Friday, January 31

Marginal Means

A marginal mean is effectively an average of expected responses. The **emmeans** package is particularly useful for making inferences about marginal means. It can also be done using **contrast** but it is not a documented feature.

```
library(trtools)
library(emmeans)
```

```
Welcome to emmeans.
Caution: You lose important information if you filter this package's results.
See '? untidy'
```

```
Attaching package: 'emmeans'
```

The following objects are masked from 'package:trtools':

```
contrast, neuralgia
```

Warning: The emmeans package contains a function called contrast which is not the same as the function of the same name in the **trtools** package, resulting in a namespace conflict if both packages are loaded. If you have both packages loaded in a given session, use trtools::contrast and emmeans::contrast to refer to a given function.

Example: Consider again the data from the platyfish study.

```
m <- lm(Percentage ~ Pair, data = Sleuth3::case0602)
summary(m)$coefficients</pre>
```

	Estimate Std.	Error	t value	Pr(> t)
(Intercept)	56.406	3.864	14.5965	5.208e-24
PairPair2	4.479	5.657	0.7919	4.308e-01
PairPair3	6.023	5.384	1.1187	2.667e-01
PairPair4	10.594	5.657	1.8727	6.485e-02
PairPair5	7.805	6.441	1.2118	2.292e-01
PairPair6	6.929	5.657	1.2250	2.243e-01

We see that there are indicator variables for male pairs 2-6. The model can be written as

```
E(Y_i) = \begin{cases} \beta_0, & \text{if the } i\text{-th observation was from the first male pair,} \\ \beta_0 + \beta_1, & \text{if the } i\text{-th observation was from the second male pair,} \\ \beta_0 + \beta_2, & \text{if the } i\text{-th observation was from the third male pair,} \\ \beta_0 + \beta_3, & \text{if the } i\text{-th observation was from the fourth male pair,} \\ \beta_0 + \beta_4, & \text{if the } i\text{-th observation was from the fifth male pair,} \\ \beta_0 + \beta_5, & \text{if the } i\text{-th observation was from the sixth male pair.} \end{cases}
```

We can use contrast to estimate the expected response for each pair.

Note how I used a shortcut to specify the pairs.

paste("Pair", 1:6, sep = "")

[1] "Pair1" "Pair2" "Pair3" "Pair4" "Pair5" "Pair6"

This can also be done using the emmeans function from the package emmeans.

```
library(emmeans)
emmeans(m, ~ Pair)
```

Pair	emmean	SE	df	lower.CL	upper.CL
Pair1	56.4	3.86	78	48.7	64.1
Pair2	60.9	4.13	78	52.7	69.1
Pair3	62.4	3.75	78	55.0	69.9
Pair4	67.0	4.13	78	58.8	75.2
Pair5	64.2	5.15	78	54.0	74.5
Pair6	63.3	4.13	78	55.1	71.6

Confidence level used: 0.95

Denote the six expected responses (one for each pair) as

$$\begin{split} \mu_1 &= \beta_0, \\ \mu_2 &= \beta_0 + \beta_1, \\ \mu_3 &= \beta_0 + \beta_2, \\ \mu_4 &= \beta_0 + \beta_3, \\ \mu_5 &= \beta_0 + \beta_4, \\ \mu_6 &= \beta_0 + \beta_5. \end{split}$$

One marginal mean would be the average expected response across the pairs. This could be written as

$$\mu = \frac{\mu_1 + \mu_2 + \mu_3 + \mu_4 + \mu_5 + \mu_6}{6} = \beta_0 + \frac{1}{6}\beta_1 + \frac{1}{6}\beta_2 + \frac{1}{6}\beta_3 + \frac{1}{6}\beta_4 + \frac{1}{6}\beta_5$$

We can estimate this quantity with lincon.

lincon(m, a = c(1, 1/6, 1/6, 1/6, 1/6, 1/6))

estimate se lower upper tvalue df pvalue (1,1/6,1/6,1/6,1/6),0 62.38 1.722 58.95 65.81 36.23 78 1.501e-50

We can also use emmeans.

emmeans(m, ~ 1)

1 emmean SE df lower.CL upper.CL overall 62.4 1.72 78 59 65.8

Results are averaged over the levels of: Pair Confidence level used: 0.95

Note that we can use the confidence interval to test the null hypothesis that $\mu = 50$. For a test statistic and p-value for this test we could write this as

