Chapter 95
The TTEST Procedure

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Overview: TTEST Procedure

The TTEST procedure performs $t$ tests and computes confidence limits for one sample, paired observations, two independent samples, and the AB/BA crossover design. Two-sided, TOST (two one-sided test) equivalence, and upper and lower one-sided hypotheses are supported for means, mean differences, and mean ratios for either normal or lognormal data.

Table 95.1 summarizes the designs, analysis criteria, hypotheses, and distributional assumptions supported in the TTEST procedure, along with the syntax used to specify them.

<table>
<thead>
<tr>
<th>Feature</th>
<th>Syntax</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Design</strong></td>
<td></td>
</tr>
<tr>
<td>One-sample</td>
<td>VAR statement</td>
</tr>
<tr>
<td>Paired</td>
<td>PAIRED statement</td>
</tr>
<tr>
<td>Two-independent-sample</td>
<td>CLASS statement, VAR statement</td>
</tr>
<tr>
<td>AB/BA crossover</td>
<td>VAR / CROSSOVER=</td>
</tr>
<tr>
<td><strong>Analysis Criterion</strong></td>
<td></td>
</tr>
<tr>
<td>Mean difference</td>
<td>PROC TTEST TEST=DFF</td>
</tr>
<tr>
<td>Mean ratio</td>
<td>PROC TTEST TEST=RATIO</td>
</tr>
<tr>
<td><strong>Hypothesis</strong></td>
<td></td>
</tr>
<tr>
<td>Two-sided</td>
<td>PROC TTEST SIDES=2</td>
</tr>
<tr>
<td>Equivalence</td>
<td>PROC TTEST TOST ( &lt;lower , &gt; upper )</td>
</tr>
<tr>
<td>Lower one-sided</td>
<td>PROC TTEST SIDES=L</td>
</tr>
<tr>
<td>Upper one-sided</td>
<td>PROC TTEST SIDES=U</td>
</tr>
<tr>
<td><strong>Distribution</strong></td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>PROC TTEST DIST=NORMAL</td>
</tr>
<tr>
<td>Lognormal</td>
<td>PROC TTEST DIST=LOGNORMAL</td>
</tr>
</tbody>
</table>

FREQ and WEIGHT statements are available. Data can be input in the form of observations or, in certain cases, summary statistics. Output includes summary statistics; confidence limits for means, standard deviations, and coefficients of variation; hypothesis tests; and a variety of graphical displays, including histograms, densities, box plots, confidence intervals, Q-Q plots, profiles, and agreement plots.
Getting Started: TTEST Procedure

One-Sample $t$ Test

A one-sample $t$ test can be used to compare a sample mean to a given value. This example, taken from Huntsberger and Billingsley (1989, p. 290), tests whether the mean length of a certain type of court case is more than 80 days by using 20 randomly chosen cases. The data are read by the following DATA step:

```plaintext
data time;
  input time @@;
datalines;
43 90 84 87 116 95 86 99 93 92
121 71 66 98 79 102 60 112 105 98
;
run;
```

The only variable in the data set, `time`, is assumed to be normally distributed. The trailing at signs (@@) indicate that there is more than one observation on a line. The following statements invoke PROC TTEST for a one-sample $t$ test:

```plaintext
ods graphics on;
proc ttest h0=80 plots(showh0) sides=u alpha=0.1;
  var time;
run;
ods graphics off;
```

The `VAR` statement indicates that the `time` variable is being studied, while the `H0=` option specifies that the mean of the `time` variable should be compared to the null value 80 rather than the default of 0. The `PLOTS(SHOWH0)` option requests that this null value be displayed on all relevant graphs. The `SIDES=U` option reflects the focus of the research question, namely whether the mean court case length is greater than 80 days, rather than different than 80 days (in which case you would use the default `SIDES=2` option). The `ALPHA=0.1` option requests 90% confidence intervals rather than the default 95% confidence intervals. The output is displayed in Figure 95.1.
Figure 95.1 One-Sample $t$ Test Results

<table>
<thead>
<tr>
<th>The TTEST Procedure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Variable: time</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>N</th>
<th>Mean</th>
<th>Std Dev</th>
<th>Std Err</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>89.8500</td>
<td>19.1456</td>
<td>4.2811</td>
<td>43.0000</td>
<td>121.0000</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Mean 90% CL Mean</th>
<th>Std Dev 90% CL Std Dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>89.8500 84.1659 Infty</td>
<td>19.1456 15.2002 26.2374</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>DF</th>
<th>t Value</th>
<th>Pr &gt; t</th>
</tr>
</thead>
<tbody>
<tr>
<td>19</td>
<td>2.30</td>
<td>0.0164</td>
</tr>
</tbody>
</table>

Summary statistics appear at the top of the output. The sample size (N), mean, standard deviation, and standard error are displayed with the minimum and maximum values of the time variable. The 90% confidence limits for the mean and standard deviation are shown next. Due to the SIDES=U option, the interval for the mean is an upper one-sided interval with a finite lower bound (84.1659 days). The limits for the standard deviation are the equal-tailed variety, per the default CI=EQUAL option in the PROC TTEST statement. At the bottom of the output are the degrees of freedom, $t$ statistic value, and $p$-value for the $t$ test. At the 10% $\alpha$ level, this test indicates that the mean length of the court cases is significantly greater than from 80 days ($t = 2.30, p = 0.0164$).

The summary panel in Figure 95.2 shows a histogram with overlaid normal and kernel densities, a box plot, the 90% confidence interval for the mean, and the null value of 80 days.
The confidence interval excludes the null value, consistent with the rejection of the null hypothesis at $\alpha = 0.1$.

The Q-Q plot in Figure 95.3 assesses the normality assumption.
The curvilinear shape of the Q-Q plot suggests a possible slight deviation from normality. You could use the UNIVARIATE procedure with the NORMAL option to numerically check the normality assumptions.

**Comparing Group Means**

If you want to compare values obtained from two different groups, and if the groups are independent of each other and the data are normally or lognormally distributed in each group, then a group $t$ test can be used. Examples of such group comparisons include the following:

- test scores for two third-grade classes, where one of the classes receives tutoring
- fuel efficiency readings of two automobile nameplates, where each nameplate uses the same fuel
- sunburn scores for two sunblock lotions, each applied to a different group of people
- political attitude scores of males and females
In the following example, the golf scores for males and females in a physical education class are compared. The sample sizes from each population are equal, but this is not required for further analysis. The scores are thought to be approximately normally distributed within gender. The data are read by the following statements:

```plaintext
data scores;
  input Gender $ Score @@;
datalines;
f 75 f 76 f 80 f 77 f 80 f 77 f 73
m 82 m 80 m 85 m 85 m 78 m 87 m 82;
run;
```

The dollar sign ($) following `Gender` in the INPUT statement indicates that `Gender` is a character variable. The trailing at signs (@@) enable the procedure to read more than one observation per line.

You can use a group *t* test to determine whether the mean golf score for the men in the class differs significantly from the mean score for the women. If you also suspect that the distributions of the golf scores of males and females have unequal variances, then you might want to specify the COCHRAN option in order to use the Cochran approximation (in addition to the Satterthwaite approximation, which is included by default). The following statements invoke PROC TTEST for the case of unequal variances, along with both types of confidence limits for the pooled standard deviation.

```plaintext
ods graphics on;
proc ttest cochran ci=equal umpu;
  class Gender;
  var Score;
run;
ods graphics off;
```

The CLASS statement contains the variable that distinguishes the groups being compared, and the VAR statement specifies the response variable to be used in calculations. The COCHRAN option produces *p*-values for the unequal variance situation by using the Cochran and Cox (1950) approximation. Equal-tailed and uniformly most powerful unbiased (UMPU) confidence intervals for *σ* are requested by the CI= option. Output from these statements is displayed in Figure 95.4 through Figure 95.7.

**Figure 95.4** Simple Statistics

<table>
<thead>
<tr>
<th>Gender</th>
<th>N</th>
<th>Mean</th>
<th>Std Dev</th>
<th>Std Err</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>f</td>
<td>7</td>
<td>76.8571</td>
<td>2.5448</td>
<td>0.9619</td>
<td>73.0000</td>
<td>80.0000</td>
</tr>
<tr>
<td>m</td>
<td>7</td>
<td>82.7143</td>
<td>3.1472</td>
<td>1.1895</td>
<td>78.0000</td>
<td>87.0000</td>
</tr>
<tr>
<td>Diff (1-2)</td>
<td></td>
<td>-5.8571</td>
<td>2.8619</td>
<td>1.5298</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Simple statistics for the two populations being compared, as well as for the difference of the means between the populations, are displayed in Figure 95.4. The Gender column indicates the population corresponding to the statistics in that row. The sample size (N), mean, standard deviation, standard error, and minimum and maximum values are displayed.

Confidence limits for means and standard deviations are shown in Figure 95.5.

**Figure 95.5 Simple Statistics**

<table>
<thead>
<tr>
<th>Gender</th>
<th>Method</th>
<th>Mean</th>
<th>95% CL Mean</th>
<th>Std Dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>f</td>
<td></td>
<td>76.8571</td>
<td>74.5036</td>
<td>79.2107</td>
</tr>
<tr>
<td>m</td>
<td></td>
<td>82.7143</td>
<td>79.8036</td>
<td>85.6249</td>
</tr>
<tr>
<td>Diff (1-2) Pooled</td>
<td>-5.8571</td>
<td>-9.1902</td>
<td>-2.5241</td>
<td>2.8619</td>
</tr>
<tr>
<td>Diff (1-2) Satterthwaite</td>
<td>-5.8571</td>
<td>-9.2064</td>
<td>-2.5078</td>
<td></td>
</tr>
</tbody>
</table>

For the mean differences, both pooled (assuming equal variances for males and females) and Satterthwaite (assuming unequal variances) 95% intervals are shown. The confidence limits for the standard deviations are of the equal-tailed variety.

The test statistics, associated degrees of freedom, and \( p \)-values are displayed in Figure 95.6.

