# One-way ANOVA with post-hoc Tukey HSD Test Calculator

## .....with Scheffé, Bonferroni and Holm multiple comparison calculation also provided

#### Your input data on k=4 independent treatments:

Treatment →	А	В	С	D
Input Data →	66.0	63.0	58.0	61.0
	65.0	68.0	64.0	72.0
	67.0	71.0	66.0	65.0
	61.0	66.0	60.0	67.0
	72.0	62.0	62.0	66.0

#### Descriptive statistics of your k=4 independent treatments:

Treatment →	А	В	С	D	Pooled Total
observations N	5	5	5	5	20
sum $\sum x_i$	331.0000	330.0000	310.0000	331.0000	1,302.0000
mean $ar{x}$	66.2000	66.0000	62.0000	66.2000	65.1000
sum of squares $\sum x_i^2$	21,975.0000	21,834.0000	19,260.0000	21,975.0000	85,044.0000
sample variance $s^2$	15.7000	13.5000	10.0000	15.7000	14.9368
sample std. dev. $s$	3.9623	3.6742	3.1623	3.9623	3.8648
std. dev. of mean $SE_{\overline{x}}$	1.7720	1.6432	1.4142	1.7720	0.8642

#### One-way ANOVA of your k=4 independent treatments:

source	sum of squares SS	degrees of freedom $ u$	mean square MS	F statistic	p-value
treatment	64.2000	3	21.4000	1.5592	0.2381
error	219.6000	16	13.7250		
total	283.8000	19			

#### **Conclusion from Anova:**

The p-value corresponing to the F-statistic of one-way ANOVA is higher than 0.05, suggesting that the treatments are not significantly different for that level of significance. The Tukey HSD test, as well as other multiple comparison tests like Scheffe or Bonferroni, might not narrow down which of the pairs of treatments are significantly different. Even though your data does not suggest the presence of significantly different treatment pairs in one-way ANOVA, we proceed witht he multiple conparison tests. In some instances, a Bonferroni test of a small set of pairs might show significance, even though 1-way ANOVA suggests that there is too much noise and randomness in your data.

#### Tukey HSD Test:

The p-value corrresponing to the F-statistic of one-way ANOVA is lower than 0.05 which strongly suggests that one or more pairs of treatments are significantly different. You have k=4 treatments, for which we shall apply Tukey's HSD test to each of the 6 pairs to pinpoint which of them exhibits statistially significant difference.

We first establish the critical value of the Tukey-Kramer HSD Q statistic based on the k=4 treatments and  $\nu=16$  degrees of freedom for the error term, for significance level  $\alpha$ = 0.01 and 0.05 (p-values) in the Studentized Range distribution. We obtain these ctitical values for Q, for  $\alpha$  of 0.01 and 0.05, as  $Q_{critical}^{\alpha=0.01,k=4,\nu=16}=5.1924$  and  $Q_{critical}^{\alpha=0.05,k=4,\nu=16}=4.0464$ , respectively. These critical values may be verified at several published tables of the inverse Studentized Range distribution, such as this table at Duke University.

Next, we establish a Tukey test statistic from our sample columns to compare with the appropriate critical value of the studentized range distribution. We take the Tukey-Kramer confidence limits as documented in the  ${\color{blue} {\rm NIST}}$   ${\color{blue} {\rm Engineering~Statistics~Handbook}}$  and make simplyfying algebraic transformation. We calculate a parameter for each pair of columns being compared, which we loosely call here as the Tukey-Kramer HSD Q-statistic, or simply the Tukey HSD Q-statistic, as:

$$Q_{i,\,j}=rac{|ar{x}_i-ar{x}_j|}{s_{i,\,j}}$$

where the denominator in the above expression is:

$$s_{i,\,j} = rac{\hat{\sigma}_{\epsilon}}{\sqrt{H_{i,\,j}}} \;\;\; i,\,j=1,\,\ldots,\,k;\;i
eq j\,.$$

The quantity  $H_{i,\,j}$  is the <u>harmonic mean</u> of the number of observations in columns labeled i and j. Note that when the sample sizes in the columns are equal, then their harmonic mean is simply the common sample size. When the sample sizes of columns in a pair being compared are different, the harmonic mean lies somewhere inbetween the two sample sizes. The relvant harmonic mean is required for applying the Tukey-Kramer procedure for columns with unequal sample sizes. The quantity  $\hat{\sigma}_{\epsilon}=3.7047$  is the square root of the Mean Square Error = 13.7250 determined in the precursor one-way ANOVA procedure. Note that  $\hat{\sigma}_{\epsilon}$  is same across all pairs being compared. The only factor that varies across pairs in the computation of  $s_{i,\,j}=\frac{\hat{\sigma}_{\epsilon}}{\sqrt{H_{i,\,j}}}$  is the denominator, which is the harmonic mean of the sample sizes being compared.

The test of whether the NIST Tukey-Kramer confidence interval includes zero is equivalent to evaluating whether  $Q_{i,j} > Q_{critical}$ , the latter determined according to the desired level of significance  $\alpha$  (p-value), the number of treatments k and the degrees of freedom for error  $\nu$ , as described above.