 $\mu = 50 \Leftrightarrow \beta_0 + \frac{1}{6}\beta_1 + \frac{1}{6}\beta_2 + \frac{1}{6}\beta_3 + \frac{1}{6}\beta_4 + \frac{1}{6}\beta_5 = 50 \Leftrightarrow \beta_0 + \frac{1}{6}\beta_1 + \frac{1}{6}\beta_2 + \frac{1}{6}\beta_3 + \frac{1}{6}\beta_4 + \frac{1}{6}\beta_5 - 50 = 0.$

Here is how we can do that with lincon.

lincon(m, a = c(1, 1/6, 1/6, 1/6, 1/6, 1/6), b = -50)

estimate se lower upper tvalue df pvalue (1,1/6,1/6,1/6,1/6,1/6),-50 12.38 1.722 8.95 15.81 7.189 78 3.439e-10 Here is how we do it with emmeans. emmeans(m, ~ 1, offset = -50, infer = TRUE) 1 emmean SE df lower.CL upper.CL t.ratio p.value overall 12.4 1.72 78 8.95 15.8 7.189 <.0001

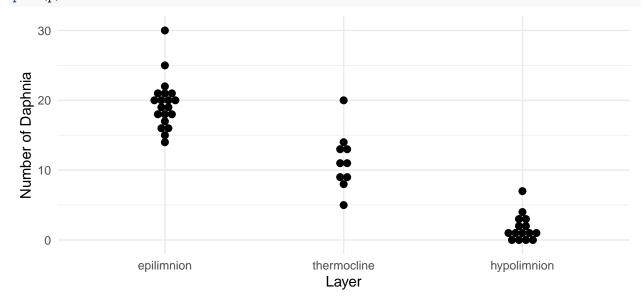
Results are averaged over the levels of: Pair Confidence level used: 0.95

By *not* listing an explanatory variable on the right-hand side of \sim , we are asking that emmeans average over that explanatory variable. Also note that the argument infer = TRUE makes the emmeans function provide both confidence intervals as well as tests.

Note: If we just want to know whether or not we would reject the null hypothesis that $\mu = 50$ we can also just look at the confidence interval for μ .

Example: Consider the following data from a survey of water fleas.

```
library(ggplot2)
p <- ggplot(daphniastrat, aes(x = layer, y = count)) +
  geom_dotplot(binaxis = "y", stackdir = "center") +
  labs(x = "Layer", y = "Number of Daphnia") + theme_minimal()
plot(p)</pre>
```



We might model these data using the following linear model.

```
m <- lm(count ~ layer, data = daphniastrat)
summary(m)$coefficients</pre>
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	19.50	0.7271	26.820	4.727e-28
layerthermocline	-8.20	1.2593	-6.512	7.293e-08
layerhypolimnion	-17.77	1.1106	-15.997	1.784e-19

So our model can be written as

 $E(Y_i) = \begin{cases} \beta_0, & \text{if the } i\text{-th observation is from the epilimnion layer,} \\ \beta_0 + \beta_1, & \text{if the } i\text{-th observation is from the thermocline layer,} \\ \beta_0 + \beta_2, & \text{if the } i\text{-th observation is from the hypolimnion layer.} \end{cases}$

Let μ_e , μ_t , and μ_h denote the expected number of daphnia per liter for the epilimnion, thermocline, and hypolimnion layers, respectively (i.e., the density of daphnia in each layer). So

$$\mu_e = \beta_0, \ \mu_t = \beta_0 + \beta_1, \ \mu_h = \beta_0 + \beta_2.$$

It is known that the volumes of the epilimnion, thermocline, and hypolimnion layers are 100, 200, and 400 kL, respectively. The density for the entire lake is then

$$\mu = \frac{100}{700}\mu_e + \frac{200}{700}\mu_t + \frac{400}{700}\mu_h = \beta_0 + \frac{2}{7}\beta_1 + \frac{4}{7}\beta_2.$$

We can estimate this with lincon or emmeans using the weights option.

```
lincon(m, a = c(1, 2/7, 4/7))
```

```
estimate se lower upper tvalue df pvalue
(1,2/7,4/7),0 7.005 0.572 5.85 8.159 12.25 42 1.907e-15
emmeans(m, ~ 1, weights = c(1/7, 2/7, 4/7))
```

```
        1
        emmean
        SE df
        lower.CL
        upper.CL

        overall
        7
        0.572
        42
        5.85
        8.16
```

```
Results are averaged over the levels of: layer
Confidence level used: 0.95
```

Note that when using **emmeans** it is important to put the weights in the correct order. We can verify the order using **level** (if the variable is a factor) or by using **weights = "slow.levels"**.