**Figure 95.6 \( t \) Tests**

| Method   | Variances | DF  | \( t \) Value | \( Pr > |t| \)  |
|----------|-----------|-----|---------------|--------------|
| Pooled   | Equal     | 12  | -3.83         | 0.0024       |
| Satterthwaite | Unequal | 11.496 | -3.83         | 0.0026       |
| Cochran  | Unequal   | 6   | -3.83         | 0.0087       |

The Method column denotes which \( t \) test is being used for that row, and the Variances column indicates what assumption about variances is being made. The pooled test assumes that the two populations have equal variances and uses degrees of freedom \( n_1 + n_2 - 2 \), where \( n_1 \) and \( n_2 \) are the sample sizes for the two populations. The remaining two tests do not assume that the populations have equal variances. The Satterthwaite test uses the Satterthwaite approximation for degrees of freedom, while the Cochran test uses the Cochran and Cox approximation for the \( p \)-value. All three tests result in highly significant \( p \)-values, supporting the conclusion of a significant difference between males’ and females’ golf scores.

The “Equality of Variances” test in Figure 95.7 reveals insufficient evidence of unequal variances (the Folded F statistic \( F' = 1.53 \), with \( p = 0.6189 \)).
The summary panel in Figure 95.8 shows comparative histograms, normal and kernel densities, and box plots, comparing the distribution of golf scores between genders.

The Q-Q plots in Output 95.9 assess the normality assumption for each gender.
The plots for both males and females show no obvious deviations from normality. You can check the assumption of normality more rigorously by using PROC UNIVARIATE with the NORMAL option; if the assumption of normality is not reasonable, you should analyze the data with the nonparametric Wilcoxon rank sum test by using PROC NPAR1WAY.

**Syntax: TTEST Procedure**

The following statements are available in PROC TTEST:

```
PROC TTEST < options > ;
   CLASS variable ;
   PAIRED variables ;
   BY variables ;
   VAR variables < / options > ;
   FREQ variable ;
   WEIGHT variable ;
```

No statement can be used more than once. There is no restriction on the order of the statements after the PROC TTEST statement.
PROC TTEST Statement

PROC TTEST <options> ;

The PROC TTEST statement invokes the procedure. Table 95.2 summarizes the options in the PROC TTEST statement by function. The options are then described fully in alphabetical order.

Table 95.2 PROC TTEST Statement Options

<table>
<thead>
<tr>
<th>Option</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Basic Options</strong></td>
<td></td>
</tr>
<tr>
<td>DATA=</td>
<td>Specifies input data set</td>
</tr>
<tr>
<td>ORDER=</td>
<td>Determines sort order of CLASS variable or CROSSOVER= treatment variables</td>
</tr>
<tr>
<td><strong>Analysis Options</strong></td>
<td></td>
</tr>
<tr>
<td>ALPHA=</td>
<td>Specifies 1− confidence level</td>
</tr>
<tr>
<td>DIST=</td>
<td>Specifies distributional assumption (normal or lognormal)</td>
</tr>
<tr>
<td>H0=</td>
<td>Specifies null value</td>
</tr>
<tr>
<td>SIDES=</td>
<td>Specifies number of sides and direction</td>
</tr>
<tr>
<td>TEST=</td>
<td>Specifies test criterion (difference or ratio)</td>
</tr>
<tr>
<td>TOST</td>
<td>Requests equivalence test and specifies bounds</td>
</tr>
<tr>
<td><strong>Displayed Output</strong></td>
<td></td>
</tr>
<tr>
<td>CI=</td>
<td>Requests confidence interval for standard deviation or CV</td>
</tr>
<tr>
<td>COCHRAN</td>
<td>Requests Cochran t test</td>
</tr>
<tr>
<td>PLOTS</td>
<td>Produces ODS statistical graphics</td>
</tr>
<tr>
<td><strong>Output Ordering</strong></td>
<td></td>
</tr>
<tr>
<td>BYVAR</td>
<td>Groups results by PAIRED or VAR variables</td>
</tr>
<tr>
<td>NOBYVAR</td>
<td>Groups results by tables</td>
</tr>
</tbody>
</table>

The following options can appear in the PROC TTEST statement.

**ALPHA=p**
specifies that confidence intervals (except test-based mean confidence intervals when the TOST option is used) are to be 100(1 − p)% confidence intervals, where 0 < p < 1. When the TOST option is used, the test-based mean confidence intervals are 100(1 − 2p)% confidence intervals. By default, PROC TTEST uses ALPHA=0.05. If p is 0 or less, or 1 or more, an error message is printed.

**BYVAR**
groups the results by the PAIRED or VAR variables. The BYVAR option is enabled by default. Note that this represents a change from previous releases for how the results are grouped with respect to variables and tables. Prior to SAS 9.2, multiple variables were included in each table, similar to the new NOBYVAR option.
**CI=EQUAL | UMPU | NONE**

**CL=EQUAL | UMPU | NONE**

specifies whether a confidence interval is displayed for \( \sigma \) and, if so, what kind. The **CI=EQUAL** option specifies an equal-tailed confidence interval, and it is the default. The **CI=UMPU** option specifies an interval based on the uniformly most powerful unbiased test of \( H_0: \sigma = \sigma_0 \). The **CI=NONE** option requests that no confidence interval be displayed for \( \sigma \). The values **EQUAL** and **UMPU** together request that both types of confidence intervals be displayed. If the value **NONE** is specified with one or both of the values **EQUAL** and **UMPU**, **NONE** takes precedence. For more information, see the section “Two-Independent-Sample Design” on page 8021.

**COCHRAN**

requests the Cochran and Cox (1950) approximation of the probability level for the unequal variances situation. For more information, see the section “Two-Independent-Sample Design” on page 8021.

**DATA=SAS-data-set**

names the SAS data set for the procedure to use. By default, PROC TTEST uses the most recently created SAS data set. The input data set can contain summary statistics of the observations instead of the observations themselves. The number, mean, and standard deviation of the observations are required for each **BY** group (one sample and paired differences) or for each class within each **BY** group (two samples). For more information about the **DATA=** option, see the section “Input Data Set of Statistics” on page 8015.

**DIST=LOGNORMAL | NORMAL**

specifies the underlying distribution assumed for the data. The default is **NORMAL**, unless **TEST=RATIO** is specified, in which case the default is **LOGNORMAL**.

**H0=m**

requests tests against a null value of \( m \), unless the **TOST** option is used, in which case \( m \) is merely used to derive the lower and upper equivalence bounds. For the crossover design, the value \( m \) applies for both treatment and period tests. By default, PROC TTEST uses \( H0=0 \) when **TEST=DIFF** (or **DIST=NORMAL** for a one-sample design) and \( H0=1 \) when **TEST=RATIO** (or **DIST=LOGNORMAL** for a one-sample design).

**NOBYVAR**

includes all **PAIRED** or **VAR** variables together in each output table. If the **NOBYVAR** option is not specified, then the **BYVAR** option is enabled, grouping the results by the **PAIRED** and **VAR** variables.

**ORDER=DATA | FORMATTED | FREQ | INTERNAL | MIXED**

specifies the order in which to sort the levels of the classification variables (which are specified in the **CLASS** statement) and treatment variables (which are specified in the **CROSSOVER=** option in the **VAR** statement). The default is **ORDER=MIXED**, which corresponds to the ordering in releases previous to SAS 9.2.

This option applies to the levels for all classification or treatment variables, except when you use the **ORDER=FORMATTED** option with numeric classification or treatment variables that have no explicit format. With this option, the levels of such variables are ordered by their internal value.

The **ORDER=** option can take the following values:
The PROC TTEST statement provides options for specifying how levels of a variable are sorted in the input data set. The `ORDER=` option determines the order in which levels are sorted. The possible values for `ORDER=` are:

<table>
<thead>
<tr>
<th>Value of ORDER=</th>
<th>Levels Sorted By</th>
</tr>
</thead>
<tbody>
<tr>
<td>DATA</td>
<td>Order of appearance in the input data set.</td>
</tr>
<tr>
<td>FORMATTED</td>
<td>External formatted value, except for numeric variables with no explicit format, which are sorted by their unformatted (internal) value.</td>
</tr>
<tr>
<td>FREQ</td>
<td>Descending frequency count; levels with the most observations come first in the order. In the event of a tie, ORDER=MIXED is used.</td>
</tr>
<tr>
<td>INTERNAL</td>
<td>Unformatted value.</td>
</tr>
<tr>
<td>MIXED</td>
<td>Same as ORDER=FORMATTED if the unformatted variable is character-valued; same as ORDER=INTERNAL otherwise (the unformatted variable is numeric-valued).</td>
</tr>
</tbody>
</table>

For FORMATTED and INTERNAL, the sort order is machine-dependent.

For more information about sorting order, see the chapter on the SORT procedure in the Base SAS Procedures Guide and the discussion of BY-group processing in SAS Language Reference: Concepts.

The `PLOTS` option controls the plots produced through ODS Graphics. When you specify only one plot request, you can omit the parentheses around the plot request. Here are some examples:

- `plots=none`
- `plots=(histogram boxplot interval qq profiles agreement)`
- `plots(unpack)=summary`
- `plots(showh0)=interval(type=pergroup)`
- `plots=(summary(unpack) interval(type=period))`

ODS Graphics must be enabled before requesting plots. For example:

```sas
ods graphics on;

proc ttest plots=all;
   var oxygen;
run;

ods graphics off;
```

For more information about enabling and disabling ODS Graphics, see the section “Enabling and Disabling ODS Graphics” on page 609 in Chapter 21, “Statistical Graphics Using ODS.”

