

# Logistic Regression

## Spider mites example

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Dose of DDT	No. survived	No. dead
0.0	18	7
0.5	19	6
1.0	12	13
1.5	5	20
2.0	6	19
2.5	2	23
3.0	1	24

# Binary vs. continuous outcomes

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Continuous: ANOVA  $\longleftrightarrow$  Regression

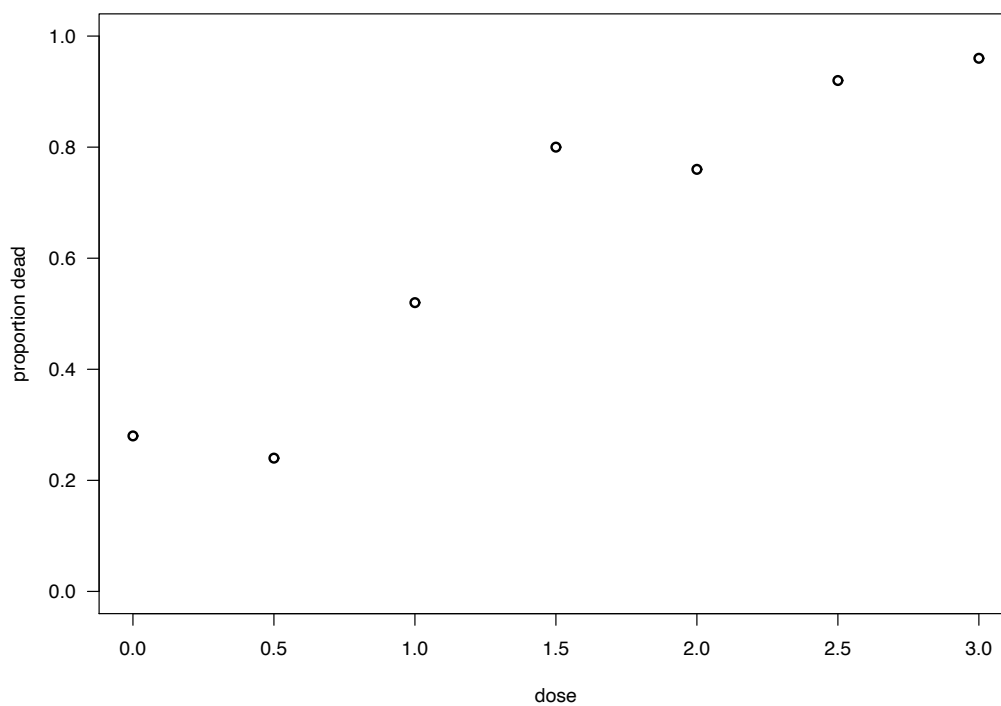
Binary:  $k \times 2$  table  $\longleftrightarrow$  ?

Goals:

- Determine the relationship between dose and  $\text{Pr}(\text{dead})$ .
- Find the dose at which  $\text{Pr}(\text{dead}) = 1/2$ .

## A plot of the data

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# Binary outcomes

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Let  $p_d = \text{Pr}(\text{dead} \mid \text{dose } d)$

$$p_d = \beta_0 + \beta_1 d ?$$

$$0 \leq p_d \leq 1 \quad \text{but} \quad -\infty \leq \beta_0 + \beta_1 d \leq \infty$$

