2/9/2010

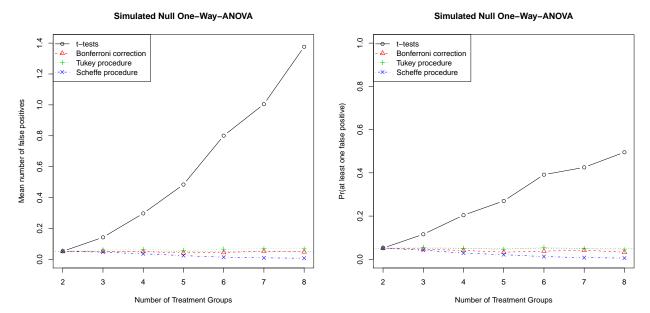
36-402/608 ADA-II H. Seltman Breakout #9 Results

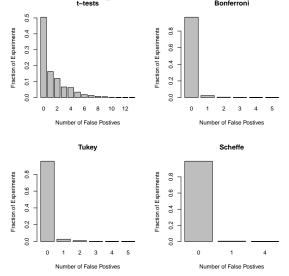
The results below are from simulation of experiments with one factor, which represents between I=2 and and I=8 different treatments to which subjects are randomized. For each experiment the null hypothesis, that $\mu_1 = \cdots = \mu_I$, is true. For each I, 1900 experiments are simulated, so the error for events that occur 5% of the time is +/-1%.

For each I, the number of possible paired tests is I(I-1)/2, which ranges from 1 to 28. For each experiment, all paired t-tests are run, and the number of positive (which must all be false positive) tests is recorded. Over all 1900 experiments the table below shows the mean number of positive tests per experiment and the mean of the indicator variable that at least one test was positive (FWAOP=fraction with at least one positive test). Similarly for each experiment the Bonferroni correction is used to control for multiple testing (based on looking at all pairs), the Tukey procedure is run for all pairs, and the Scheffe procedure is run (which is valid for all possible contrasts simultaneously).

t.test		-Bonferroni-		Tukey		Scheffe	
I Count	FWAOP	Count	FWAOP	\mathtt{Count}	FWAOP	\mathtt{Count}	FWAOP
2 0.052	0.052	0.052	0.052	0.052	0.052	0.052	0.052
3 0.143	0.116	0.052	0.047	0.061	0.055	0.046	0.042
4 0.297	0.204	0.050	0.041	0.064	0.051	0.036	0.029
5 0.484	0.270	0.042	0.033	0.057	0.046	0.024	0.022
6 0.801	0.392	0.044	0.038	0.066	0.055	0.014	0.013
7 1.005	0.425	0.053	0.042	0.068	0.049	0.009	0.008
8 1.376	0.495	0.048	0.033	0.068	0.044	0.007	0.006

Here are plots of all of the counts and of the FWAOP values:





And here is a plot of the distribution of counts for I=8:

Question 1: What do the plots show?

Example to demonstrate tests and code:

```
dtf = read.csv("Experiment.csv")
# treat
            score
# A:8
                : 7.905
        Min.
        1st Qu.: 9.594
# B:8
# C:8
        Median :10.100
# D:8
        Mean
                :10.074
        3rd Qu.:10.538
# E:8
# F:8
        Max.
                :12.225
e0 = aov(score~treat, dtf)
summary(e0)
#
              Df Sum Sq Mean Sq F value
                                             Pr(>F)
                                  6.4438 0.0001581 ***
# treat
               5 69.372 13.8744
# Residuals
              42 90.431
                          2.1531
```

The standard procedure is to "focus" your power by used contrasts that are planned (i.e., chosen *before* looking at the data.

For this experiment, the (orthogonal) planned contrasts, based on the underlying science, are to compare ABC vs. DEF, AB vs. C, DE vs. F, A vs. B, and D vs. E.

```
library(gmodels) # for fit.contrast()
planC = rbind(ABCvsDEF = c(1/3, 1/3, 1/3, -1/3, -1/3, -1/3),
              ABvsC = c(1/2, 1/2, -1, 0, 0, 0), AvsB = c(1, -1, 0, 0, 0, 0),
              DEvsF = c(0,0,0,1/2,1/2,-1), DvsE = c(0,0,0,1,-1,0))
round( fit.contrast(e0, "treat", planC), 3)
#
                 Estimate Std. Error t value Pr(>|t|)
# treatABCvsDEF
                   -0.411
                               0.424
                                      -0.971
                                                  0.337
# treatABvsC
                   -0.720
                               0.635
                                      -1.134
                                                  0.263
# treatAvsB
                   -3.999
                               0.734 -5.450
                                                  0.000
# treatDEvsF
                    0.161
                               0.635
                                        0.253
                                                  0.801
# treatDvsE
                    0.345
                               0.734
                                                  0.641
                                        0.470
```

Question 2: What do we conclude? Why can't we do more testing the same way?

The naive "all t-tests approach" starts with finding biggest mean difference, doing a t-test, and repeating until a non-significant one is found.

