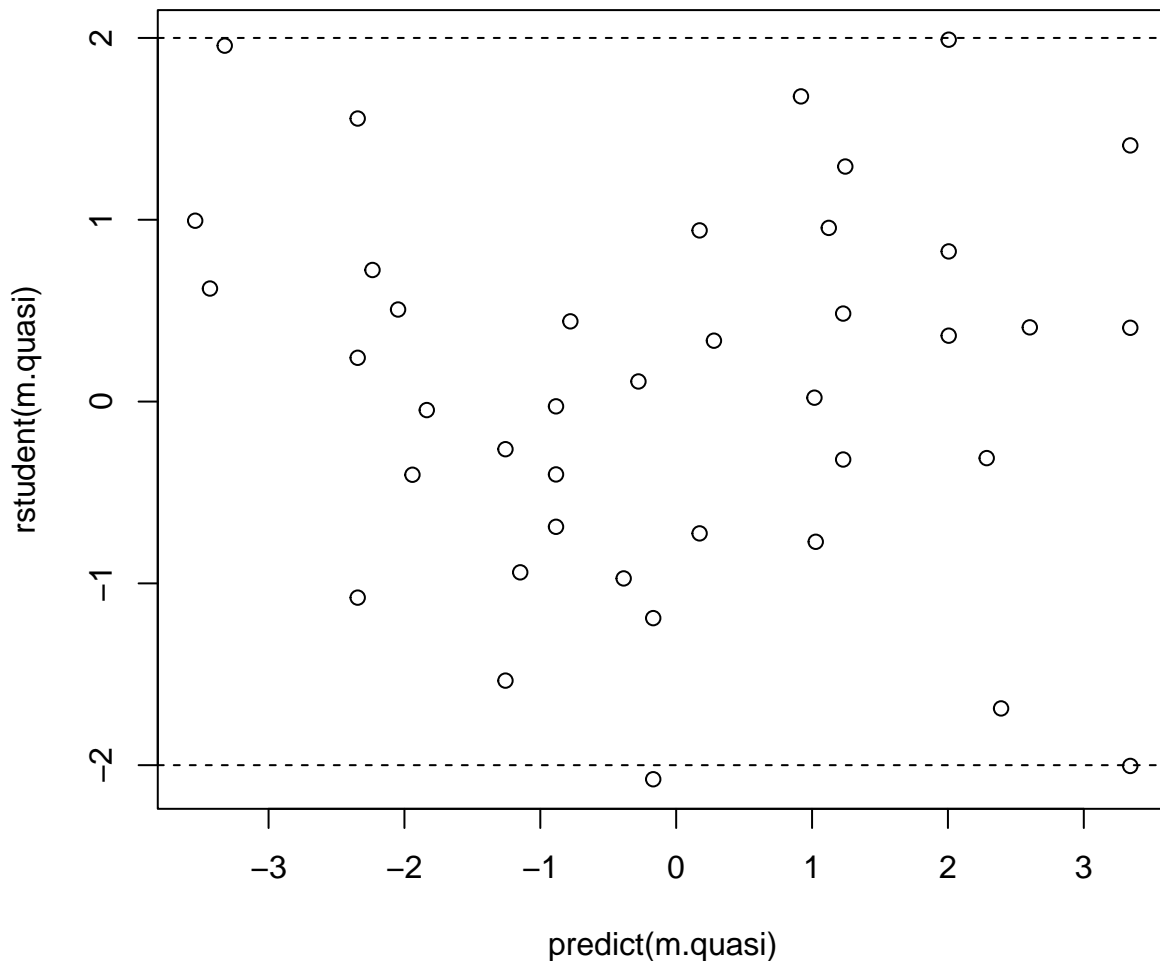
$$\hat{\phi} = \frac{1}{n-p} \sum_{i=1}^n \frac{(y_i - \hat{y}_i)^2}{V(\hat{y}_i)},$$

which is analogous to the estimate of  $\sigma^2$  in a normal linear model. This is a quasi-likelihood approach because the variance structures with  $\phi \neq 1$  do not correspond to a binomial or Poisson distribution. This kind of quasi-likelihood can be done with `glm` by using `quasipoisson` or `quasibinomial` instead of `poisson` or `binomial`, respectively, when specifying the `family` argument.