```
levels(daphniastrat$layer)
```

```
[1] "epilimnion" "thermocline" "hypolimnion"
emmeans(m, ~ 1, weights = "show.levels")
```

emmeans are obtained by averaging over these factor combinations

```
layer
1 epilimnion
2 thermocline
3 hypolimnion
```

We can estimate the expected number of daphnia per liter for each layer.

```
emmeans(m, ~ layer)
```

layeremmeanSEdflower.CLupper.CLepilimnion19.500.7274218.03320.97thermocline11.301.030429.22513.38hypolimnion1.730.840420.0393.43

```
Confidence level used: 0.95
```

```
trtools::contrast(m, a = list(layer = c("epilimnion","thermocline","hypolimnion")),
    cnames = c("epilimnion","thermocline","hypolimnion"))
```

lower upper tvalue df estimate pvalue se 19.500 0.7271 18.03274 20.967 26.820 42 4.727e-28 epilimnion 11.300 1.0282 thermocline 9.22498 13.375 10.990 42 6.221e-14 1.733 0.8395 0.03909 3.428 2.065 42 4.517e-02 hypolimnion We can also do inferences concerning the differences between pairs of layers. pairs(emmeans(m, ~ layer), adjust = "none") contrast estimate SE df t.ratio p.value epilimnion - thermocline 8.20 1.26 42 6.512 <.0001 17.77 1.11 42 15.997 epilimnion - hypolimnion <.0001 thermocline - hypolimnion 9.57 1.33 42 7.207 <.0001 trtools::contrast(m, a = list(layer = c("epilimnion","epilimnion","thermocline")), b = list(layer = c("thermocline", "hypolimnion", "hypolimnion")), cnames = c("E-T","E-H", "T-H")) se lower upper tvalue df estimate pvalue E-T 8.200 1.259 5.659 10.74 6.512 42 7.293e-08

E-18.2001.2395.03910.740.312427.293e-08E-H17.7671.11115.52520.0115.997421.784e-19T-H9.5671.3276.88812.257.207427.363e-09

The adjust = "none" option for pairs specifies that no adjustment be made to confidence intervals or tests for the family-wise Type I error rate.¹

Something to note when using the weights argument with the emmeans function is that the weights that are used must sum to one, and if they do not they will be normalized so that they do. For example, the following provide the same result.

emmeans(m, ~ 1, weights = c(1/7, 2/7, 4/7))

 1
 emmean
 SE
 df
 lower.CL
 upper.CL

 overall
 7
 0.572
 42
 5.85
 8.16

Results are averaged over the levels of: layer Confidence level used: 0.95

emmeans(m, ~ 1, weights = c(1, 2, 4)) # original weights multiplied by 7

1emmeanSE df lower.CL upper.CLoverall70.572425.858.16

Results are averaged over the levels of: layer Confidence level used: 0.95

If you want to use weights that do not sum to one, you can use the **contrast** function from the **emmeans** package (different from the function of the same name from **trtools**).

¹The family-wise Type I error rate is the probability of making at least one Type I error. If it is desired that the family-wise Type I error rate be no greater than α (default is 0.05), then some adjustment can be made. This adjustment is seen in the p-values and confidence intervals. The most general method is to use adjust = "mvt". Some special cases are more widely known such as Tukey (adjust = "tukey") and Bonferroni (adjust = "bonferroni"), but the adjustment based on the multivariate t-distribution (adjust = "mvt") is the most general and accurate. Note that an adjustment will produce "simultaneous" confidence intervals. A method of producing simultaneous confidence intervals has the property that the probability that all of the confidence intervals will contain the quantities being estimated is equal to the specified confidence level (95% by default). The multivariate t-distribution adjustment is perhaps not as well known, so a reference that you can cite is Edwards, D. & Berry, J. T. (1987). The efficiency of simulation-based multiple comparisons. Biometrics, 43(4), 913–928.

emmeans(m, ~1, weights = c(1/7, 2/7, 4/7)) SE df lower.CL upper.CL 1 emmean 7 0.572 42 5.85 8.16 overall Results are averaged over the levels of: layer Confidence level used: 0.95 emmeans::contrast(emmeans(m, ~layer), method = list(layer = c(1/7, 2/7, 4/7)), infer = TRUE) SE df lower.CL upper.CL t.ratio p.value contrast estimate 8.16 12.245 <.0001 7 0.572 42 5.85 layer

Confidence level used: 0.95

But suppose we wanted to estimate the *number* of daphnia in the lake (τ) . It can be shown that this is

 $\tau = 700000\mu = 700000 \left(\frac{1}{7}\mu_e + \frac{2}{7}\mu_t + \frac{4}{7}\mu_h\right) = 100000\mu_e + 200000\mu_t + 400000\mu_h.$

Note that there are 700kL in the lake, which is 700000L (which is the scale used for the observations). This can be estimated as follows.

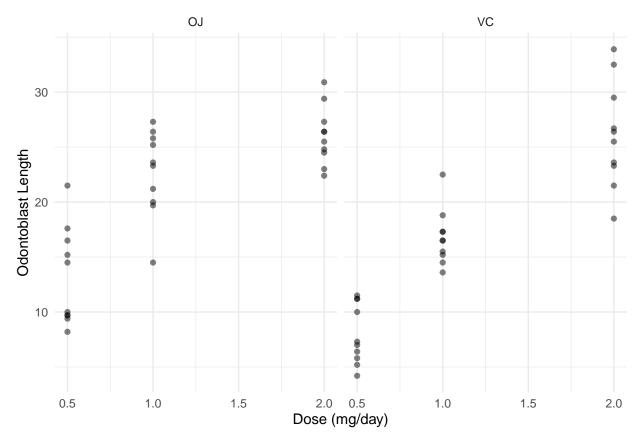
Confidence level used: 0.95

Another approach is to use lincon but your weights/coefficients will be different since they are applied to β_0 , β_1 , and β_2 .

Marginal Means and "Main Effects"

Consider data from a randomized experiment with guinea pigs administered one of three doses of vitamin C (0.5, 1, or 2 mg/day) via one of two supplement methods: orange juice (OJ) or ascorbic acid (VC).