If ODS Graphics is enabled but you do not specify the PLOTS option, then PROC TTEST produces a default set of plots. (NOTE: The graphical results are unavailable if your input data set contains summary statistics rather than observation values.)
For a one-sample design, the default plots are the following:

- summary plot (histogram with overlaid normal and kernel densities, box plot, and confidence interval band)
- Q-Q plot

For a two-independent-sample design, the default plots are the following:

- summary plot (comparative histograms with overlaid densities and box plots)
- Q-Q plot

For a paired design, the default plots are the following:

- summary plot (histogram, densities, box plot, and confidence interval) of the difference or ratio
- Q-Q plot of the difference or ratio
- profiles plot
- agreement plot

For a crossover design, the default plots are the following:

- comparative histograms with overlaid densities by treatment and period
- comparative box plots by treatment and period
- Q-Q plots by treatment and period
- profiles over treatment plot
- agreement of treatments plot

For more detailed descriptions of plots, see the section “Interpreting Graphs” on page 8032.

The global plot options include the following:

ONLY
suppresses the default plots. Only plots specifically requested are displayed.

SHOWH0
SHOWNULL
shows the null value (as specified by the H0= option in the PROC TTEST statement) in all relevant plots. For one-sample and paired designs, the null value can appear in SUMMARY, BOX, and INTERVAL. For two-independent-sample and crossover designs, the null value can appear only in INTERVAL.

UNPACKPANEL
UNPACK
suppresses paneling. By default, multiple plots can appear in some output panels. Specify UNPACKPANEL to get each plot in a separate panel. You can specify PLOTS(UNPACKPANEL) to unpack the default plots. You can also specify UNPACK as a suboption with SUMMARY.
The plot requests include the following:

**ALL**
produces all appropriate plots. You can specify other options with **ALL**; for example, to request all plots and specify that intervals should be for the period difference in a crossover design, specify `PLOTS=(ALL INTERVAL(TYPE=PERIOD))`.

**AGREEMENT**
requests an agreement plot. This plot is produced by default for paired and crossover designs, the only designs for which the **AGREEMENT** option is valid.

For paired designs, the second response in each pair is plotted against the first response. See the section “Agreement Plots for Paired Designs” on page 8032 for further details.

For crossover designs, the **AGREEMENT** plot request has the following options:

**TYPE=PERIOD**
plots the response in the second period against the response in the first period. See the section “Period Agreement Plots for Crossover Designs” on page 8033 for further details.

**TYPE=TREATMENT**
plots the response associated with the second treatment against the response associated with the first treatment. This is the default **TYPE=** option for crossover designs. See the section “Treatment Agreement Plots for Crossover Designs” on page 8033 for further details.

**BOX**
requests a box plot or comparative box plots. This plot is produced by default for crossover designs. For other designs, a box plot appears as part of the **SUMMARY** plot by default.

For one-sample and paired designs, a confidence interval for the mean is shown as a band in the background, along with the equivalence bounds if the **TOST** option is used in the **PROC TTEST** statement.

For a two-independent-sample design, comparative box plots (one for each class) are shown. For a crossover design, comparative box plots for all four combinations of the two treatments and two periods are shown.

See the section “Box Plots” on page 8033 for further details.

**HISTOGRAM**
requests a histogram or comparative histograms with overlaid normal and kernel densities. This plot is produced by default for crossover designs. For other designs, it appears as part of the **SUMMARY** plot by default.
For one-sample and paired designs, the histogram and densities are based on the test criterion (which is the mean difference or ratio for a paired design). For a two-independent-sample design, comparative histograms (one for each class) are shown. For a crossover design, histograms for all four combinations of the two treatments and two periods are shown.

See the section “Histograms” on page 8034 for further details.

**INTERVAL**
**INTERVALPLOT**
requests plots of confidence interval for means.

For a two-independent-sample design, the INTERVAL plot request has the following options:

**TYPE=PERGROUP**
shows two separate two-sided confidence intervals, one for each class. This option cannot be used along with the SHOWH0 global plot option.

**TYPE=TEST**
shows pooled and Satterthwaite confidence intervals. This is the default TYPE= option for two-independent-sample designs.

For a crossover design, the INTERVAL plot request has the following options:

**TYPE=PERGROUP**
shows four separate two-sided intervals, one for each treatment-by-period combination. This option cannot be used along with the SHOWH0 global plot option.

**TYPE=PERIOD**
shows pooled and Satterthwaite confidence intervals for the period difference or ratio. This option is invalid if the IGNOREPERIOD option is used in the VAR statement.

**TYPE=TREATMENT**
shows pooled and Satterthwaite confidence intervals for the treatment difference or ratio. This is the default TYPE= option for crossover designs.

See the section “Confidence Intervals” on page 8034 for further details.

**NONE**
suppresses all plots.

**PROFILES**
**PROFILESPLOT**
requests a profiles plot. This plot is produced by default for paired and crossover designs, the only designs for which the PROFILES option is valid.

For paired designs, a line is drawn for each observation from left to right connecting the first response to the second response. See the section “Profiles for Paired Designs” on page 8034 for further details.

For crossover designs, the PROFILES plot request has the following options:
PROC TTEST Statement

TYPE=PERIOD
shows response profiles over period, connecting the first period on the left to the second period on the right for each subject. See the section “Profiles over Period for Crossover Designs” on page 8034 for further details.

TYPE=TREATMENT
shows response profiles over treatment values, connecting the first treatment on the left to the second treatment on the right for each observation. This is the default TYPE= option for crossover designs. See the section “Profiles over Treatment for Crossover Designs” on page 8035 for further details.

QQ
QQPLOT
requests a normal quantile-quantile (Q-Q) plot. This plot is produced by default for all designs. For two-sample designs, separate plots are shown for each class in a single panel. For crossover design, separate plots are shown for each treatment-by-period combination in a single panel.

See the section “Q-Q Plots” on page 8035 for further details.

SUMMARY
SUMMARYPLOT
requests HISTOGRAM and BOX plots together in a single panel, sharing common X axes. This plot is produced by default for one-sample, paired, and two-independent-sample designs, the only designs for which the SUMMARY option is valid. See the documentation for the BOX and HISTOGRAM plot requests for details. The SUMMARY plot request has the following option:

UNPACK
plots histograms with overlaid densities in one panel and box plots (along with confidence interval bands, if one-sample or paired design) in another. Note that specifying PLOTS(ONLY)=SUMMARY(UNPACK) is exactly the same as specifying PLOTS(ONLY)=(BOX HISTOGRAM).

SIDES=2 | L | U
SIZED=2 | L | U
SIDE=2 | L | U
specifies the number of sides (or tails) and direction of the statistical tests and test-based confidence intervals. The values are interpreted as follows:

SIDES=2 (the default) specifies two-sided tests and confidence intervals for means.
SIDES=L specifies lower one-sided tests, in which the alternative hypothesis indicates a mean less than the null value, and lower one-sided confidence intervals between minus infinity and the upper confidence limit.
SIDES=U specifies upper one-sided tests, in which the alternative hypothesis indicates a mean greater than the null value, and upper one-sided confidence intervals between the lower confidence limit and infinity.
Chapter 95: The TTEST Procedure

TEST=DIFF | RATIO

specifies the test criterion. Use TEST=DIFF to test the difference of means and TEST=RATIO to test the ratio of means. The default is DIFF, unless DIST=LOGNORMAL is specified, in which case the default is RATIO. This option is ignored for one-sample designs.

TOST ( <lower ,> upper )
requests Schuirman’s TOST equivalence test. The upper equivalence bound must be specified. If TEST=DIFF, then the default value for the lower equivalence bound is $2m - upper$, where $m$ is the value of the H0= option. If TEST=RATIO, then the default value for lower is $m / upper$.

BY Statement

BY variables ;

You can specify a BY statement with PROC TTEST to obtain separate analyses on observations in groups that are defined by the BY variables. When a BY statement appears, the procedure expects the input data set to be sorted in order of the BY variables. If you specify more than one BY statement, only the last one specified is used.

If your input data set is not sorted in ascending order, use one of the following alternatives:

- Sort the data by using the SORT procedure with a similar BY statement.
- Specify the NOTSORTED or DESCENDING option in the BY statement for the TTEST procedure. The NOTSORTED option does not mean that the data are unsorted but rather that the data are arranged in groups (according to values of the BY variables) and that these groups are not necessarily in alphabetical or increasing numeric order.
- Create an index on the BY variables by using the DATASETS procedure (in Base SAS software).

For more information about BY-group processing, see the discussion in SAS Language Reference: Concepts. For more information about the DATASETS procedure, see the discussion in the Base SAS Procedures Guide.

CLASS Statement

CLASS variable ;

A CLASS statement giving the name of the classification (or grouping) variable must accompany the PROC TTEST statement in the two-independent-sample case. It should be omitted for the one-sample, paired, and AB/BA crossover designs. If it is used without the VAR statement, all numeric variables in the input data set (except those that appear in the CLASS, BY, FREQ, or WEIGHT statement) are included in the analysis.
The classification variable must have two, and only two, levels. PROC TTEST divides the observations into the two groups for the t test by using the levels of this variable. You can use either a numeric or a character variable in the CLASS statement.

Class levels are determined from the formatted values of the CLASS variable. Thus, you can use formats to define group levels. See the discussions of the FORMAT procedure, the FORMAT statement, formats, and informats in SAS Language Reference: Dictionary.

### FREQ Statement

**FREQ variable ;**

The variable in the FREQ statement identifies a variable that contains the frequency of occurrence of each observation. PROC TTEST treats each observation as if it appears n times, where n is the value of the FREQ variable for the observation. If the value is not an integer, only the integer portion is used. If the frequency value is less than 1 or is missing, the observation is not used in the analysis. When the FREQ statement is not specified, each observation is assigned a frequency of 1. The FREQ statement cannot be used if the DATA= data set contains statistics instead of the original observations.

### PAIRED Statement

**PAIRED PairLists ;**

The PairLists in the PAIRED statement identifies the variables to be compared in paired comparisons. You can use one or more PairLists. Variables or lists of variables are separated by an asterisk (*) or a colon (:). The asterisk requests comparisons between each variable on the left with each variable on the right. The colon requests comparisons between the first variable on the left and the first on the right, the second on the left and the second on the right, and so forth. The number of variables on the left must equal the number on the right when the colon is used. The differences are calculated by taking the variable on the left minus the variable on the right for both the asterisk and colon. A pair formed by a variable with itself is ignored. Use the PAIRED statement only for paired comparisons. The CLASS and VAR statements cannot be used with the PAIRED statement.