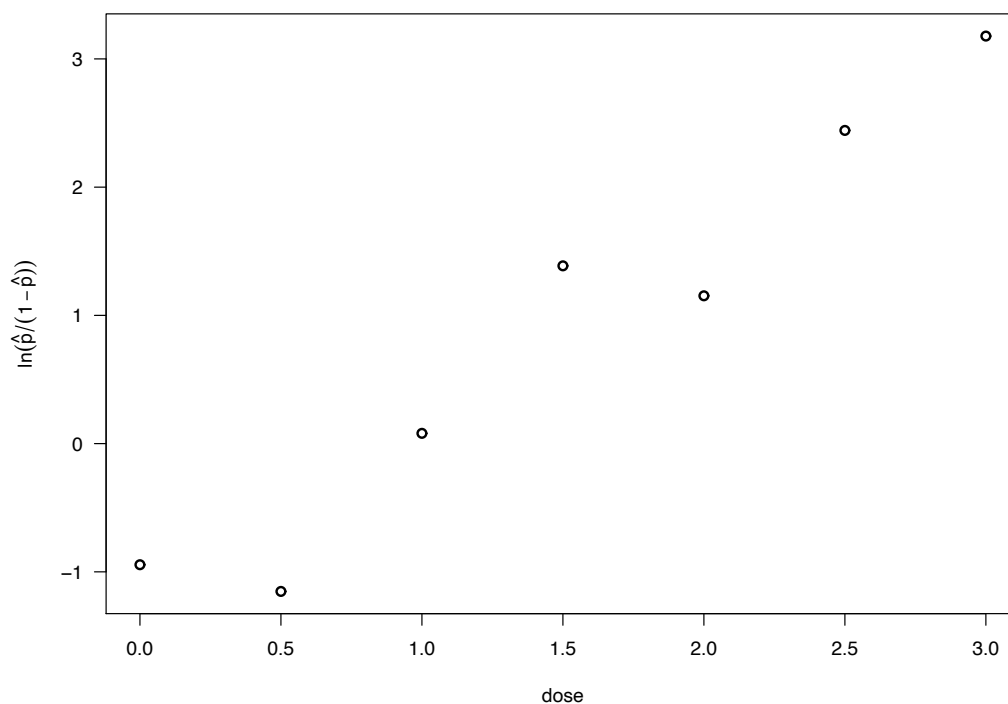
Odds of death:  $0 \leq \frac{p_d}{1 - p_d} \leq \infty$

Log odds of death:  $-\infty \leq \ln\left(\frac{p_d}{1 - p_d}\right) \leq \infty$

→  $\ln\left(\frac{p}{1 - p}\right)$  is also called **logit(p)** or the **logistic function**.

## logit( $\hat{p}_d$ ) vs d

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## Logistic regression

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$$\ln\left(\frac{p_d}{1-p_d}\right) = \beta_0 + \beta_1 d$$

Try least squares, regressing  $\ln\left(\frac{\hat{p}_d}{1-\hat{p}_d}\right)$  on the dose  $d$ ?

Problems:

- What if  $\hat{p}_d = 0$  or  $1$ ?
- $SD(\hat{p}_d)$  is not constant with  $d$ .

## Maximum likelihood

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Assume that

- $y_d \sim \text{Binomial}(n_d, p_d)$ ,
- $y_d$  independent,
- $\text{logit}(p_d) = \ln\left(\frac{p_d}{1-p_d}\right) = \beta_0 + \beta_1 d$

Note: 
$$p_d = \frac{e^{\beta_0 + \beta_1 d}}{1 + e^{\beta_0 + \beta_1 d}}$$

Likelihood:

$$L(\beta_0, \beta_1 | \mathbf{y}) = \prod_d p_d^{y_d} (1 - p_d)^{(n_d - y_d)}$$

# Logistic regression in R

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Logistic regression is a special case of a *generalized linear model*.