**Example:** Consider again the rotifer model.

```
m.quasi <- glm(cbind(y, total - y) ~ species + density + species:density,
  family = quasibinomial, data = myrotifer)
plot(predict(m.quasi), rstudent(m.quasi), main = "Residual Plot")
abline(h = c(-2,2), lty = 2)
```

### Residual Plot





Note: You **cannot** compare the residual deviance to the residual degrees of freedom as a diagnostic to determine if using quasi-likelihood was successful, but standardized residuals are still appropriate.

How does this impact our inferences?

```
m.binom <- glm(cbind(y, total - y) ~ species + density + species:density,
  family = binomial, data = myrotifer)
cbind(summary(m.binom)$coefficients, confint(m.binom))
```

	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(summary(m.quasi)$coefficients, confint(m.quasi))
```

	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

*# odds ratios for effect of a 0.01 unit increase in density*

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

	estimate	lower	upper
kc	2.967	2.751	3.200
pm	2.877	2.607	3.174

*# odds ratios for effect of a 0.01 unit increase in density*

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

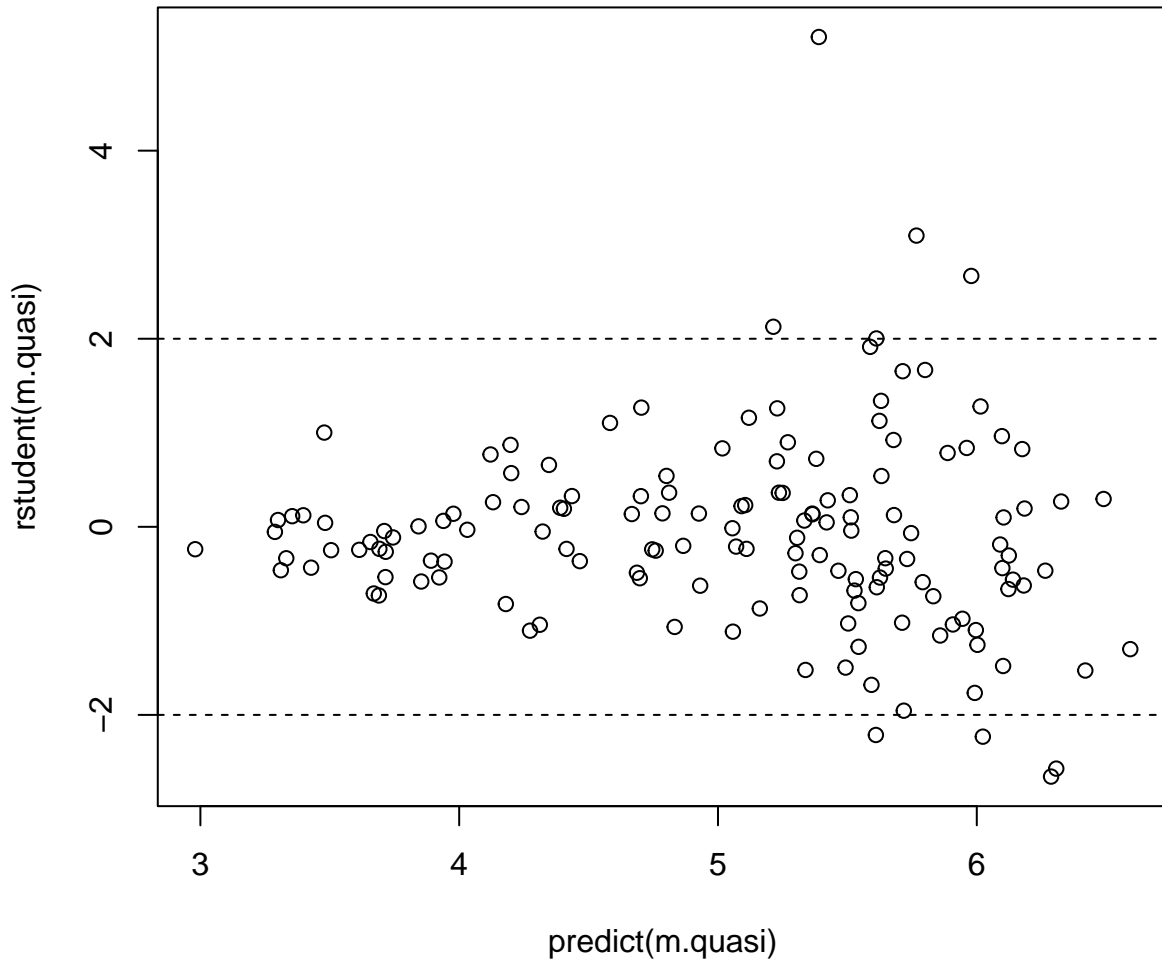
	estimate	lower	upper
kc	2.967	2.220	3.965
pm	2.877	1.973	4.195

Note that point estimates are unchanged, but standard errors, tests, and confidence intervals are affected.

**Example:** Now let's try the same approach with trawling data.