Examples of the use of the asterisk and the colon are shown in Table 95.3.
Table 95.3 PAIRED Statement in the TTEST Procedure

<table>
<thead>
<tr>
<th>These PAIRED statements...</th>
<th>yield these comparisons</th>
</tr>
</thead>
<tbody>
<tr>
<td>PAIRED A*B;</td>
<td>A-B</td>
</tr>
<tr>
<td>PAIRED A<em>B C</em>D;</td>
<td>A-B and C-D</td>
</tr>
<tr>
<td>PAIRED (A B) * (C D);</td>
<td>A-C, A-D, B-C, and B-D</td>
</tr>
<tr>
<td>PAIRED (A B) * (C B);</td>
<td>A-C, A-B, and B-C</td>
</tr>
<tr>
<td>PAIRED (A1-A2) : (B1-B2);</td>
<td>A1-B1 and A2-B2</td>
</tr>
</tbody>
</table>

**VAR Statement**

\[ \text{VAR variables } / \text{ options} ; \]

The **VAR** statement names the variables to be used in the analyses. One-sample comparisons are conducted when the **VAR** statement is used without the **CROSSOVER=** option or **CLASS** statement. Two-independent-sample comparisons are conducted when the **VAR** statement is used with a **CLASS** statement.

An AB/BA crossover analysis is conducted when the **CROSSOVER=** option is used in the **VAR** statement. In this case, you must specify an even number of variables. Each set of two variables represents the responses in the first and second periods of the AB/BA crossover design. For example, if you use the **CROSSOVER=** option and specify `VAR x1 x2 x3 x4`, then you will get two analyses. One analysis will have `x1` as the period 1 response and `x2` as the period 2 response. The other analysis will have `x3` as the period 1 response and `x4` as the period 2 response.

The **VAR** statement cannot be used with the **PAIRED** statement. If the **VAR** statement is omitted, all numeric variables in the input data set (except a numeric variable appearing in the **BY**, **CLASS**, **FREQ**, or **WEIGHT** statement) are included in the analysis.

You can specify the following options after a slash (/):

- **CROSSOVER=** (variable1 variable2)
  
  Specifies the variables representing the treatment applied in each of the two periods in an AB/BA crossover design. The treatment variables must have two, and only two, levels. For any given observation, the levels for the two variables must be different, due to the restrictions of the AB/BA crossover design. You can use either numeric or character variables.

  Treatment levels are determined from the formatted values of the variables. Thus, you can use formats to define the treatment levels. See the discussions of the **FORMAT** procedure, the **FORMAT** statement, formats, and informats in *SAS Language Reference: Dictionary*.

- **IGNOREPERIOD**
  
  Indicates that the period effect shall be ignored—that is, assumed to be equal to 0 (if **TEST=DIFF**) or 1 (if **TEST=RATIO**). This assumption increases the degrees of freedom for the test of the treatment
difference by one and is usually more powerful, but it risks incorrect results if there is actually a period effect.

**WEIGHT Statement**

`WEIGHT variable ;`

The **WEIGHT** statement weights each observation in the input data set by the value of the **WEIGHT** variable. The values of the **WEIGHT** variable can be nonintegral, and they are not truncated. Observations with negative, zero, or missing values for the **WEIGHT** variable are not used in the analyses. Each observation is assigned a weight of 1 when the **WEIGHT** statement is not used. The **WEIGHT** statement cannot be used with an input data set of summary statistics.

**Details: TTEST Procedure**

**Input Data Set of Statistics**

PROC TTEST accepts data containing either observation values or summary statistics. Observation values are supported for all analyses, whereas summary statistics are supported only for a subset of analyses. If the analysis involves the paired design, the AB/BA crossover design, or the lognormal distributional assumption (DIST=LOGNORMAL), then observation values must be used. The graphical results are unavailable if your input data set contains summary statistics rather than raw observed values.

PROC TTEST assumes that the DATA= data set contains statistics if it contains a character variable with name _TYPE_ or _STAT_. The TTEST procedure expects this character variable to contain the names of statistics. If both _TYPE_ and _STAT_ variables exist and are of type character, PROC TTEST expects _TYPE_ to contain the names of statistics including ‘N’, ‘MEAN’, and ‘STD’ for each BY group (or for each class within each BY group for two-sample t tests). If no ‘N’, ‘MEAN’, or ‘STD’ statistics exist, an error message is printed.

**FREQ, WEIGHT, and PAIRED statements cannot be used with input data sets of statistics. BY, CLASS, and VAR statements are the same regardless of data set type. For paired comparisons, see the _DIF_ values for the _TYPE_ = T observations in output produced by the OUTSTATS= option in the PROC COMPARE statement (see the Base SAS Procedures Guide).**

**Missing Values**

An observation is omitted from the calculations if it has a missing value for either the CLASS variable, a CROSSOVER= variable, a PAIRED variable, the variable to be tested (in a one-sample or two-independent-
sample design), or either of the two response variables (in a crossover design). If more than one variable or pair of variables is listed in the VAR statement, a missing value in one variable or pair does not eliminate the observation from the analysis of other nonmissing variables or variable pairs.

**Computational Methods**

This section describes the computational formulas for the estimates, confidence limits, and tests for each analysis in the TTEST procedure. The first subsection defines some common notation. The second subsection discusses the distinction between arithmetic and geometric means. The third subsection explains the concept of the coefficient of variation. The following four subsections address the four supported designs (one-sample, paired, two-independent-sample, and AB/BA crossover). The content in each of those subsections is divided into separate discussions according to different values of the DIST= and TEST= options in the PROC TTEST statement. The last subsection describes TOST equivalence analyses.

**Common Notation**

Table 95.4 displays notation for some of the commonly used symbols.

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>( \mu )</td>
<td>Population value of (arithmetic) mean</td>
</tr>
<tr>
<td>( \mu_0 )</td>
<td>Null value of test (value of H0= option in PROC TTEST statement)</td>
</tr>
<tr>
<td>( \sigma^2 )</td>
<td>Population variance</td>
</tr>
<tr>
<td>( \sigma )</td>
<td>Population value of standard deviation</td>
</tr>
<tr>
<td>( \gamma )</td>
<td>Population value of geometric mean</td>
</tr>
<tr>
<td>CV</td>
<td>Population value of coefficient of variation (ratio of population standard deviation and population arithmetic mean)</td>
</tr>
<tr>
<td>( \alpha )</td>
<td>Value of ALPHA= option in PROC TTEST statement</td>
</tr>
<tr>
<td>( t_{p,v} )</td>
<td>( p )th percentile of ( t ) distribution with ( v ) degrees of freedom (d.f.)</td>
</tr>
<tr>
<td>( F_{p,v_1,v_2} )</td>
<td>( p )th percentile of ( F ) distribution with ( v_1 ) numerator d.f. and ( v_2 ) denominator d.f.</td>
</tr>
<tr>
<td>( \chi^2_{p,v} )</td>
<td>( p )th percentile of chi-square distribution with ( v ) d.f.</td>
</tr>
</tbody>
</table>

**Arithmetic and Geometric Means**

The *arithmetic mean* (more commonly called simply the *mean*) of the distribution of a random variable \( X \) is its expected value, \( E(X) \). The arithmetic mean is the natural parameter of interest for a normal distribution because the distribution of the difference of normal random variables has a known normal distribution, and the arithmetic mean of a normal difference is equal to the difference of the individual arithmetic means. (No such convenient property holds for geometric means with normal data, with either differences or ratios.)
The usual estimate of an arithmetic mean is the sum of the values divided by the number of values:

$$\text{arithmetic mean} = \frac{1}{n} \sum_{i=1}^{n} y_i$$

The geometric mean of the distribution of a random variable $X$ is $\exp(E(\log(X)))$, the exponentiation of the mean of the natural logarithm. The geometric mean is the natural parameter of interest for a lognormal distribution because the distribution of a ratio of lognormal random variables has a known lognormal distribution, and the geometric mean of a lognormal ratio is equal to the ratio of the individual geometric means. (No such convenient property holds for arithmetic means with lognormal data, with either differences or ratios.)

The usual estimate of a geometric mean is the product of the values raised to the power $1/n$, where $n$ is the number of values:

$$\text{geometric mean} = \left( \prod_{i=1}^{n} y_i \right)^{1/n}$$

**Coefficient of Variation**

The coefficient of variation (abbreviated “CV”) of the distribution of a random variable $X$ is the ratio of the standard deviation to the (arithmetic) mean, or $\sqrt{\text{Var}(X)}/E(X)$. Conceptually, it is a measure of the variability of $X$ expressed in units corresponding to the mean of $X$.

For lognormal data, the CV is the natural measure of variability (rather than the standard deviation) because the CV is invariant to multiplication of a lognormal variable by a constant. For a two-independent-sample design, the assumption of equal CVs on a lognormal scale is analogous to the assumption of equal variances on the normal scale. When the CVs of two independent samples of lognormal data are assumed equal, the pooled estimate of variability is used.