Function in R: `glm()`

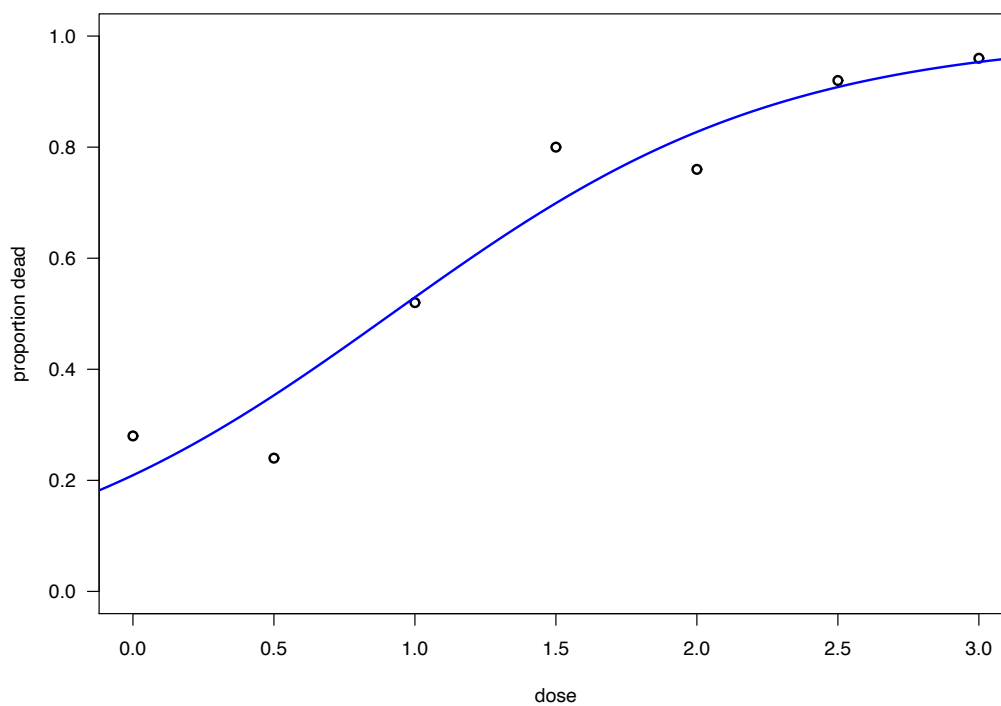
```
> glm.out <- glm(n.dead/n ~ dose, weights=n, data=spiders,  
                family=binomial(link=logit))
```

```
> summary(glm.out)$coef
```

	Est	SE	z-val	P-val
(Intercept)	-1.33	0.33	-4.06	<0.001
dose	1.44	0.23	6.29	<0.001

## Fitted curve

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## Interpretation of $\beta$ 's

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$$\ln\left(\frac{p_d}{1-p_d}\right) = \beta_0 + \beta_1 d$$

$\beta_0$  = log odds when dose = 0

Note:  $\beta_0 = 0 \longrightarrow p_0 = \frac{1}{2}$

$\beta_1$  = change in log odds with unit increase in dose

Note:  $\beta_1 = 0 \longrightarrow$  survival unrelated to dose.

## LD50

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LD50 = dose at which  $\Pr(\text{dead} \mid \text{dose}) = \frac{1}{2}$ .

$$\ln\left(\frac{1/2}{1-1/2}\right) = \beta_0 + \beta_1(\text{LD50})$$

$$0 = \beta_0 + \beta_1(\text{LD50})$$

$$\text{LD50} = -\beta_0/\beta_1$$

$$\widehat{\text{LD50}} = -\hat{\beta}_0/\hat{\beta}_1$$

$$\widehat{\text{SE}}(\widehat{\text{LD50}}) \approx |\widehat{\text{LD50}}| \sqrt{\left(\frac{\widehat{\text{SE}}(\hat{\beta}_0)}{\hat{\beta}_0}\right)^2 + \left(\frac{\widehat{\text{SE}}(\hat{\beta}_1)}{\hat{\beta}_1}\right)^2 - 2 \frac{\widehat{\text{cov}}(\hat{\beta}_0, \hat{\beta}_1)}{\hat{\beta}_0 \hat{\beta}_1}}$$

