20.9. Donner Party.

For females: $\log (\text{odds}) = 3.2 - (0.078 \times age)$

For males: $\log (odds) = 1.6 - (0.078 \times age)$

- (a) The estimated probabilities of survival are:
 - For 25 year-old men:

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 1.6 - (0.078)(25) = -0.35$$
$$\hat{\pi} = \frac{\exp(-0.35)}{1 + \exp(-0.35)} = 0.4134$$

For 50 year-old men:

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 1.6 - (0.078)(50) = -2.3$$
$$\hat{\pi} = \frac{\exp(-2.3)}{1 + \exp(-2.3)} = 0.0911$$

• For 25 year-old women:

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 3.2 - (0.078)(25) = 1.25$$
$$\hat{\pi} = \frac{\exp(1.25)}{1 + \exp(1.25)} = 0.7773$$

• For 50 year-old women:

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 3.2 - (0.078)(50) = -0.7$$
$$\hat{\pi} = \frac{\exp(-0.7)}{1 + \exp(-0.7)} = \mathbf{0.3318}$$

(b) When the estimated probability of survival is $\hat{\pi} = 0.5$, then

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = \log\left(\frac{0.5}{0.5}\right) = \log(1) = 0$$

(i) For men, the age corresponding to $\hat{\pi} = 0.5$ is

$$0=1.6-(0.078\times age)\Rightarrow age=\mathbf{20.51}$$

(ii) For women, the age corresponding to $\hat{\pi} = 0.5$ is

$$0 = 3.2 - (0.078 \times age) \Rightarrow age = 41.03$$

20.10. Odds Ratio.

Let ω_A = the odds at A, and ω_B = the odds at B. Then

$$\log(\omega_A) - \log(\omega_B) = (\beta_0 + \beta_1 A + \beta_2 x_2 + \dots) - (\beta_0 + \beta_1 B + \beta_2 x_2 + \dots)$$
$$= \beta_1 A - \beta_1 B = \beta_1 (A - B)$$

Note, however, that the left-hand side can be written

$$\log(\omega_A) - \log(\omega_B) = \log\left(\frac{\omega_A}{\omega_B}\right)$$

and so taking the anti-logarithm of both sides yields

$$\frac{\omega_A}{\omega_B} = \exp\left(\beta_1(A - B)\right)$$

20.11

> shuttle=data.frame(TEMP=ex2011\$Temp,FAILURE=ex2011\$Failure)

(a) R output given below:

- > mshut <- glm(FAILURE~TEMP,family=binomial,data=shuttle)</pre>
- > summary(mshut)

Call:

glm(formula = FAILURE ~ TEMP, family = binomial, data = shuttle)

Deviance Residuals:

Coefficients:

Estimate Std. Error z value Pr(>|z|)
(Intercept) 10.87535 5.70291 1.907 0.0565 .
TEMP -0.17132 0.08344 -2.053 0.0400 *

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

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 28.975 on 23 degrees of freedom Residual deviance: 23.030 on 22 degrees of freedom

AIC: 27.030

Number of Fisher Scoring iterations: 4

The fitted logistic regression model is

$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 10.875 - 0.171(temp)$$

(b) Wish to test H_0 : $\beta_1 = 0$ vs. H_a : $\beta_1 < 0$. The test statistic is

$$z = \frac{\hat{\beta}_1}{\text{SE}(\hat{\beta}_1)} = \frac{-0.171}{0.083} = -2.053$$

The one-sided p-value is

$$P(Z < -2.053) = 0.020$$

We reject H_0 and conclude that the odds of O-ring failure decrease with temperature.

> pnorm(-2.053)
[1] 0.02003629

(c) Wish to test H_0 : $\beta_1 = 0$ vs. H_a : $\beta_1 \neq 0$, using the drop-in-deviance test. The test statistic is the drop in deviance:

$$\chi^2 = D_{reduced} - D_{full} = 28.975 - 23.030 = 5.945$$

Under H_0 , χ^2 follows a chi-squared distribution with 23 - 22 = 1 degree of freedom. The p-value for this test is

$$P(\chi_1^2 > 5.945) = 0.015$$

Hence we reject H_0 and conclude the full model (where $\beta_1 \neq 0$) is a better fit.

> 1-pchisq(5.945,df=1) [1] 0.01475909

(d) A 95% confidence interval for β_1 is given by

$$\hat{\beta}_1 \pm z(.975) \text{SE}(\hat{\beta}_1) = -0.1713 \pm (1.96)(0.0834)$$

= -0.335 to -0.0078.