**One-Sample Design**

Define the following notation:

- $n^* =$ number of observations in data set
- $y_i =$ value of $i$th observation, $i \in \{1, \ldots, n^*\}$
- $f_i =$ frequency of $i$th observation, $i \in \{1, \ldots, n^*\}$
- $w_i =$ weight of $i$th observation, $i \in \{1, \ldots, n^*\}$
- $n = $ sample size $= \sum_{i} f_i$
Normal Data (DIST=NORMAL)

The mean estimate $\bar{y}$, standard deviation estimate $s$, and standard error SE are computed as follows:

$$\bar{y} = \frac{\sum_{i}^{n^*} f_i w_i y_i}{\sum_{i}^{n^*} f_i w_i}$$

$$s = \left( \frac{\sum_{i}^{n^*} f_i w_i (y_i - \bar{y})^2}{n - 1} \right)^{\frac{1}{2}}$$

$$SE = \frac{s}{\sum_{i}^{n^*} f_i w_i}$$

The $100(1 - \alpha)$% confidence interval for the mean $\mu$ is

$$\left( \bar{y} - t_{1-\frac{\alpha}{2},n-1}SE , \bar{y} + t_{1-\frac{\alpha}{2},n-1}SE \right)$$

SIDES=2

$$(-\infty , \bar{y} + t_{1-\alpha,n-1}SE)$$

SIDES=L

$$\left( \bar{y} - t_{1-\alpha,n-1}SE , \infty \right)$$

SIDES=U

The $t$ value for the test is computed as

$$t = \frac{\bar{y} - \mu_0}{SE}$$

The $p$-value of the test is computed as

$$p\text{-value} = \begin{cases} P \left( t^2 > F_{1-\alpha,1,n-1} \right) , & \text{2-sided} \\ P \left( t < t_{\alpha,n-1} \right) , & \text{lower 1-sided} \\ P \left( t > t_{1-\alpha,n-1} \right) , & \text{upper 1-sided} \end{cases}$$

The equal-tailed confidence interval for the standard deviation (CI=EQUAL) is based on the acceptance region of the test of $H_0: \sigma = \sigma_0$ that places an equal amount of area ($\frac{\alpha}{2}$) in each tail of the chi-square distribution:

$$\left\{ \chi^2_{\frac{\alpha}{2},n-1} \leq \frac{(n - 1)s^2}{\sigma^2_0} \leq \chi^2_{1-\frac{\alpha}{2},n-1} \right\}$$

The acceptance region can be algebraically manipulated to give the following $100(1 - \alpha)$% confidence interval for $\sigma^2$:

$$\left( \frac{(n - 1)s^2}{\chi^2_{1-\frac{\alpha}{2},n-1}} , \frac{(n - 1)s^2}{\chi^2_{\frac{\alpha}{2},n-1}} \right)$$

Taking the square root of each side yields the $100(1 - \alpha)$% CI=EQUAL confidence interval for $\sigma$:

$$\left( \left( \frac{(n - 1)s^2}{\chi^2_{1-\frac{\alpha}{2},n-1}} \right)^{\frac{1}{2}} , \left( \frac{(n - 1)s^2}{\chi^2_{\frac{\alpha}{2},n-1}} \right)^{\frac{1}{2}} \right)$$

The other confidence interval for the standard deviation (CI=UMPU) is derived from the uniformly most powerful unbiased test of $H_0: \sigma = \sigma_0$ (Lehmann 1986). This test has acceptance region

$$\left\{ c_1 \leq \frac{(n - 1)s^2}{\sigma^2_0} \leq c_2 \right\}$$
where the critical values \( c_1 \) and \( c_2 \) satisfy
\[
\int_{c_1}^{c_2} f_{n-1}(y) dy = 1 - \alpha
\]
and
\[
\int_{c_1}^{c_2} y f_{n-1}(y) dy = (n - 1)(1 - \alpha)
\]
where \( f_v(y) \) is the p.d.f. of the chi-square distribution with \( v \) degrees of freedom. This acceptance region can be algebraically manipulated to arrive at
\[
P \left\{ \frac{(n - 1)s^2}{c_2} \leq \sigma^2 \leq \frac{(n - 1)s^2}{c_1} \right\} = 1 - \alpha
\]
where \( c_1 \) and \( c_2 \) solve the preceding two integrals. To find the area in each tail of the chi-square distribution to which these two critical values correspond, solve \( c_1 = \chi^2_{1-\alpha_1,n-1} \) and \( c_2 = \chi^2_{\alpha_2,n-1} \) for \( \alpha_1 \) and \( \alpha_2 \); the resulting \( \alpha_1 \) and \( \alpha_2 \) sum to \( \alpha \). Hence, a 100(1 - \( \alpha \))% confidence interval for \( \sigma^2 \) is given by
\[
\left( \frac{(n - 1)s^2}{\chi^2_{1-\alpha_2,n-1}}, \frac{(n - 1)s^2}{\chi^2_{\alpha_1,n-1}} \right)
\]
Taking the square root of each side yields the 100(1 - \( \alpha \))% CI=UMPU confidence interval for \( \sigma \):
\[
\left( \sqrt{\frac{(n - 1)s^2}{\chi^2_{1-\alpha_2,n-1}}}, \sqrt{\frac{(n - 1)s^2}{\chi^2_{\alpha_1,n-1}}} \right)^{\frac{1}{2}}
\]

**Lognormal Data (DIST=LOGNORMAL)**

The DIST=LOGNORMAL analysis is handled by log-transforming the data and null value, performing a DIST=NORMAL analysis, and then transforming the results back to the original scale. This simple technique is based on the properties of the lognormal distribution as discussed in Johnson, Kotz, and Balakrishnan (1994, Chapter 14).

Taking the natural logarithms of the observation values and the null value, define
\[
z_i = \log(y_i), \quad i \in \{1, \ldots, n^*\}
\]
\[
\gamma_0 = \log(\mu_0)
\]

First a DIST=NORMAL analysis is performed on \( z_i \) in place of \( y_i \). The geometric mean estimate \( \hat{\gamma} \) and CV estimate \( \hat{C_V} \) of the original lognormal data are computed as follows:
\[
\hat{\gamma} = \exp(\bar{y})
\]
\[
\hat{C_V} = \left( \exp(s^2) - 1 \right)^{\frac{1}{2}}
\]

The \( t \) value and \( p \)-value remain the same. The confidence limits for the geometric mean and CV on the original lognormal scale are computed from the confidence limits for the arithmetic mean and standard deviation in the DIST=NORMAL analysis on the log-transformed data, in the same way that \( \hat{\gamma} \) is derived from \( \bar{y} \) and \( \hat{C_V} \) is derived from \( s \).
Paired Design

Define the following notation:

\[ n^* = \text{number of observations in data set} \]
\[ y_{1i} = \text{value of } i\text{th observation for first PAIRED variable, } i \in \{1, \ldots, n^*\} \]
\[ y_{2i} = \text{value of } i\text{th observation for second PAIRED variable, } i \in \{1, \ldots, n^*\} \]
\[ f_i = \text{frequency of } i\text{th observation, } i \in \{1, \ldots, n^*\} \]
\[ w_i = \text{weight of } i\text{th observation, } i \in \{1, \ldots, n^*\} \]
\[ n = \text{sample size} = \sum_{i}^{n^*} f_i \]

Normal Difference (DIST=NORMAL TEST=DIFF)

The analysis is the same as the analysis for the one-sample design in the section “Normal Data (DIST=NORMAL)” on page 8018 based on the differences

\[ d_i = y_{1i} - y_{2i} \quad i \in \{1, \ldots, n^*\} \]

Lognormal Ratio (DIST=LOGNORMAL TEST=RATIO)

The analysis is the same as the analysis for the one-sample design in the section “Lognormal Data (DIST=LOGNORMAL)” on page 8019 based on the ratios

\[ r_i = \frac{y_{1i}}{y_{2i}} \quad i \in \{1, \ldots, n^*\} \]

Normal Ratio (DIST=NORMAL TEST=RATIO)

The hypothesis \( H_0: \mu_1/\mu_2 = \mu_0 \), where \( \mu_1 \) and \( \mu_2 \) are the means of the first and second PAIRED variables, respectively, can be rewritten as \( H_0: \mu_1 - \mu_0 \mu_2 = 0 \). The \( t \) value and \( p \)-value are computed in the same way as in the one-sample design in the section “Normal Data (DIST=NORMAL)” on page 8018 based on the transformed values

\[ z_i = y_{1i} - \mu_0 y_{2i} \quad i \in \{1, \ldots, n^*\} \]

Estimates and confidence limits are not computed for this situation.
Two-Independent-Sample Design

Define the following notation:

- \( n_1^* \) = number of observations at first class level
- \( n_2^* \) = number of observations at second class level
- \( y_{1i} \) = value of \( i \)th observation at first class level, \( i \in \{1, \ldots, n_1^*\} \)
- \( y_{2i} \) = value of \( i \)th observation at second class level, \( i \in \{1, \ldots, n_2^*\} \)
- \( f_{1i} \) = frequency of \( i \)th observation at first class level, \( i \in \{1, \ldots, n_1^*\} \)
- \( f_{2i} \) = frequency of \( i \)th observation at second class level, \( i \in \{1, \ldots, n_2^*\} \)
- \( w_{1i} \) = weight of \( i \)th observation at first class level, \( i \in \{1, \ldots, n_1^*\} \)
- \( w_{2i} \) = weight of \( i \)th observation at second class level, \( i \in \{1, \ldots, n_2^*\} \)

\[ n_1 = \text{sample size for first class level} = \sum_{i} f_{1i} \]

\[ n_2 = \text{sample size for second class level} = \sum_{i} f_{2i} \]

**Normal Difference (DIST=NORMAL TEST=DIFF)**

Observations at the first class level are assumed to be distributed as \( N(\mu_1, \sigma_1^2) \), and observations at the second class level are assumed to be distributed as \( N(\mu_2, \sigma_2^2) \), where \( \mu_1, \mu_2, \sigma_1, \) and \( \sigma_2 \) are unknown.

The within-class-level mean estimates (\( \bar{y}_1 \) and \( \bar{y}_2 \)), standard deviation estimates (\( s_1 \) and \( s_2 \)), standard errors (\( SE_1 \) and \( SE_2 \)), and confidence limits for means and standard deviations are computed in the same way as for the one-sample design in the section “Normal Data (DIST=NORMAL)” on page 8018.