```
m.quasi <- glm(totabund ~ period * meandepth + offset(log(sweptarea)),
  family = quasipoisson, data = fishing)
plot(predict(m.quasi), rstudent(m.quasi), main = "Residual Plot")
abline(h = c(-2, 2), lty = 2)
```

## Residual Plot



That was maybe somewhat less successful. Note the “megaphone” pattern. The assumed variance structure is

$$\text{Var}(Y_i) = \phi E(Y_i).$$

We could relax this by assuming instead

$$\text{Var}(Y_i) = \phi E(Y_i)^p.$$

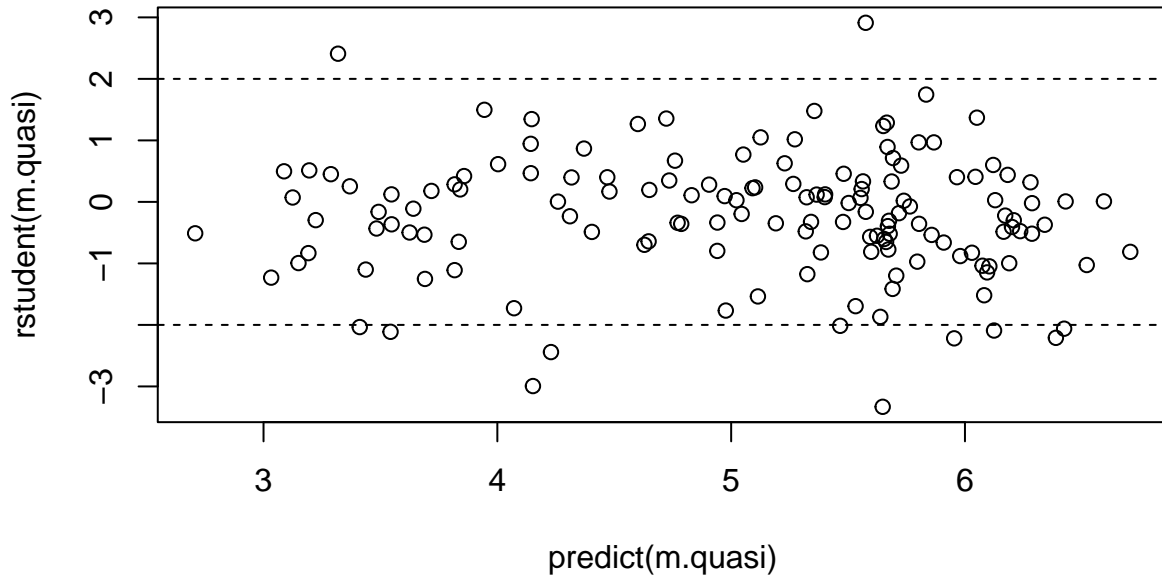
for some  $p > 1$ . If  $p = 1, 2$ , or  $3$  then we can use `quasi`. Here we are using it for  $p = 2$ .

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

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	-3.250e+00	1.592e-01	-20.4180	3.187e-44
period2000-2002	-6.041e-01	2.720e-01	-2.2212	2.791e-02
meandepth	-1.041e-03	5.866e-05	-17.7403	5.988e-38
period2000-2002:meandepth	7.272e-05	9.992e-05	0.7278	4.679e-01