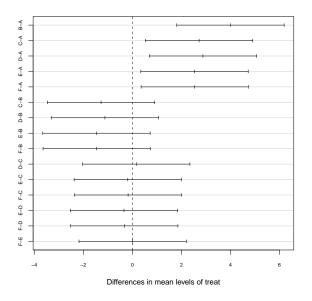
```
with(dtf, tapply(score, treat, mean))
#
         Α
                  В
                           С
                                    D
                                              F.
                                                       F
# 10.04231 14.04113 12.76204 12.91898 12.57440 12.58586
with(dtf, t.test(score[treat=="A"], score[treat=="B"]))
\# t = -7.0826, df = 13.972, p-value = 5.547e-06
with(dtf, t.test(score[treat=="A"], score[treat=="D"]))
# t = -3.7533, df = 11.791, p-value = 0.002837
with(dtf, t.test(score[treat=="A"], score[treat=="C"]))
\# t = -3.5927, df = 11.912, p-value = 0.003736
with(dtf, t.test(score[treat=="A"], score[treat=="F"]))
# t = -3.9848, df = 13.549, p-value = 0.001440
with(dtf, t.test(score[treat=="A"], score[treat=="E"]))
\# t = -4.0317, df = 13.67, p-value = 0.001294
with(dtf, t.test(score[treat=="B"], score[treat=="E"]))
\# t = 2.3791, df = 13.466, p-value = 0.03276
with(dtf, t.test(score[treat=="B"], score[treat=="F"]))
# t = 2.3212, df = 13.322, p-value = 0.03673
with(dtf, t.test(score[treat=="B"], score[treat=="C"]))
# t = 1.7112, df = 11.598, p-value = 0.1136
```

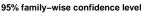
Question 3: Why is this equivalent to testing *all* pairs?

To test (snoop) beyond the planned comparisons (or if there were no planned comparisons, which is generally a bad idea), *if* you decide (before looking at the data) that only paired tests are worthwhile, then the Tukey procedure is used. Ignore any differences already tested as planned. This is based on the distribution of the maximum of paired differences under the ANOVA null hypothesis.

```
> print(TukeyHSD(e0),digits=3)
 Tukey multiple comparisons of means
    95% family-wise confidence level
Fit: aov(formula = score ~ treat, data = dtf)
$treat
       diff
               lwr
                     upr p adj
    3.9988
B-A
             1.809 6.189 0.000
    2.7197
             0.530 4.910 0.007
C-A
    2.8767
             0.686 5.067 0.004
D-A
E-A
    2.5321
             0.342 4.722 0.015
F-A 2.5435
            0.353 4.734 0.015
C-B -1.2791 -3.469 0.911 0.512
D-B -1.1221 -3.312 1.068 0.648
E-B -1.4667 -3.657 0.723 0.360
F-B -1.4553 -3.645 0.735 0.369
D-C
    0.1569 -2.033 2.347 1.000
E-C -0.1876 -2.378 2.003 1.000
F-C -0.1762 -2.366 2.014 1.000
E-D -0.3446 -2.535 1.846 0.997
F-D -0.3331 -2.523 1.857 0.997
F-E 0.0115 -2.179 2.202 1.000
```

> plot(TukeyHSD(e0, cex.axis=0.7)) # Without cex.axis, some labels are hidden.





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Question 4: How do you interpret the plot?

Question 5: Multiple choice: with the Tukey procedure the power to detect a true difference is larger/smaller/same compared to all t-tests.

Now suppose you have a fixed set of contrasts (pairs, triples, etc.) that you plan to test. The Bonferroni procedure says that if there are m tests, then test them at $\alpha' = \alpha/m$. If some of the m are planned, use the fit.contrast() results for those tests. E.g. we plan to test all pairs plus all "2 vs. 3" comparisons, with none having higher interest than any others. There are choose(6,2)=15 pairs and 6·choose(5,2)=60 ways to set up 2 vs. 3 comparisons, so m = 75. Reject only those tests with $p \leq (0.05/75) = 0.000667$. You could list all 75 tests and test them

using fit.contrast() with either 75 function calls, or with some cleverly thought about sets of orthogonal contrasts. Practically you might just test those that look likely to show large differences, e.g., for pairs we can reinterpret the t-tests above (or slightly better use contrasts with the pooled SE, as below), and for the "2 vs. 3" tests, start with two highest vs. three lowest and the three highest vs. two lowest:

```
fit.contrast(e0, "treat", rbind(A.B=c(1,-1,0,0,0,0), A.C=c(1,0,-1,0,0,0)))
#
           Estimate Std. Error t value Pr(>|t|)
# treatA.B -3.99882
                       0.73368 -5.45037
                                         0.00000
# treatA.C -2.71972
                       0.73368 -3.70698
                                        0.00061
fit.contrast(e0, "treat", rbind(A.D=c(1,0,0,-1,0,0), A.E=c(1,0,0,0,-1,0)))
            Estimate Std. Error
                                  t value
                                              Pr(>|t|)
# treatA.D -2.876669
                     0.7336774 -3.920890 0.0003200433
# treatA.E -2.532089 0.7336774 -3.451230 0.0012846660
fit.contrast(e0, "treat", rbind(BD.AEF=c(-1/2,1/3,0,-1/2,1/3,1/3),
                                AE.BCD=c(1/2,-1/3,-1/3,-1/3,1/2,0)))
#
               Estimate Std. Error
                                     t value
                                                 Pr(>|t|)
# treatBD.AEF
               1.586482
                         0.4735867
                                    3.349929 0.0017165505
# treatAE.BCD -1.932358 0.4735867 -4.080262 0.0001967149
```

Question 6: Which contrast would you try next?

For the intense data snooping approach of *all* possible contrasts, the Scheffe procedure is used. (It may also be used instead of Bonferroni for a fixed size set of contrasts if it has a more favorable cutoff.) The cutoff value for statistical significance on the F scale is, in R code: $(I-1)*qF(1-\alpha, I-1, N-I)$, where N is the total sample size, and N-I is the within-group df. We can take the square root to get on the t-test scale: > sqrt(5 * qf(0.95, 5, 42))
3.491198

So we are "safe" rejecting any contrasts with t-values less than -3.49 or greater than +3.49 if we are an intense snooper.

For example we might be (post hoc) interested in A vs all others, and we can fairly test this with:

Because $-5.16 \le -3.49$ we can say that we have sufficient evidence to claim that the population mean of A is lower than the average of the other population means.