The mean difference \( \mu_1 - \mu_2 = \mu_d \) is estimated by

\[ \bar{y}_d = \bar{y}_1 - \bar{y}_2 \]

Under the assumption of equal variances (\( \sigma_1^2 = \sigma_2^2 \)), the pooled estimate of the common standard deviation is

\[ s_p = \left( \frac{(n_1 - 1)s_1^2 + (n_2 - 1)s_2^2}{n_1 + n_2 - 2} \right)^{\frac{1}{2}} \]

The pooled standard error (the estimated standard deviation of \( \bar{y}_d \) assuming equal variances) is

\[ SE_p = s_p \left( \frac{1}{\sum_{i=1}^{n_1^*} f_{1i} w_{1i}} + \frac{1}{\sum_{i=1}^{n_2^*} f_{2i} w_{2i}} \right)^{\frac{1}{2}} \]

The pooled 100(1 - \( \alpha \))% confidence interval for the mean difference \( \mu_d \) is

\[ (\bar{y}_d - t_{1-\frac{\alpha}{2},n_1+n_2-2}SE_p \ , \ \bar{y}_d + t_{1-\frac{\alpha}{2},n_1+n_2-2}SE_p ) \] \( \text{SIDES=2} \)

\[ (-\infty \ , \ \bar{y}_d + t_{1-\alpha,n_1+n_2-2}SE_p ) \] \( \text{SIDES=L} \)

\[ (\bar{y}_d - t_{1-\alpha,n_1+n_2-2}SE_p \ , \ \infty) \] \( \text{SIDES=U} \)
The \( t \) value for the pooled test is computed as

\[
t_p = \frac{\bar{y}_d - \mu_0}{SE_p}
\]

The \( p \)-value of the test is computed as

\[
p\text{-value} = \begin{cases} 
    P \left( t^2_p > F_{1-\alpha,1,n_1+n_2-2} \right), & \text{2-sided} \\
    P \left( t_p < t_{\alpha,n_1+n_2-2} \right), & \text{lower 1-sided} \\
    P \left( t_p > t_{1-\alpha,n_1+n_2-2} \right), & \text{upper 1-sided}
\end{cases}
\]

Under the assumption of unequal variances (the Behrens-Fisher problem), the unpooled standard error is computed as

\[
SE_u = \left( \frac{s_1^2}{\sum_{i=1}^{n_1} w_i} + \frac{s_2^2}{\sum_{i=1}^{n_2} w_i} \right)^{\frac{1}{2}}
\]

Satterthwaite’s (1946) approximation for the degrees of freedom, extended to accommodate weights, is computed as

\[
df_u = \frac{SE^4_u}{\left( \sum_{i=1}^{n_1} w_i \right)^2 + \left( \sum_{i=1}^{n_2} w_i \right)^2}
\]

The unpooled Satterthwaite 100(1 - \( \alpha \))% confidence interval for the mean difference \( \mu_d \) is

\[
\left( \bar{y}_d - t_{1-\frac{\alpha}{2},df_u} SE_u, \bar{y}_d + t_{1-\frac{\alpha}{2},df_u} SE_u \right), \quad \text{SIDES=2}
\]

\[
\left( -\infty, \bar{y}_d + t_{1-\alpha,df_u} SE_u \right), \quad \text{SIDES=L}
\]

\[
\left( \bar{y}_d - t_{1-\alpha,df_u} SE_u, \infty \right), \quad \text{SIDES=U}
\]

The \( t \) value for the unpooled Satterthwaite test is computed as

\[
t_u = \frac{\bar{y}_d - \mu_0}{SE_u}
\]

The \( p \)-value of the unpooled Satterthwaite test is computed as

\[
p\text{-value} = \begin{cases} 
    P \left( t^2_u > F_{1-\alpha,1,df_u} \right), & \text{2-sided} \\
    P \left( t_u < t_{\alpha,df_u} \right), & \text{lower 1-sided} \\
    P \left( t_u > t_{1-\alpha,df_u} \right), & \text{upper 1-sided}
\end{cases}
\]

When the COCHRAN option is specified in the PROC TTEST statement, the Cochran and Cox (1950) approximation of the \( p \)-value of the \( t_u \) statistic is the value of \( p \) such that

\[
t_u = \frac{t_1}{\sqrt{\sum_{i=1}^{n_1} w_i}} + \frac{t_2}{\sqrt{\sum_{i=1}^{n_2} w_i}}
\]

\[
\left( \frac{s_1^2}{\sum_{i=1}^{n_1} w_i} \right) + \frac{s_2^2}{\sum_{i=1}^{n_2} w_i}
\]

\[
\left( \frac{s_2^2}{\sum_{i=1}^{n_1} w_i} \right) + \frac{s_2^2}{\sum_{i=1}^{n_2} w_i}
\]
where $t_1$ and $t_2$ are the critical values of the $t$ distribution corresponding to a significance level of $p$ and sample sizes of $n_1$ and $n_2$, respectively. The number of degrees of freedom is undefined when $n_1 \neq n_2$. In general, the Cochran and Cox test tends to be conservative (Lee and Gurland 1975).

The 100(1 $- \alpha$)% CI=EQUAL and CI=UMPU confidence intervals for the common population standard deviation $\sigma$ assuming equal variances are computed as discussed in the section “Normal Data (DIST=NORMAL)” on page 8018 for the one-sample design, except replacing $s^2$ by $\sigma^2_p$ and $(n - 1)$ by $(n_1 + n_2 - 1)$.

The folded form of the $F$ statistic, $F'$, tests the hypothesis that the variances are equal (Steel and Torrie 1980), where

$$F' = \frac{\max(s_1^2, s_2^2)}{\min(s_1^2, s_2^2)}$$

A test of $F'$ is a two-tailed $F$ test because you do not specify which variance you expect to be larger. The $p$-value gives the probability of a greater $F$ value under the null hypothesis that $\sigma_1^2 = \sigma_2^2$. Note that this test is not very robust to violations of the assumption that the data are normally distributed, and thus it is not recommended without confidence in the normality assumption.

**Lognormal Ratio (DIST=LOGNORMAL TEST=RATIO)**

The DIST=LOGNORMAL analysis is handled by log-transforming the data and null value, performing a DIST=NORMAL analysis, and then transforming the results back to the original scale. See the section “Normal Data (DIST=NORMAL)” on page 8018 for the one-sample design for details on how the DIST=NORMAL computations for means and standard deviations are transformed into the DIST=LOGNORMAL results for geometric means and CVs. As mentioned in the section “Coefficient of Variation” on page 8017, the assumption of equal CVs on the lognormal scale is analogous to the assumption of equal variances on the normal scale.

**Normal Ratio (DIST=NORMAL TEST=RATIO)**

The distributional assumptions, equality of variances test, and within-class-level mean estimates ($\bar{y}_1$ and $\bar{y}_2$), standard deviation estimates ($s_1$ and $s_2$), standard errors (SE$_1$ and SE$_2$), and confidence limits for means and standard deviations are the same as in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design.

The mean ratio $\mu_1 / \mu_2 = \mu_r$ is estimated by

$$\hat{\mu}_r = \bar{y}_1 / \bar{y}_2$$

No estimates or confidence intervals for the ratio of standard deviations are computed.

Under the assumption of equal variances ($\sigma_1^2 = \sigma_2^2$), the pooled confidence interval for the mean ratio is the
Fieller (1954) confidence interval, extended to accommodate weights. Let

\[ a_p = \frac{s_p^2 f_1 f_2 n_1 n_2 -2 - \bar{y}_2^2}{\sum_{i=1}^{n_2} f_2 w_i} \]
\[ b_p = \bar{y}_1 \bar{y}_2 \]
\[ c_p = \frac{s_p^2 f_1 f_2 n_1 n_2 -2 - \bar{y}_1^2}{\sum_{i=1}^{n_2} f_1 w_i} \]

where \( s_p \) is the pooled standard deviation defined in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design. If \( a_p \geq 0 \) (which occurs when \( \bar{y}_2 \) is too close to zero), then the pooled two-sided 100(1 - \( \alpha \))% Fieller confidence interval for \( \mu_r \) does not exist. If \( a < 0 \), then the interval is

\[ \left( -\frac{b_p}{a_p} + \frac{(b_p^2 - a_p c_p)^{\frac{1}{2}}}{a_p}, -\frac{b_p}{a_p} - \frac{(b_p^2 - a_p c_p)^{\frac{1}{2}}}{a_p} \right) \]

For the one-sided intervals, let

\[ a_p^* = \frac{s_p^2 f_1 f_2 n_1 n_2 -2 - \bar{y}_2^2}{\sum_{i=1}^{n_2} f_2 w_i} \]
\[ c_p^* = \frac{s_p^2 f_1 f_2 n_1 n_2 -2 - \bar{y}_1^2}{\sum_{i=1}^{n_2} f_1 w_i} \]

which differ from \( a_p \) and \( c_p \) only in the use of \( \alpha \) in place of \( \alpha/2 \). If \( a_p^* \geq 0 \), then the pooled one-sided 100(1 - \( \alpha \))% Fieller confidence intervals for \( \mu_r \) do not exist. If \( a_p^* < 0 \), then the intervals are

\[ \left( -\infty, -\frac{b_p}{a_p^*} - \frac{(b_p^2 - a_p^* c_p^*)^{\frac{1}{2}}}{a_p^*} \right), \text{ SIDES=L} \]
\[ \left( -\frac{b_p}{a_p^*} + \frac{(b_p^2 - a_p^* c_p^*)^{\frac{1}{2}}}{a_p^*}, \infty \right), \text{ SIDES=U} \]

The pooled \( t \) test assuming equal variances is the Sasabuchi (1988a, 1988b) test. The hypothesis \( H_0: \mu_r = \mu_o \) is rewritten as \( H_0: \mu_1 - \mu_0 \mu_2 = 0 \), and the pooled \( t \) test in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design is conducted on the original \( y_{1i} \) values (\( i \in \{1, \ldots, n_1^* \} \)) and transformed values of \( y_{2i}^* \)

\[ y_{2i}^* = \mu_0 y_{2i} , \quad i \in \{1, \ldots, n_2^* \} \]

with a null difference of 0. The \( t \) value for the Sasabuchi pooled test is computed as

\[ t_p = \frac{\bar{y}_1 - \mu_0 \bar{y}_2}{s_p \left( \frac{1}{\sum_{i=1}^{n_1^*} f_i w_i} + \frac{\mu_0^2}{\sum_{i=1}^{n_2^*} f_2 w_2 i} \right)^{\frac{1}{2}}} \]
The $p$-value of the test is computed as

$$p\text{-value} = \begin{cases} P \left( t_p^2 > F_{1-\alpha, n_1+n_2-2} \right), & \text{2-sided} \\ P \left( t_p < t_{\alpha, n_1+n_2-2} \right), & \text{lower 1-sided} \\ P \left( t_p > t_{1-\alpha, n_1+n_2-2} \right), & \text{upper 1-sided} \end{cases}$$