```
plot(predict(m.quasi), rstudent(m.quasi), main = "Residual Plot")
abline(h = c(-2,2), lty = 2)
```

## Residual Plot



Note that `quasi(link = "log", variance = "mu")` is the same as `quasipoisson`. For more options consider `family = tweedie`. The `tweedie` family defines power functions for link and variance functions of the form

$$E(Y_i)^q = \eta_i \quad \text{and} \quad \text{Var}(Y_i) = \phi E(Y_i)^p,$$

where  $E(Y_i)^0 \equiv \log E(Y_i)$  when using `tweedie` (not mathematically of course — this is just for interface purposes). For example, to replicate the quasi-likelihood model above we can use the following.

```
library(statmod) # for tweedie "family"
m.tweedie <- glm(totabund ~ period * meandepth + offset(log(sweptarea)),
  family = tweedie(link.power = 0, var.power = 2), data = fishing)
summary(m.tweedie)$coefficients
```

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	-3.250e+00	1.592e-01	-20.4180	3.187e-44
period2000-2002	-6.041e-01	2.720e-01	-2.2212	2.791e-02
meandepth	-1.041e-03	5.866e-05	-17.7403	5.988e-38
period2000-2002:meandepth	7.272e-05	9.992e-05	0.7278	4.679e-01

The powers  $p$  and  $q$  are not required to be integers when using `tweedie`.

Whether or not we use quasi-likelihood will affect the standard errors, as well as tests and confidence intervals. Failing to account for substantial over-dispersion can result in biased standard errors, and thus incorrect tests and confidence intervals. Estimates of parameters (of functions thereof such as what we get from `contrast`) may or may not change, depending on the variance structure.

```
m.poisson <- glm(totabund ~ period * meandepth + offset(log(sweptarea)),
  family = poisson, data = fishing)
# rate ratios for year
trtools::contrast(m.poisson,
  a = list(sweptarea = 1, meandepth = c(1000,2000,3000,4000,5000), period = "2000-2002"),
  b = list(sweptarea = 1, meandepth = c(1000,2000,3000,4000,5000), period = "1977-1989"),
  cnames = c("1000m", "2000m", "3000m", "4000m", "5000m"), tf = exp)
```

	estimate	lower	upper
1000m	0.5277	0.5100	0.5460

2000m	0.6020	0.5861	0.6183
3000m	0.6869	0.6565	0.7187
4000m	0.7837	0.7293	0.8421
5000m	0.8941	0.8087	0.9885

```
trtools::contrast(m.tweedie,
  a = list(sweptarea = 1, meandepth = c(1000,2000,3000,4000,5000), period = "2000-2002"),
  b = list(sweptarea = 1, meandepth = c(1000,2000,3000,4000,5000), period = "1977-1989"),
  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
4000m	0.7311	0.4905	1.0897
5000m	0.7863	0.4458	1.3867

## Inferences With Quasi-Likelihood

Using quasi-likelihood instead of maximum likelihood changes how inferences are made in several ways.

1. The standard errors are multiplied by  $\sqrt{\hat{\phi}}$ . If  $\hat{\phi} > 1$  (which it probably is if over-dispersion is present) then the standard errors will be larger (and thus failing to account for over-dispersion leads us to usually underestimate them). Note that this adjustment is made automatically when using quasi-likelihood.
2. Wald confidence intervals and tests for a single parameter or function of parameters are based on the  $t$  distribution rather than the standard normal distribution. The  $t$  distribution is believed to provide more accurate results, although it is still an approximation.
3. Using `confint` or `anova` use the  $F$  distribution rather than the  $\chi^2$  distribution. The underlying test statistic is similar to the  $F$  test statistic used in normal linear models. When using `anova` you should use `test = "F"` rather than `test = "LRT"` if you are using quasi-likelihood.
4. Function in `emmeans` do not adjust the degrees of freedom for estimating the dispersion parameter when using quasi-likelihood. This does not make much difference unless  $n$  is small. But you can specify it manually via the `df` argument (use the degrees of freedom for the residual deviance from `summary` or extract it with `modelname$df.residual`). But `contrast` and `lincon` do not require manual specification, although you can via the `df` argument for those functions.

```
library(emmeans)
m.quasi <- glm(cbind(y, total - y) ~ species + density + species:density,
  family = quasibinomial, data = myrotifer)
trtools::contrast(m.quasi,
  a = list(species = c("kc", "pm"), density = 0.02),
  b = list(species = c("kc", "pm"), density = 0.01),
  cnames = c("kc", "pm"), tf = exp)
```