# Estimating LD50 in R

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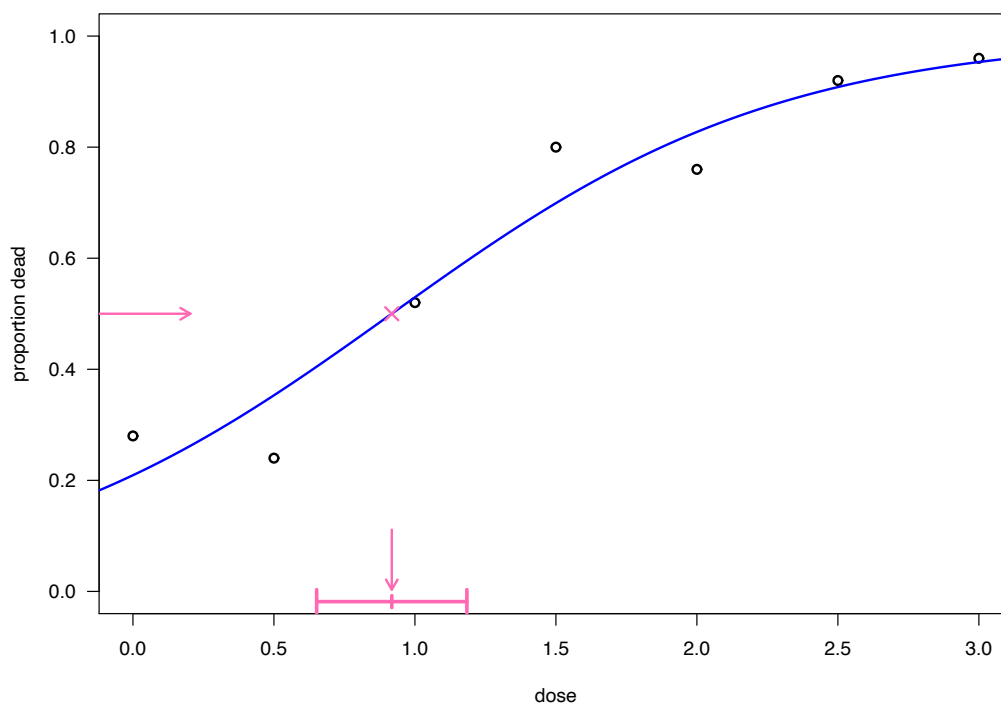
```
> glm.out <- glm(n.dead/n ~ dose, weights=n, data=spiders,
  family=binomial(link=logit))
> glm.sum <- summary(glm.out)

> co <- glm.out$coef
> ld50 <- -co[1]/co[2]
> se.co <- glm.sum$coef[,2]
> cov.co <- glm.sum$cov.scaled[1,2]
> se.ld50 <- abs(ld50) * sqrt( (se.co[1]/co[1])^2 +
  (se.co[2]/co[2])^2 -
  2*cov.co/(co[1]*co[2]) )

> ld50
  0.92
> se.ld50
  0.14
> ld50 + c(-1,1) * qnorm(0.975) * se.ld50
  0.65  1.18
```

## LD50

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## Another example

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Tobacco budworm, *Heliothis virescens*

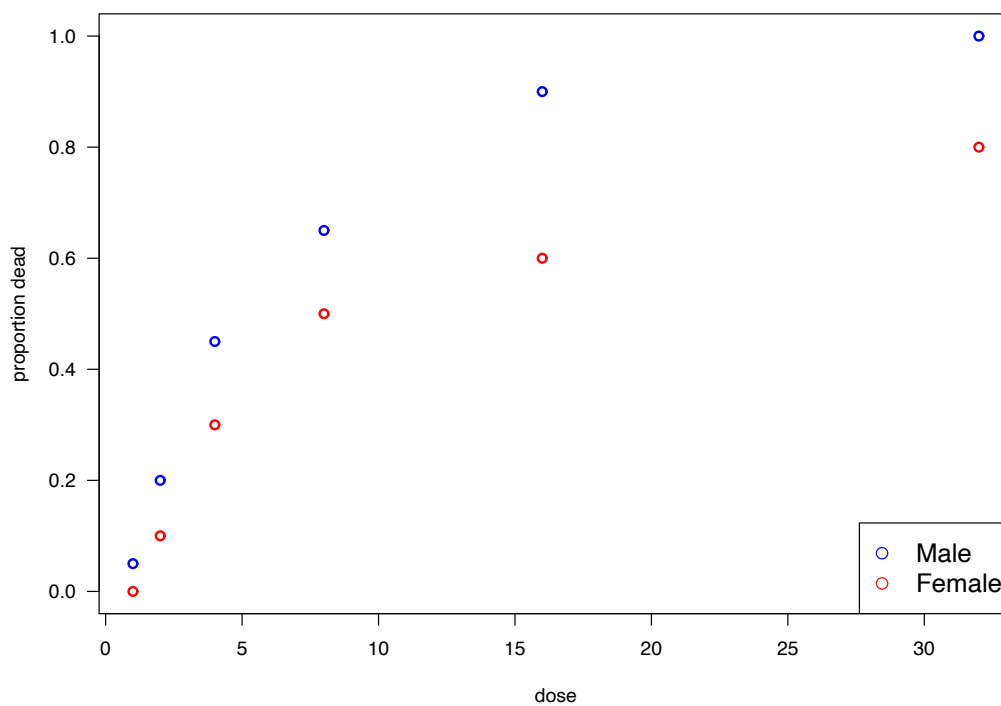
Batches of 20 male and 20 female worms were given a 3-day dose of pyrethroid *trans*-cypermethrin

The no. dead or “knocked down” in each batch was noted.