Under the assumption of unequal variances, the unpooled Satterthwaite-based confidence interval for the mean ratio $\mu_r$ is computed according to the method in Dilba, Schaarschmidt, and Hothorn (2006), extended to accommodate weights. The degrees of freedom are computed as

$$\text{df}_u = \frac{\left( \frac{s_1^2}{\sum_{i=1}^{n_1^*} f_{1i} w_{1i}} + \frac{\hat{\mu}_r^2 s_2^2}{\sum_{i=1}^{n_2^*} f_{2i} w_{2i}} \right)^2}{(n_1-1) \left( \sum_{i=1}^{n_1^*} f_{1i} w_{1i} \right)^2 + (n_2-1) \left( \sum_{i=1}^{n_2^*} f_{2i} w_{2i} \right)^2}$$

Note that the estimate $\hat{\mu}_r = \bar{y}_1 / \bar{y}_2$ is used in $\text{df}_u$. Let

$$a_u = \frac{s_1^2 f_1^2 - \frac{g_s \text{df}_u}{2}}{\sum_{i=1}^{n_1^*} f_{1i} w_{1i}} - \frac{\bar{y}_2^2}{2},$$

$$b_u = \frac{\bar{y}_1 \bar{y}_2}{2},$$

$$c_u = \frac{s_1^2 f_1^2 - \frac{g_s \text{df}_u}{2}}{\sum_{i=1}^{n_1^*} f_{1i} w_{1i}} - \frac{\bar{y}_1^2}{2},$$

where $s_1$ and $s_2$ are the within-class-level standard deviations defined in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design. If $a_u \geq 0$ (which occurs when $\bar{y}_2$ is too close to zero), then the unpooled Satterthwaite-based two-sided 100$(1 - \alpha)\%$ confidence interval for $\mu_r$ does not exist. If $a_u < 0$, then the interval is

$$\left( -\frac{b_u}{a_u} + \left( \frac{b_u^2 - a_u c_u}{a_u} \right)^{\frac{1}{2}}, -\frac{b_u}{a_u} - \left( \frac{b_u^2 - a_u c_u}{a_u} \right)^{\frac{1}{2}} \right)$$

The $t$ test assuming unequal variances is the test derived in Tamhane and Logan (2004). The hypothesis $H_0$: $\mu_r = \mu_0$ is rewritten as $H_0$: $\mu_1 - \mu_0 \mu_2 = 0$, and the Satterthwaite $t$ test in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design is conducted on the original $y_{1i}$ values ($i \in \{1, \ldots, n_1^* \}$) and transformed values of $y_{2i}$

$$y_{2i}^* = \mu_0 y_{2i}, \quad i \in \{1, \ldots, n_2^* \}$$

with a null difference of 0. The degrees of freedom used in the unpooled $t$ test differs from the $\text{df}_u$ used in the unpooled confidence interval. The mean ratio $\mu_0$ under the null hypothesis is used in place of the estimate $\hat{\mu}_r$:

$$\text{df}_u^* = \frac{\left( \frac{s_1^2}{\sum_{i=1}^{n_1^*} f_{1i} w_{1i}} + \frac{\hat{\mu}_r^2 s_2^2}{\sum_{i=1}^{n_2^*} f_{2i} w_{2i}} \right)^2}{(n_1-1) \left( \sum_{i=1}^{n_1^*} f_{1i} w_{1i} \right)^2 + (n_2-1) \left( \sum_{i=1}^{n_2^*} f_{2i} w_{2i} \right)^2}$$
The \( t \) value for the Satterthwaite-based unpooled test is computed as

\[
t_u = \frac{\bar{y}_1 - \mu_0 \bar{y}_2}{\left( \frac{s_1^2}{\sum_{i=1}^{n_1} f_i w_{1i}} + \frac{\mu_0^2 s_2^2}{\sum_{i=1}^{n_2} f_{2i} w_{2i}} \right)^{1/2}}
\]

The \( p \)-value of the test is computed as

\[
p-value = \begin{cases} P \left( t_u > F_{1-\alpha, 1, df_u} \right), & \text{2-sided} \\ P \left( t_u < t_{\alpha, df_u} \right), & \text{lower 1-sided} \\ P \left( t_u > t_{1-\alpha, df_u} \right), & \text{upper 1-sided} \end{cases}
\]

**AB/BA Crossover Design**

Let “A” and “B” denote the two treatment values. Define the following notation:

- \( n_1^* \) = number of observations with treatment sequence AB
- \( n_2^* \) = number of observations with treatment sequence BA
- \( y_{11i} \) = response value of \( i \)th observation in sequence AB during period 1, \( i \in \{1, \ldots, n_1^*\} \)
- \( y_{12i} \) = response value of \( i \)th observation in sequence AB during period 2, \( i \in \{1, \ldots, n_1^*\} \)
- \( y_{21i} \) = response value of \( i \)th observation in sequence BA during period 1, \( i \in \{1, \ldots, n_2^*\} \)
- \( y_{22i} \) = response value of \( i \)th observation in sequence BA during period 2, \( i \in \{1, \ldots, n_2^*\} \)

So \( \{y_{11i}, \ldots, y_{11n_1^*}\} \) and \( \{y_{12i}, \ldots, y_{12n_1^*}\} \) are all observed at treatment level A, and \( \{y_{12i}, \ldots, y_{12n_1^*}\} \) and \( \{y_{21i}, \ldots, y_{21n_2^*}\} \) are all observed at treatment level B.

Define the **period difference** for an observation as the difference between period 1 and period 2 response values:

\[
p_{d_{kji}} = y_{k1i} - y_{k2i}
\]

for \( k \in \{1, 2\} \) and \( i \in \{1, \ldots, n_k^*\} \). Similarly, the **period ratio** is the ratio between period 1 and period 2 response values:

\[
p_{r_{kji}} = y_{k1i} / y_{k2i}
\]

The **crossover difference** for an observation is the difference between treatment A and treatment B response values:

\[
c_{d_{kji}} = \begin{cases} y_{k1i} - y_{k2i}, & k = 1 \\ y_{k2i} - y_{k1i}, & k = 2 \end{cases}
\]

Similarly, the **crossover ratio** is the ratio between treatment A and treatment B response values:

\[
c_{r_{kji}} = \begin{cases} y_{k1i} / y_{k2i}, & k = 1 \\ y_{k2i} / y_{k1i}, & k = 2 \end{cases}
\]
In the absence of the IGNOREPERIOD option in the PROC TTEST statement, the data are split into two groups according to treatment sequence and analyzed as a two-independent-sample design. If DIST=NORMAL, then the analysis of the treatment effect is based on the half period differences \( \{pd_{kji}/2\} \), and the analysis for the period effect is based on the half crossover differences \( \{cd_{kji}/2\} \). The computations for the normal difference analysis are the same as in the section “Normal Difference (DIST=NORMAL TEST=DIFF)” on page 8021 for the two-independent-sample design. The normal ratio analysis without the IGNOREPERIOD option is not supported for the AB/BA crossover design. If DIST=LOGNORMAL, then the analysis of the treatment effect is based on the square root of the period ratios \( \{\sqrt{pr_{kji}}\} \), and the analysis for the period effect is based on the square root of the crossover ratios \( \{\sqrt{cr_{kji}}\} \). The computations are the same as in the section “Lognormal Ratio (DIST=LOGNORMAL TEST=RATIO)” on page 8023 for the two-independent-sample design.

If the IGNOREPERIOD option is specified, then the treatment effect is analyzed as a paired analysis on the (treatment A, treatment B) response value pairs, regardless of treatment sequence. So the set of pairs is taken to be the concatenation of \( \{(y_{111}, y_{121}), \ldots, (y_{11n_1}, y_{12n_1})\} \) and \( \{(y_{211}, y_{221}), \ldots, (y_{21n_2}, y_{22n_2})\} \). The computations are the same as in the section “Paired Design” on page 8020.

See Senn (2002, Chapter 3) for a more detailed discussion of the AB/BA crossover design.

**TOST Equivalence Test**

The hypotheses for an equivalence test are

\[
H_0: \mu < \theta_L \quad \text{or} \quad \mu > \theta_U \\
H_1: \theta_L \leq \mu \leq \theta_U
\]

where \( \theta_L \) and \( \theta_U \) are the lower and upper bounds specified in the TOST option in the PROC TTEST statement, and \( \mu \) is the analysis criterion (mean, mean ratio, or mean difference, depending on the analysis). Following the two one-sided tests (TOST) procedure of Schuirmann (1987), the equivalence test is conducted by performing two separate tests:

\[
H_{a0}: \mu < \theta_L \\
H_{a1}: \mu \geq \theta_L
\]

and

\[
H_{b0}: \mu > \theta_U \\
H_{b1}: \mu \leq \theta_U
\]

The overall \( p \)-value is the larger of the two \( p \)-values of those tests.

Rejection of \( H_0 \) in favor of \( H_1 \) at significance level \( \alpha \) occurs if and only if the 100(1 - 2\( \alpha \))% confidence interval for \( \mu \) is contained completely within \( (\theta_L, \theta_U) \). So, the 100(1 - 2\( \alpha \))% confidence interval for \( \mu \) is displayed in addition to the usual 100(1 - \( \alpha \))% interval.

See Phillips (1990), Diletti, Hauschke, and Steinijans (1991), and Hauschke et al. (1999) for further discussion of equivalence testing for the designs supported in the TTEST procedure.
Displayed Output

For an AB/BA crossover design, the “CrossoverVarInfo” table shows the variables specified for the response and treatment values in each period of the design.