#### post-hoc Tukey HSD Test Calculator results:

k=4 treatments

degrees of freedom for the error term  $\nu=16$  Critical values of the Studentized Range Q statistic:

$$Q_{critical}^{\alpha=0.01,k=4,\nu=16}=$$
 = 5.1924  $Q_{critical}^{\alpha=0.05,k=4,\nu=16}=$  = 4.0464

We present below color coded results (red for insignificant, green for significant) of evaluating whether  $Q_{i,j} > Q_{critical}$  for all relevant pairs of treatments. In addition, we also present the significance (p-value) of the observed Q-statistic  $Q_{i,j}$ . The algorithm used here to calculate the critical values of the studentized range distribution, as well as p-values corresponding to an observed value of  $Q_{i,j}$ , is that of <u>Gleason (1999)</u>. This is an improvement over the <u>Copenhaver-Holland (1988) algorithm</u> deployed in <u>the R statistical package</u>.

#### **Tukey HSD results**

treatments pair	Tukey HSD Q statistic	Tukey HSD p-value	Tukey HSD inferfence
A vs B	0.1207	0.8999947	insignificant
A vs C	2.5350	0.3126036	insignificant
A vs D	0.0000	0.8999947	insignificant
B vs C	2.4143	0.3526589	insignificant
B vs D	0.1207	0.8999947	insignificant
C vs D	2.5350	0.3126036	insignificant

#### Scheffé multiple comparison

We define a statistic named T as the ratio of unsigned contrast mean to contrast standard error, as explained in the <u>NIST Engineering Statistics Handbook page for Scheffe's method</u>. It can be show that for contrasts that are treatment pairs (i,j) with unit coefficients,

$$T_{i,j} = rac{Q_{i,j}}{\sqrt{2}}$$

where  $Q_{i,j}$  is the Q-statistic that was created for the Tukey HSD test. This T-statistic has interesting properties.

The same  ${\color{blue} {\rm NIST~Engineering~Statistics~Handbook~page~for~Scheffe's~method~provides~a}}$  formula which directly leads to the Scheffé p-value corresponding to an observed value of T as:

$$1-F\left(rac{T^2}{k-1},k-1,
u
ight)$$

where F() is the cumulative F distribution with its two degrees of freedom parameters k-1 and  $\nu$ . Note that k is the number of treatments and  $\nu$  is the degrees of freedom of error that were established earlier.

The Scheffé p-value of the observed T-statistic  $T_{i,j}$  is shown below for all relevant pairs of treatments, along with color coded Scheffé inference (red for insignificant, green for significant) based on the p-value.

#### Scheffé results

treatments pair	Scheffé $T$ -statistic	Scheffé p-value	Scheffé inferfence
A vs B	0.0854	0.9998274	insignificant
A vs C	1.7925	0.3891847	insignificant
A vs D	0.0000	1.0000000	insignificant
B vs C	1.7072	0.4305041	insignificant
B vs D	0.0854	0.9998274	insignificant
C vs D	1.7925	0.3891847	insignificant

#### Bonferroni and Holm multiple comparison

The same statistic T for the Scheffé method, along with the number of contrasts (pairs) q being simultaneously compared, leads to the Bonferroni formula. The <u>NIST Engineering Statistics Handbook page for Bonferroni method</u> provides a formula which directly leads to the Bonferroni p-value corresponding to an observed value of T in the context of simultaneous comparison of q contrasts as:

context of simultaneous comparison of q contrasts as: Bonferroni p-value:  $P_{i,j}^{Bonferroni}=P_{i,j}^{unadjusted}q$  where

$$P_{i,j}^{unadjusted} = \left[1 - t\left(rac{T^2}{k-1},
u
ight)
ight]2$$

and where  $t\left(\right)$  is the cumulative Student's t distribution with its degree of freedom parameter  $\nu$ . Note that  $\nu$  is the degrees of freedom of error that were established earlier. Also note that the p-value of Bonferroni simultaneous comparison is directly proportional to q, the number of contrasts (pairs) being simultaneously compared.

The Holm procedure described in <u>Aickin and Gensler (1996) review paper</u> requires sorting the  $P_{i,j}^{unadjusted}$  as above in *ascending* order and determining the sort rank  $R_{i,j}$  of each unique pair (i,j). These sort ranks run from 1 through q. The Holm p-value for comparing a given pair (i,j) in the context of multiple comparison of q such pairs simultaneously is:

Holm p-value:  $P_{i,j}^{Holm} = P_{i,j}^{unadjusted} \left(q - R_{i,j} + 1
ight)$ 

In this first combined Bonferroni and Holm table below, we consider all possible contrasts (pairs) for simultaneous comparion, thus q=6. The Bonferoni and Holm p-values of the observed T-statistic  $T_{i,j}$  for all relevant q=6 pairs of treatments is shown below, along with color coded Bonferroni and Holm inferences (red for insignificant, green for significant) based on the p-value.