```
p <- ggplot(ToothGrowth, aes(x = dose, y = len)) +
geom_point(alpha = 0.5) + facet_wrap(~supp) +
labs(x = "Dose (mg/day)", y = "Odontoblast Length") + theme_minimal()
plot(p)</pre>
```



Here we are going to model dose as a categorical variable so we need to coerce it to a factor. Perhaps the safest approach is to create a new variable.

ToothGrowth\$dosef <- factor(ToothGrowth\$dose)</pre>

Note: Whether a variable is a numeric, a factor, or something else can be seen use str (for "structure").

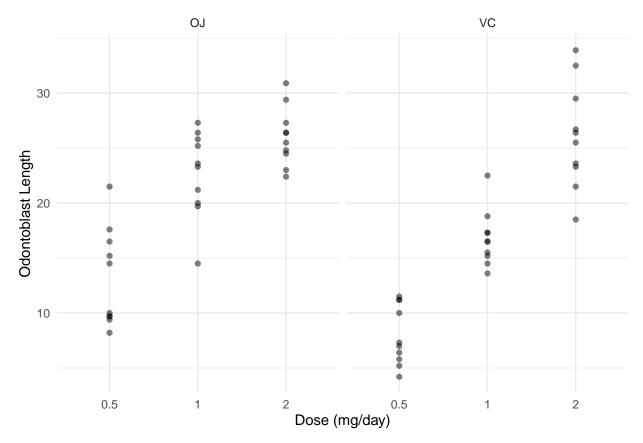
str(ToothGrowth)

```
'data.frame': 60 obs. of 4 variables:
$ len : num 4.2 11.5 7.3 5.8 6.4 10 11.2 11.2 5.2 7 ...
$ supp : Factor w/ 2 levels "OJ","VC": 2 2 2 2 2 2 2 2 2 2 2 ...
$ dose : num 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 ...
$ dosef: Factor w/ 3 levels "0.5","1","2": 1 1 1 1 1 1 1 1 1 ...
```

Notice that ggplot responds differently.