	estimate	lower	upper
kc	2.967	2.220	3.965
pm	2.877	1.973	4.195

```
pairs(emmeans(m.quasi, ~density|species, at = list(density = c(0.02, 0.01))),
  type = "response"), infer = TRUE) # wrong df
```

species = kc:									
contrast		odds.ratio	SE	df	asympt.LCL	asympt.UCL	null	z.ratio	p.value
density0.02 / density0.01		2.97	0.424	Inf	2.24	3.93	1	7.606	<.0001

```
species = pm:
  contrast          odds.ratio    SE  df asymp.LCL asymp.UCL null z.ratio p.value
density0.02 / density0.01      2.88 0.535 Inf      2.00      4.14    1   5.681 <.0001
```

Confidence level used: 0.95

Intervals are back-transformed from the log odds ratio scale

Tests are performed on the log odds ratio scale

```
pairs(emmeans(m.quasi, ~density|species, at = list(density = c(0.02, 0.01)),
  type = "response"), infer = TRUE, df = m.quasi$df.residual) # correct df
```

```
species = kc:
  contrast          odds.ratio    SE df lower.CL upper.CL null t.ratio p.value
density0.02 / density0.01      2.97 0.424 36      2.22      3.96    1   7.606 <.0001
```

```
species = pm:
  contrast          odds.ratio    SE df lower.CL upper.CL null t.ratio p.value
density0.02 / density0.01      2.88 0.535 36      1.97      4.20    1   5.681 <.0001
```

Degrees-of-freedom method: user-specified

Confidence level used: 0.95

Intervals are back-transformed from the log odds ratio scale

Tests are performed on the log odds ratio scale

Admittedly it does not make much difference here.

## Misspecified Mean Structures and over-dispersion

A poorly specified *mean structure* may be mistaken for over-dispersion.

```
library(trtools)
ceriodaphniastrain$strain <- factor(ceriodaphniastrain$strain, labels = c("a","b"))
m <- glm(count ~ strain + sqrt(concentration), family = poisson, data = ceriodaphniastrain)
summary(m)
```

Call:

```
glm(formula = count ~ strain + sqrt(concentration), family = poisson,
    data = ceriodaphniastrain)
```

Coefficients:

```
              Estimate Std. Error z value Pr(>|z|)
(Intercept)    4.5284     0.0400  113.08 < 2e-16 ***
strainb        -0.2750     0.0484   -5.68 1.3e-08 ***
sqrt(concentration) -1.6576     0.0474 -34.99 < 2e-16 ***
---
```