	Dose					
Sex	1	2	4	8	16	32
Male	1	4	9	13	18	20
Female	0	2	6	10	12	16

## A plot of the data

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# Analysis in R

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## Assume no sex difference

```
> glm.out <- glm(n.dead/n ~ dose, weights=n, data=worms,  
                family=binomial(link=logit))
```

```
> summary(glm.out)$coef  
              Est      SE  z-val  P-val  
(Intercept) -1.57   0.23  -6.8  <0.001  
dose         0.153  0.022   6.8  <0.001
```

## Assume sexes completely different

```
> glm.outB <- glm(n.dead/n ~ sex*dose, weights=n, data=worms,  
                 family=binomial(link=logit))
```

```
> summary(glm.outB)$coef  
              Est      SE  z-val  P-val  
(Intercept) -1.72   0.32  -5.3  <0.001  
sexmale      -0.21   0.51  -0.4   0.68  
dose         0.116  0.024   4.9  <0.001  
sexmale:dose 0.182  0.067   2.7   0.007
```

# Analysis in R (continued)

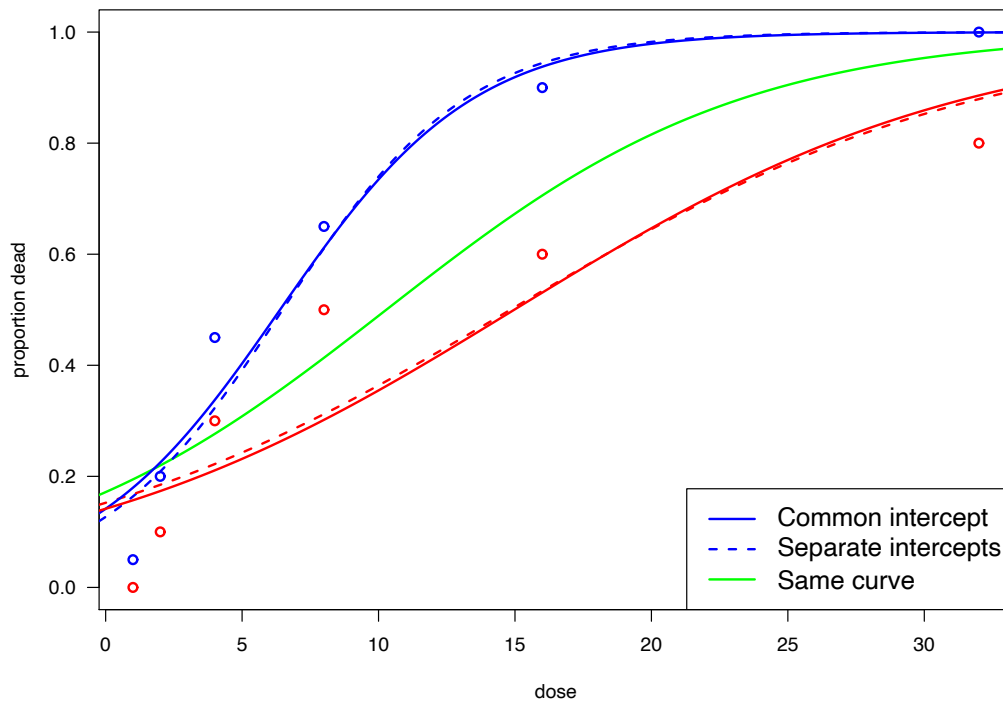
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## Different slopes but common “intercept”