```
summary(ToothGrowth)
```

```
len
                supp
                              dose
                                        dosef
Min.
        : 4.2
                OJ:30
                        Min.
                                :0.50
                                        0.5:20
 1st Qu.:13.1
                VC:30
                         1st Qu.:0.50
                                        1 :20
Median :19.2
                         Median :1.00
                                        2 :20
Mean
       :18.8
                         Mean
                               :1.17
 3rd Qu.:25.3
                         3rd Qu.:2.00
Max.
        :33.9
                         Max.
                                :2.00
p <- ggplot(ToothGrowth, aes(x = dosef, y = len)) +</pre>
  geom_point(alpha = 0.5) + facet_wrap(~supp) +
  labs(x = "Dose (mg/day)", y = "Odontoblast Length") + theme_minimal()
plot(p)
```



Now consider the following linear model.

 x_{i4}

m <- lm(len ~ dosef + supp + dosef:supp, data = ToothGrowth)
summary(m)\$coefficients</pre>

	Estimate S	Std.	Error	t value	Pr(> t)
(Intercept)	13.23		1.148	11.5208	3.603e-16
dosef1	9.47		1.624	5.8312	3.176e-07
dosef2	12.83		1.624	7.9002	1.430e-10
suppVC	-5.25		1.624	-3.2327	2.092e-03
dosef1:suppVC	-0.68		2.297	-0.2961	7.683e-01
dosef2:suppVC	5.33		2.297	2.3207	2.411e-02

The model is

$$E(Y_i) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3} + \beta_4 x_{i4} + \beta_5 x_{i5}$$

where

$$\begin{aligned} x_{i1} &= \begin{cases} 1, & \text{if dose is 1 mg/day,} \\ 0, & \text{otherwise,} \end{cases} \\ x_{i2} &= \begin{cases} 1, & \text{if dose is 2 mg/day,} \\ 0, & \text{otherwise,} \end{cases} \\ x_{i3} &= \begin{cases} 1, & \text{if supplement type is VC} \\ 0, & \text{otherwise,} \end{cases} \\ &= x_{i1}x_{i3} = \begin{cases} 1, & \text{if dose is 1 mg/day and supplement type is VC,} \\ 0, & \text{otherwise,} \end{cases} \end{aligned}$$

$$x_{i5} = x_{i2}x_{i3} = \begin{cases} 1, & \text{if dose is } 2 \text{ mg/day and supplement type is VC,} \\ 0, & \text{otherwise.} \end{cases}$$

We can write this model case-wise.

.

$$E(Y_i) = \begin{cases} \beta_0, & \text{if dose is } 0.5 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_1, & \text{if dose is } 1 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_2, & \text{if dose is } 2 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_3, & \text{if dose is } 0.5 \text{ mg/day and supplement type is VC,} \\ \beta_0 + \beta_1 + \beta_3 + \beta_4, & \text{if dose is } 1 \text{ mg/day and supplement type is VC,} \\ \beta_0 + \beta_2 + \beta_3 + \beta_5, & \text{if dose is } 2 \text{ mg/day and supplement type is VC.} \end{cases}$$

Note that if we omitted the interaction term so that the model formula is $len \sim dosef + supp$, then we would have the model

$$E(Y_i) = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3},$$

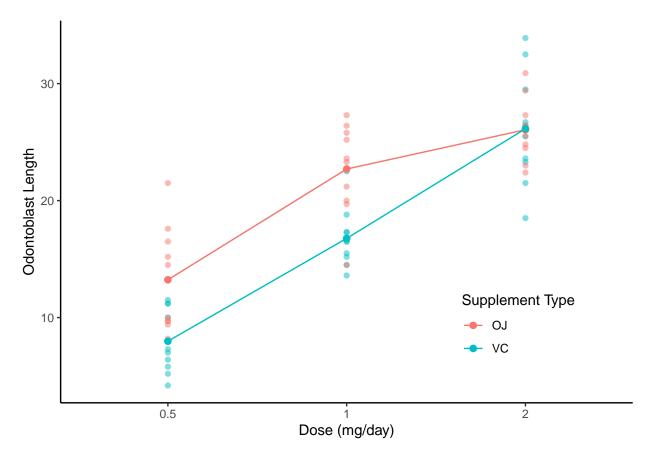
which can be written case-wise as

	$\beta_0,$	if dose is 0.5 mg/day and supplement type is OJ, if dose is 1 mg/day and supplement type is OJ, if dose is 2 mg/day and supplement type is OJ, if dose is 0.5 mg/day and supplement type is VC, if dose is 1 mg/day and supplement type is VC, if dose is 2 mg/day and supplement type is VC.
$\beta_0 + \beta_0 $	$\beta_0 + \beta_1,$	if dose is 1 mg/day and supplement type is OJ,
	$\beta_0 + \beta_2,$	if dose is 2 mg/day and supplement type is OJ,
$L(I_i) =$	$\beta_0 + \beta_3,$	if dose is 0.5 mg/day and supplement type is VC,
	$\beta_0 + \beta_1 + \beta_3$	if dose is 1 mg/day and supplement type is VC,
	$\beta_0 + \beta_2 + \beta_3$	if dose is 2 mg/day and supplement type is VC.

Here is a visualization of the data and the model with the interaction.