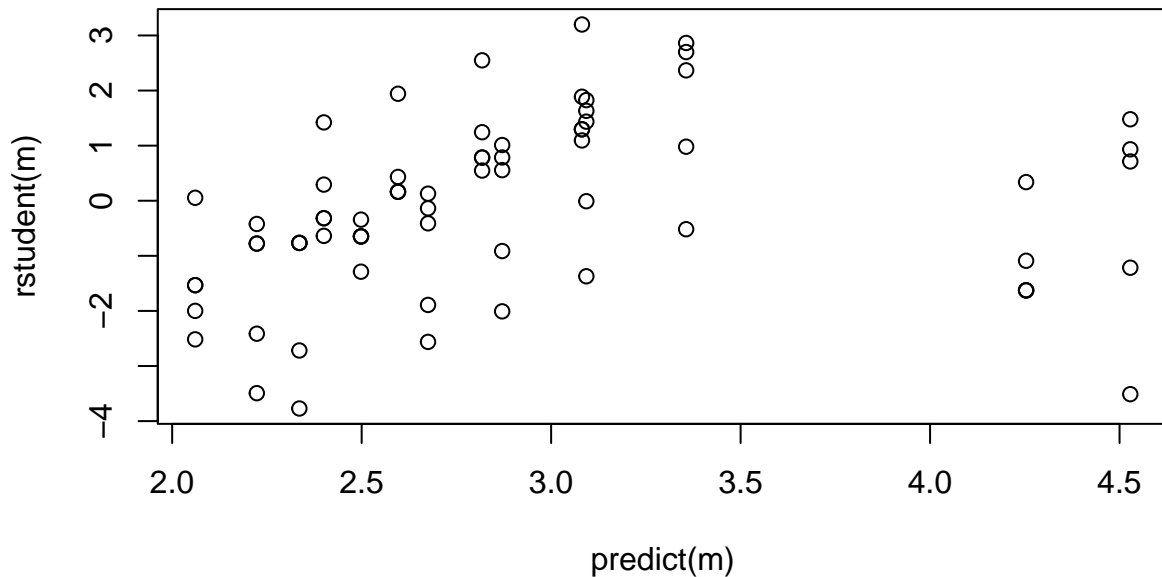
```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

(Dispersion parameter for poisson family taken to be 1)

```
Null deviance: 1359.38  on 69  degrees of freedom
Residual deviance: 164.28  on 67  degrees of freedom
AIC: 493.9
```

Number of Fisher Scoring iterations: 4

```
plot(predict(m), rstudent(m))
```



```
m <- glm(count ~ strain + concentration, family = poisson, data = ceriodaphniastrain)
summary(m)
```

Call:

```
glm(formula = count ~ strain + concentration, family = poisson,
     data = ceriodaphniastrain)
```

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	4.4546	0.0391	113.82	< 2e-16 ***
strainb	-0.2750	0.0484	-5.68	1.3e-08 ***
concentration	-1.5431	0.0466	-33.11	< 2e-16 ***

---

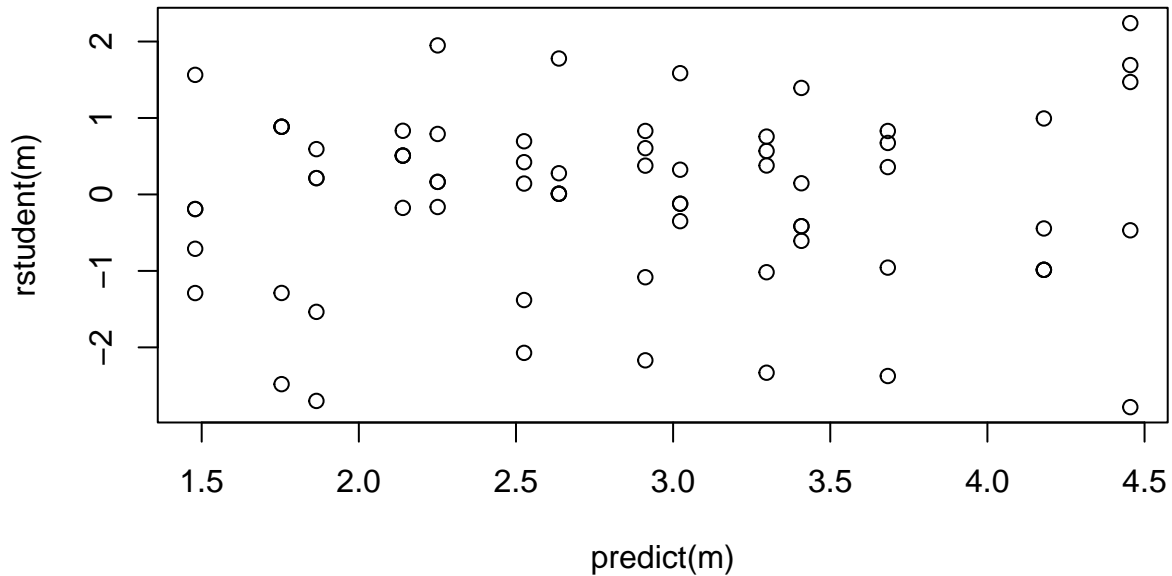
Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)

Null deviance: 1359.381 on 69 degrees of freedom  
Residual deviance: 86.376 on 67 degrees of freedom  
AIC: 416

Number of Fisher Scoring iterations: 4