#### Bonferroni and Holm results: all pairs simultaineously compared

treatments pair	Bonferroni and Holm $T ext{-statistic}$	Bonferroni p-value	Bonferroni inferfence	Holm p-value	Holm inferfence
A vs B	0.0854	5.5982150	insignificant	2.7991075	insignificant
A vs C	1.7925	0.5518603	insignificant	0.5518603	insignificant
A vs D	0.0000	6.0000000	insignificant	1.0000000	insignificant
B vs C	1.7072	0.6427252	insignificant	0.4284835	insignificant
B vs D	0.0854	5.5982150	insignificant	1.8660717	insignificant
C vs D	1.7925	0.5518603	insignificant	0.4598836	insignificant

In this second Bonferroni and Holm table below, we consider a subset of contrasts (pairs) for simultaneous comparion, of only pairs relative to treatment A. Such a situation may be relevant when treatment A is the control, and the experimenter is interested only in differences of treatments relative to control, thus q=3. The Bonferoni and Holm p-values of the observed T-statistic  $T_{i,j}$  for q=3 relevant pairs of treatments, along with color coded Bonferroni inference (red for insignificant, green for significant) based on the p-value.

#### Bonferroni and Holm results: only pairs relative to A simultaineously compared

treatments pair	Bonferroni and Holm $T ext{-statistic}$	Bonferroni p-value	Bonferroni inferfence	Holm p-value	Holm inferfence
A vs B	0.0854	2.7991075	insignificant	1.8660717	insignificant
A vs C	1.7925	0.2759302	insignificant	0.2759302	insignificant
A vs D	0.0000	3.0000000	insignificant	1.0000000	insignificant

#### How to repeat & verify post-hoc Tukey HSD calculation by hand in Microsoft Excel

Microsoft Excel lacks built-in functions relating to the studentized range distribution, so even though it calculates the Mean Square Error in one-way ANOVA, whose square root is  $\hat{\sigma}_{\epsilon}$ , and is aware of all the sample sizes and degrees of freedom, it is unable to conduct the next step of post-hoc Tukey HSD comparison of treatments. To manually conduct post-hoc Tukey HSD test calculation, you would take the mean squared error from the Excel's one-way ANOVA output, then take its square root to determine  $\hat{\sigma}_{\epsilon}$ . You would then divide this  $\hat{\sigma}_{\epsilon}$  by the square root of  $H_{i,j}$ , the harmonic mean of the relevant sample columns being compared, resulting in  $s_{i,j}$ , for each pair of columns (i,j). Excel has a built-in function HARMEAN(n1, n2,...) that calculates the harmonic mean. With these on hand, you would determine  $Q_{i,j} = \frac{|\bar{x}_i - \bar{x}_j|}{s_{i,j}}$ . Microsoft Excel provides the relevant sample column averages (means) to calculate the numerator. You would finally compare whether  $Q_{i,j} > Q_{critical}$ . For this comparison, you would obtain the critical values for the appropriate number of degrees of freedom of error (shown in Microsoft Excel) and the number of treatments (columns) from tables of the (inverse) studentized range distribution widely available on the web.

### How to repeat & verify post-hoc Scheffé, Bonferroni and Holm calculations by hand in Microsoft Excel

For Scheffé, Bonferroni and Holm steps, calculate the T-statistic  $T_{i,\,j}$  for all pairs by taking  $Q_{i,\,j}$  from the earlier Tukey HSD step and dividing it by  $\sqrt{2}$ . Calculate the Scheffé p-value of the observed T-statistic by using Excel's built-in formula for the F distribution which has the form F.DIST (x,deg\_freedom1,deg\_freedom2,cumulative). Set

its first argument x as  $\frac{T^2}{k-1}$ . The second and third arguments are k-1 and  $\nu$ . The fourth argument is set to 1. The Scheffé p-value is calculated in Excel by the formula as 1-F.DIST(x,  $k-1, \nu, 1$ ).

For the Bonferroni and Holm comparison, take the same T-statistic  $T_{i,j}$  that was determined for the Scheffé step above. Determine q, the number of pairs that are being simultaneously compared. Calculate the Bonferroni p-value of the observed T-statistic by using Excel's <u>built-in formula for the t-distribution T.DIST(x,deg\_freedom, cumulative)</u>. The unadusted p-value  $P^{unadjusted}$  is calculated in Excel by the formula as  $(1-T.DIST(T,\nu,TRUE))^2$ . The Bonferroni p-value is calculated for each pair (i,j) as  $P^{Bonferroni} = P^{unadjusted}q$ . The q-element array of the  $P^{unadjusted}_{i,j}$  is sorted in ascending order to determine the sort rank  $R_{i,j}$  for each given pair (i,j). These sort ranks run from 1 through q. The Holm p-value is calculated for each pair (i,j) as  $P^{Holm}_{i,j} = P^{unadjusted}_{i,j} (q - R_{i,j} + 1)$ .

### R code and Tutorial for conducting Tukey HSD, Scheffé, Bonferroni and Holm methods

A tutorial for the solving the demo example using the free open-source academic-research-grade <u>R statistical</u> <u>package</u> together with complete <u>R code and output is provided here</u>. The output of the demo example in this web calculator for all three methods of multiple comparison are fully reproduced in R, thus further establishing the validity of the formula and methodology discussed earlier.

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