```
d <- expand.grid(dosef = levels(ToothGrowth$dosef), supp = levels(ToothGrowth$supp))
d$yhat <- predict(m, newdata = d)</pre>
```

```
p <- ggplot(ToothGrowth, aes(x = dosef, y = len, color = supp)) +
geom_point(alpha = 0.5) + theme_classic() +
theme(legend.position = "inside", legend.position.inside = c(0.8,0.2)) +
geom_point(aes(y = yhat), size = 2, data = d) +
geom_line(aes(y = yhat, group = supp), data = d) +
labs(x = "Dose (mg/day)", y = "Odontoblast Length", color = "Supplement Type")
plot(p)</pre>
```



Consider again the model with the interaction so that

$$E(Y_i) = \begin{cases} \beta_0, & \text{if dose is } 0.5 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_1, & \text{if dose is } 1 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_2, & \text{if dose is } 2 \text{ mg/day and supplement type is OJ,} \\ \beta_0 + \beta_3, & \text{if dose is } 0.5 \text{ mg/day and supplement type is VC,} \\ \beta_0 + \beta_1 + \beta_3 + \beta_4, & \text{if dose is } 1 \text{ mg/day and supplement type is VC,} \\ \beta_0 + \beta_2 + \beta_3 + \beta_5, & \text{if dose is } 2 \text{ mg/day and supplement type is VC.} \end{cases}$$

The "cell means" are

$$\mu_{\rm OJ,0.5} = \beta_0,\tag{1}$$

$$\mu_{\mathrm{OJ},1.0} = \beta_0 + \beta_1,\tag{2}$$

$$\mu_{\rm OJ,2.0} = \beta_0 + \beta_2,\tag{3}$$

$$\mu_{\rm VC,0.5} = \beta_0 + \beta_3,\tag{4}$$

$$\mu_{\rm VC,1.0} = \beta_0 + \beta_1 + \beta_3 + \beta_4,\tag{5}$$

$$\mu_{\rm VC,2.0} = \beta_0 + \beta_1 + \beta_3 + \beta_5. \tag{6}$$

The "marginal means" for supplement type are

$$\mu_{\rm OJ} = \frac{\mu_{\rm OJ,0.5} + \mu_{\rm OJ,1.0} + \mu_{\rm OJ,2.0}}{3} = \beta_0 + \frac{1}{3}\beta_1 + \frac{1}{3}\beta_2,$$

and

$$\mu_{\rm VC} = \frac{\mu_{\rm VC,0.5} + \mu_{\rm VC,1.0} + \mu_{\rm OJ,2.0}}{3} = \beta_0 + \frac{1}{3}\beta_1 + \frac{1}{3}\beta_2 + \beta_3 + \frac{1}{3}\beta_4 + \frac{1}{3}\beta_5.$$

We can estimate them using lincon.

lincon(m, a = c(1, 1/3, 1/3, 0, 0, 0))

estimate se lower upper tvalue df pvalue (1,1/3,1/3,0,0,0),0 20.66 0.663 19.33 21.99 31.17 54 3.359e-36 lincon(m, a = c(1,1/3,1/3,1,1/3,1/3))

estimate se lower upper tvalue df pvalue (1,1/3,1/3,1,1/3,1/3),0 16.96 0.663 15.63 18.29 25.59 54 7.306e-32

But we can also do it using emmeans.

emmeans(m, ~supp)

 supp
 emmean
 SE
 df
 lower.CL
 upper.CL

 OJ
 20.7
 0.663
 54
 19.3
 22.0

 VC
 17.0
 0.663
 54
 15.6
 18.3

Results are averaged over the levels of: dosef Confidence level used: 0.95

Now suppose we want to estimate the "main effect" which is

$$\mu_{\rm OJ} - \mu_{\rm VC} = \frac{\mu_{\rm OJ,0.5} + \mu_{\rm OJ,1.0} + \mu_{\rm OJ,2.0}}{3} - \frac{\mu_{\rm VC,0.5} + \mu_{\rm VC,1.0} + \mu_{\rm OJ,2.0}}{3} = -\beta_3 - \frac{1}{3}\beta_4 - \frac{1}{3}\beta_5.$$

We can do this using lincon.

lincon(m, a = c(0,0,0,-1,-1/3,-1/3))

estimate se lower upper tvalue df pvalue (0,0,0,-1,-1/3,-1/3),0 3.7 0.9376 1.82 5.58 3.946 54 0.0002312

But we can also use functions from the **emmeans** package.

pairs(emmeans(m, ~supp), infer = TRUE)

contrast estimateSE df lower.CL upper.CL t.ratio p.valueOJ - VC3.7 0.938 541.825.583.9460.0002

Results are averaged over the levels of: dosef Confidence level used: 0.95

The main effect of dose concerns differences among the marginal means of dose defined as $\mu_{0.5}$, μ_1 and μ_2 where

 $\mu_{0.5} = \frac{\mu_{\text{OJ},0.5} + \mu_{\text{VC},0.5}}{2}, \quad \mu_1 = \frac{\mu_{\text{OJ},1} + \mu_{\text{VC},1}}{2}, \quad \mu_2 = \frac{\mu_{\text{OJ},2} + \mu_{\text{VC},2}}{2}.$

emmeans(m, ~ dosef)

dosefemmeanSEdflower.CLupper.CL0.510.60.812548.9812.2119.70.8125418.1121.4226.10.8125424.4727.7

Results are averaged over the levels of: supp Confidence level used: 0.95

pairs(emmeans(m, ~ dosef), adjust = "none")

contrast estimate SE df t.ratio p.value dosef0.5 - dosef1 -9.13 1.15 54 -7.951 <.0001 dosef0.5 - dosef2 -15.49 1.15 54 -13.493 <.0001 dosef1 - dosef2 -6.37 1.15 54 -5.543 <.0001

Results are averaged over the levels of: supp

Main Effects in Anova Tables

In ANOVA tables the test of the "main effect" is the (joint) null hypothesis that all pairwise differences are zero. For the variable dose the null hypothesis is $\mu_{0.5} = \mu_1 = \mu_2$. This can be done using the test function.