```
plot(predict(m), rstudent(m))
```



## Quasi-Likelihood and Nonlinear Regression

Quasi-likelihood for a GLM is essentially the same as using (nonlinear) regression with iteratively weighted least squares to account for heteroscedasticity. The weights are

$$w_i = \frac{1}{V(\hat{y}_i)},$$

where  $V$  is the variance function.

**Example:** Consider the model for the trawling data where the variance is proportional to  $E(Y_i)^2$ . To estimate this model using iteratively weighted least squares we use weights of  $w_i = 1/E(Y_i)^2$ .

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

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	-3.250e+00	1.592e-01	-20.4180	3.187e-44
period2000-2002	-6.041e-01	2.720e-01	-2.2212	2.791e-02
meandepth	-1.041e-03	5.866e-05	-17.7403	5.988e-38
period2000-2002:meandepth	7.272e-05	9.992e-05	0.7278	4.679e-01

```
fishing$w <- 1
for (i in 1:10) {
  m.iwls <- nls(totabund ~ exp(b0 + b1*(period == "2000-2002") + b2*meandepth +
    b3*(period == "2000-2002")*meandepth + log(sweptarea)), data = fishing,
    start = list(b0 = -3, b1 = -0.6, b2 = 0, b3 = 0), weights = w)
  fishing$w <- 1 / predict(m.iwls)^2
}
summary(m.iwls)$coefficients
```

	Estimate	Std. Error	t value	Pr(> t )
b0	-3.250e+00	1.592e-01	-20.4179	3.189e-44
b1	-6.041e-01	2.720e-01	-2.2213	2.790e-02
b2	-1.041e-03	5.866e-05	-17.7402	5.991e-38
b3	7.273e-05	9.992e-05	0.7279	4.679e-01

**Example:** Consider the model for the rotifer data. Here the variance is proportional to  $E(Y_i)[1 - E(Y_i)]/m_i$  (recall that  $m_i$  is the “total possible” for the counts). To estimate this model using iteratively weighted least squares we use weights of

$$w_i = \frac{m_i}{E(Y_i)[1 - E(Y_i)]}.$$

```
m.binomial <- glm(cbind(y, total - y) ~ species * density,
  family = quasibinomial, data = myrotifer)
summary(m.binomial)$coefficients
```

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	-114.352	14.95	-7.6472	4.736e-09
speciespm	4.629	24.46	0.1893	8.509e-01
density	108.746	14.30	7.6056	5.358e-09
speciespm:density	-3.077	23.46	-0.1312	8.964e-01

```
myrotifer$w <- 1
for (i in 1:20) {
  m <- nls(y/total ~ plogis(b0 + b1*(species == "pm") + b2*density +
    b3*(species == "pm")*density), data = myrotifer, weights = w,
    start = list(b0 = -114, b1 = 4.6, b2 = 109, b3 = -3))
  myrotifer$yhat <- predict(m)
  myrotifer$w <- myrotifer$total / (myrotifer$yhat * (1 - myrotifer$yhat))
}
summary(m)$coefficients
```

	Estimate	Std. Error	t value	Pr(> t )
b0	-114.338	14.95	-7.6485	4.718e-09
b1	4.614	24.46	0.1887	8.514e-01
b2	108.732	14.29	7.6069	5.338e-09
b3	-3.063	23.46	-0.1306	8.968e-01

Note that `plogis` is the function  $e^x/(1 + e^x)$ . The model can be written as

$$E(Y_i) = \frac{e^{\eta_i}}{1 + e^{\eta_i}}$$

where  $Y_i$  is the observed *proportion*, and

$$\eta_i = \beta_0 + \beta_1 s_i + \beta_2 d_i + \beta_3 s_i d_i,$$

where  $s_i$  is an indicator variable for the `pm` species, and  $d_i$  is the density.

Using iteratively weighted least squares is not necessary if we can use `quasi` or `tweedie`, but it is a useful option for cases where the variance structure is outside what can be done with `quasi` or `tweedie` (although one can *program* new variance structures).