(e) The estimated logit of failure probability at 31° F is

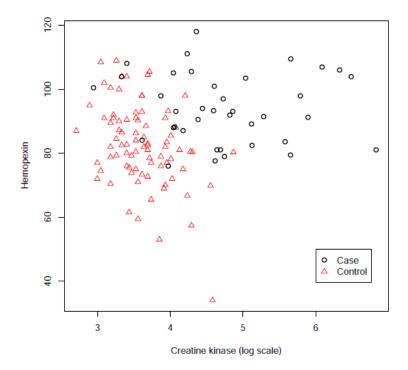
$$\log\left(\frac{\hat{\pi}}{1-\hat{\pi}}\right) = 10.875 - 0.1713(31) = 5.56$$

(This corresponds to an estimated probability of O-ring failure of 0.9962).

(f) The data used to fit the logistic regression model contained temperatures ranging from 51° F to 81° F. To make a prediction based on an observation falling so far from these data is an extrapolation and can yield very inaccurate results.

20.12

- (a) Scatterplot provided on the following page.
 - > ex2012\$lck <- with(ex2012,log(CK))
 - > attach(ex2012)
 - > plot(H~lck,xlab="Creatine kinase (log scale)",ylab="Hemopexin",
 col=as.numeric(GROUP),pch=as.numeric(GROUP))
 - > legend(6,50,col=c(1,2),pch=c(1,2),legend=c("Case","Control"))



There is a clear separation between cases and controls, in terms of these two enzymes. So yes, these enzymes will be useful predictors of whether a woman is a carrier.

(b) R output given below:

```
> m1 <- glm(GROUP~CK+I(CK^2),family=binomial)</pre>
  > summary(m1)
  Call:
  glm(formula = GROUP ~ CK + I(CK^2), family = binomial)
  Deviance Residuals:
      Min 1Q Median 3Q
                                             Max
  -2.50614 -0.03892 0.37943 0.51824
                                         2.27518
  Coefficients:
               Estimate Std. Error z value Pr(>|z|)
  (Intercept) 4.181e+00 7.272e-01 5.749 8.96e-09 ***
            -5.805e-02 1.301e-02 -4.460 8.18e-06 ***
  I(CK^2)
             5.060e-05 3.286e-05 1.540
                                             0.124
  Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
      Null deviance: 149.840 on 119 degrees of freedom
  Residual deviance: 85.435 on 117 degrees of freedom
  AIC: 91.435
  The CK-squared term does not differ significantly from 0.
R output given below:
> m2 <- glm(GROUP~lck+I(lck^2),family=binomial)</pre>
> summary(m2)
Call:
glm(formula = GROUP ~ lck + I(lck^2), family = binomial)
Deviance Residuals:
          1Q Median
    Min
                                3Q
                                        Max
-2.39251 -0.03075 0.38037 0.50190 2.28852
Coefficients:
       Estimate Std. Error z value Pr(>|z|)
(Intercept) -9.830 16.309 -0.603 0.547
                     8.366 1.024
            8.568
                                    0.306
lck
I(lck^2)
            -1.453
                     1.064 -1.365
                                     0.172
   Null deviance: 149.84 on 119 degrees of freedom
Residual deviance: 84.98 on 117 degrees of freedom
AIC: 90.98
```

The squared term does not significantly differ from 0.

The model with the untransformed CK has more significant p-values for the terms. However, the AIC is smaller for the model with the log-transformed CK. Also, the distribution of log(CK) is more symmetric than the distribution of CK, which is highly skewed to the right.

(c) R output given below:

```
> m3 <- glm(GROUP~lck+H,family=binomial)
> summary(m3)
```

Call:

glm(formula = GROUP ~ lck + H, family = binomial)

Deviance Residuals:

Min 1Q Median 3Q Max -2.60372 -0.09903 0.16697 0.38782 1.89707

Coefficients:

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

(Intercept) 28.91300 5.80030 4.985 6.20e-07 ***

lck -4.02041 0.82909 -4.849 1.24e-06 ***

H -0.13652 0.03654 -3.736 0.000187 ***

--
Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1

Null deviance: 149.840 on 119 degrees of freedom Residual deviance: 61.992 on 117 degrees of freedom

AIC: 67.992

(d) Wish to test H_0 : $\beta_1 = \beta_2 = 0$ vs. H_a : some $\beta_j \neq 0$, j = 1, 2. The test statistic is

$$\chi^2 = 149.840 - 61.992 = 87.848$$

Under H_0 , χ^2 follows a chi-squared distribution on 119 - 117 = 2 degrees of freedom. The *p*-value for this test is **approximately zero**, so we reject H_0 and conclude that $\log(\text{CK})$ and H are useful predictors.

> 1-pchisq(87.848,df=2) [1] 0

(e) The difference in log odds is

$$\begin{split} (28.913 - 4.0204(\log(80)) - 0.1365(85)) - (28.913 - 4.0204(\log(300)) - 0.1365(100)) \\ = 4.0204(\log(300) - \log(80)) + 0.1365(100 - 85) = 7.36 \end{split}$$

so the odds ratio is

$$e^{7.36} = 1574.67$$

The odds she is a carrier are over 1,500 times as great.