```
test(pairs(emmeans(m, ~ dosef)), joint = TRUE)
```

```
df1 df2 F.ratio p.value note
2 54 92.000 <.0001 d
```

d: df1 reduced due to linear dependence

This is the traditional main effect that is sometimes reported in an "ANOVA table" such as that produced by Anova from the **car** package.

```
library(car)
m <- lm(len ~ dosef + supp + dosef:supp, data = ToothGrowth,</pre>
  contrast = list(dosef = contr.sum, supp = contr.sum))
Anova(m, type = 3)
Anova Table (Type III tests)
Response: len
            Sum Sq Df F value Pr(>F)
(Intercept) 21236 1 1610.39 < 2e-16 ***
dosef
              2426 2
                         92.00 < 2e-16 ***
                         15.57 0.00023 ***
               205 1
supp
               108 2
                          4.11 0.02186 *
dosef:supp
Residuals
               712 54
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
The option contrast = list(dosef = contr.sum, supp = contr.sum) is necessary here for the Anova
function to do the correct calculations.<sup>2</sup>
The test of the main effect of supplement method was given by
pairs(emmeans(m, ~ supp), infer = TRUE)
 contrast estimate
                       SE df lower.CL upper.CL t.ratio p.value
OJ - VC
               3.7 0.938 54
                                  1.82
                                           5.58
                                                   3.946 0.0002
Results are averaged over the levels of: dosef
Confidence level used: 0.95
We do not need a joint test here since there are only two marginal means, but here it is anyway.
test(pairs(emmeans(m, ~ supp)), joint = TRUE)
```

df1 df2 F.ratio p.value

 $^{^{2}}$ I am demonstrating here what is sometimes called inferences based on Type III sums of squares. Another common approach is to use what is called Type II sums of squares. This can be done with the **Anova** function with **type = 2**. For inferences based on Type II sums of squares with the functions from the **emmeans** package an extra step is needed (email me for an example if you really want to know how to do it).

1 54 15.572 0.0002