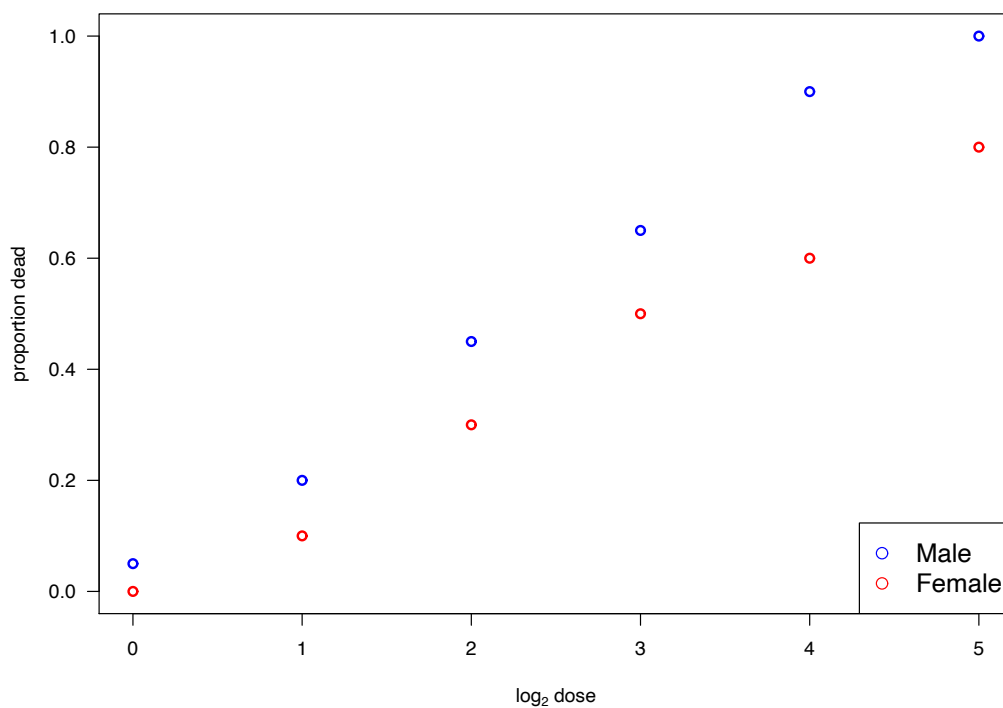
```
> glm.outC <- glm(n.dead/n ~ dose + sex:dose, weights=n,  
                 data=worms, family=binomial(link=logit))
```

```
> summary(glm.out)$coef  
              Est      SE  z-val  P-val  
(Intercept) -1.80   0.25  -7.2  <0.001  
dose         0.120  0.021   5.6  <0.001  
dose:sexmale 0.161  0.044   3.7  <0.001
```

## Fitted curves



## Plot using $\log_2$ dose



## Use $\log_2$ of the dose

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### Assume no sex difference

```
> glm.out <- glm(n.dead/n ~ dose, weights=n, data=worms,  
                family=binomial(link=logit))
```

```
> summary(glm.out)$coef  
              Est      SE    z-val    P-val  
(Intercept) -2.77  0.37    -7.6   <0.001  
dose          1.01  0.12     8.1   <0.001
```

### Assume sexes completely different

```
> glm.outB <- glm(n.dead/n ~ sex*dose, weights=n, data=worms,  
                 family=binomial(link=logit))
```

```
> summary(glm.outB)$coef  
              Est      SE    z-val    P-val  
(Intercept) -2.99  0.55    -5.4   <0.001  
sexmale       0.17  0.78    -0.2    0.82  
dose          0.91  0.17     5.4   <0.001  
sexmale:dose  0.35  0.27     1.3    0.19
```

## Use $\log_2$ of the dose (continued)

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### Different slopes but common “intercept”

```
> glm.outC <- glm(n.dead/n ~ dose + sex:dose, weights=n,  
                 data=worms, family=binomial(link=logit))
```

```
> summary(glm.out)$coef  
              Est      SE    z-val    P-val  
(Intercept) -2.91  0.39    -7.5   <0.001  
dose          0.88  0.13     6.9   <0.001  
dose:sexmale  0.41  0.12     3.3    0.001
```

# Fitted curves

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