The summary statistics in the “Statistics” table and confidence limits in the “ConfLimits” table are displayed for certain variables and/or transformations or subgroups of the variables in the analysis, depending on the design. For a one-sample design, summary statistics are displayed for all variables in the analysis. For a paired design, statistics are displayed for the difference if you specify the TEST=DIFF option in the PROC TTEST statement, or for the ratio if you specify TEST=RATIO. For a two-independent-sample design, the statistics for each of the two groups and for the difference (if TEST=DIFF) or ratio (if TEST=RATIO) are displayed. For an AB/BA crossover design, statistics are displayed for each of the four cells in the design (all four combinations of the two periods and two treatments). If the IGNOREPERIOD option is absent, then if TEST=DIFF is specified, statistics are displayed for the treatment difference within each sequence and overall, and also for the period difference. If TEST=RATIO, statistics are displayed for the treatment ratio within each sequence and overall, and also for the period ratio. If the IGNOREPERIOD option is specified in the VAR statement, then statistics are displayed for the overall treatment difference if TEST=DIFF or for the overall treatment ratio if TEST=RATIO.

The “Statistics” table displays the following summary statistics:

- the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement
- the name of the classification variable (if two-independent-sample design) or treatment and period (if AB/BA crossover design)
- N, the number of nonmissing values
- the (arithmetic) Mean, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement
- the Geometric Mean, displayed if the DIST=LOGNORMAL option is specified in the PROC TTEST statement
- Std Dev, the standard deviation, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement
- the Coefficient of Variation, displayed if the DIST=LOGNORMAL option is specified in the PROC TTEST statement
- Std Err, the standard error of the mean, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement
- the Minimum value
- the Maximum value

The “ConfLimits” table displays the following:

- the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement
- the name of the classification variable (if two-independent-sample design) or treatment and period (if AB/BA crossover design)

- the (arithmetic) Mean, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement

- the Geometric Mean, displayed if the DIST=LOGNORMAL option is specified in the PROC TTEST statement

- $100(1 - \alpha)\%$ CL Mean, the lower and upper confidence limits for the mean. Separate pooled and Satterthwaite confidence limits are shown for the difference or ratio transformations in two-independent-sample designs and AB/BA crossover designs without the IGNOREPERIOD option.

- Std Dev, the standard deviation, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement

- the Coefficient of Variation, displayed if the DIST=LOGNORMAL option is specified in the PROC TTEST statement

- $100(1 - \alpha)\%$ CL Std Dev, the equal-tailed confidence limits for the standard deviation, displayed if the DIST=NORMAL and CI=EQUAL options are specified in the PROC TTEST statement

- $100(1 - \alpha)\%$ UMPU CL Std Dev, the UMPU confidence limits for the standard deviation, displayed if the DIST=NORMAL and CI=UMPU options are specified in the PROC TTEST statement

- $100(1 - \alpha)\%$ CL CV, the equal-tailed confidence limits for the coefficient of variation, displayed if the DIST=LOGNORMAL and CI=EQUAL options are specified in the PROC TTEST statement

- $100(1 - \alpha)\%$ UMPU CL CV, the UMPU confidence limits for the coefficient of variation, displayed if the DIST=LOGNORMAL and CI=UMPU options are specified in the PROC TTEST statement

The confidence limits in the “EquivLimits” table and test results in the “TTests” and “EquivTests” tables are displayed only for the test criteria—that is, the variables or transformations being tested. For a one-sample design, results are displayed for all variables in the analysis. For a paired design, results are displayed for the difference if you specify the TEST=DIFF option in the PROC TTEST statement, or for the ratio if you specify TEST=RATIO. For a two-independent-sample design, the results for the difference (if TEST=DIFF) or ratio (if TEST=RATIO) are displayed. For an AB/BA crossover design, results are displayed for the treatment difference (if TEST=DIFF) or ratio (if TEST=RATIO). If the IGNOREPERIOD option is absent, then results are also displayed for the period difference (if TEST=DIFF) or ratio (if TEST=RATIO).

The “EquivLimits” table, produced only if the TOST option is specified in the PROC TTEST statement, displays the following:

- the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement

- the (arithmetic) Mean, displayed if the DIST=NORMAL option is specified in the PROC TTEST statement

- the Geometric Mean, displayed if the DIST=LOGNORMAL option is specified in the PROC TTEST statement
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- Lower Bound, the lower equivalence bound for the mean specified in the TOST option in the PROC TTEST statement

- 100(1 – 2α)% CL Mean, the lower and upper confidence limits for the mean relevant to the equivalence test. Separate pooled and Satterthwaite confidence limits are shown for two-independent-sample designs and AB/BA crossover designs without the IGNOREPERIOD option.

- Upper Bound, the upper equivalence bound for the mean specified in the TOST option in the PROC TTEST statement

- Assessment, the result of the equivalence test at the significance level specified by the ALPHA= option in the PROC TTEST statement, either “Equivalent” or “Not equivalent”

The “TTests” table is produced only if the TOST option is not specified in the PROC TTEST statement. Separate results for pooled and Satterthwaite tests (and also the Cochran and Cox test, if the COCHRAN option is specified in the PROC TTEST statement) are displayed for two-independent-sample designs and AB/BA crossover designs without the IGNOREPERIOD option. The table includes the following results:

- the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement
- t Value, the t statistic for comparing the mean to the null value as specified by the H0= option in the PROC TTEST statement
- DF, the degrees of freedom
- the p-value, the probability of obtaining a t statistic at least as extreme as the observed t value under the null hypothesis

The “EquivTests” table is produced only if the TOST option is specified in the PROC TTEST statement. Separate results for pooled and Satterthwaite tests are displayed for two-independent-sample designs and AB/BA crossover designs without the IGNOREPERIOD option. Each test consists of two separate one-sided tests. The overall p-value is the larger p-value from these two tests. The table includes the following results:

- the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement
- Null, the lower equivalence bound for the Upper test or the upper equivalence bound for the Lower test, as specified by the TOST option in the PROC TTEST statement
- t Value, the t statistic for comparing the mean to the Null value
- DF, the degrees of freedom
- the p-value, the probability of obtaining a t statistic at least as extreme as the observed t value under the null hypothesis

The “Equality” table gives the results of the test of equality of variances. It is displayed for two-independent-sample designs and AB/BA crossover designs without the IGNOREPERIOD option. The table includes the following results:
• the name of the variable(s), displayed if the NOBYVAR option is used in the PROC TTEST statement
• Num DF and Den DF, the numerator and denominator degrees of freedom
• F Value, the $F'$ (folded) statistic
• Pr > F, the probability of a greater $F'$ value. This is the two-tailed $p$-value.

ODS Table Names

PROC TTEST assigns a name to each table it creates. You can use these names to reference the table when using the Output Delivery System (ODS) to select tables and create output data sets. These names are listed in Table 95.5. For more information about ODS, see Chapter 20, “Using the Output Delivery System.”

<table>
<thead>
<tr>
<th>ODS Table Name</th>
<th>Description</th>
<th>Syntax</th>
</tr>
</thead>
<tbody>
<tr>
<td>ConfLimits</td>
<td>$100(1 - \alpha)%$ confidence limits for means, standard deviations, and/or coefficients of variation</td>
<td>By default</td>
</tr>
<tr>
<td>Equality</td>
<td>Tests for equality of variance</td>
<td>CLASS statement or VAR / CROSSOVER=</td>
</tr>
<tr>
<td>EquivLimits</td>
<td>$100(1 - 2\alpha)%$ confidence limits for means</td>
<td>PROC TTEST TOST</td>
</tr>
<tr>
<td>EquivTests</td>
<td>Equivalence $t$ tests</td>
<td>PROC TTEST TOST</td>
</tr>
<tr>
<td>Statistics</td>
<td>Univariate summary statistics</td>
<td>By default</td>
</tr>
<tr>
<td>TTests</td>
<td>$t$ tests</td>
<td>By default</td>
</tr>
</tbody>
</table>

ODS Graphics

Statistical procedures use ODS Graphics to create graphs as part of their output. ODS Graphics is described in detail in Chapter 21, “Statistical Graphics Using ODS.”

Before you create graphs, ODS Graphics must be enabled (for example, with the ODS GRAPHICS ON statement). For more information about enabling and disabling ODS Graphics, see the section “Enabling and Disabling ODS Graphics” on page 609 in Chapter 21, “Statistical Graphics Using ODS.”

The overall appearance of graphs is controlled by ODS styles. Styles and other aspects of using ODS Graphics are discussed in the section “A Primer on ODS Statistical Graphics” on page 608 in Chapter 21, “Statistical Graphics Using ODS.”

ODS Graph Names

You can reference every graph produced through ODS Graphics with a name. The names of the graphs that PROC TTEST generates are listed in Table 95.6, along with the required statements and options.
### Table 95.6 Graphs Produced by PROC TTEST

<table>
<thead>
<tr>
<th>ODS Graph Name</th>
<th>Plot Description</th>
<th>Option</th>
</tr>
</thead>
<tbody>
<tr>
<td>AgreementOfPeriods</td>
<td>Plot of period 2 against period 1 response values for an AB/BA crossover design</td>
<td>VAR / CROSSOVER=</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PLOTS=AGREEMENT(TYPE=PERIOD)</td>
</tr>
<tr>
<td>AgreementOfTreatments</td>
<td>Plot of second treatment against first treatment response values for an AB/BA</td>
<td>VAR / CROSSOVER=</td>
</tr>
<tr>
<td></td>
<td>crossover design</td>
<td>PLOTS=AGREEMENT</td>
</tr>
<tr>
<td>AgreementPlot</td>
<td>Plot of second response against first response for a paired design</td>
<td>PAIRED statement</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PLOTS=AGREEMENT</td>
</tr>
<tr>
<td>BoxPlot</td>
<td>Box plots, also with confidence band for one-sample or paired design</td>
<td>PLOTS=BOX</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PLOTS=SUMMARY(UNPACK)</td>
</tr>
<tr>
<td>Histogram</td>
<td>Histograms with overlaid kernel densities, and also normal densities if DIST=NORMAL</td>
<td>PLOTS=HISTOGRAM</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PLOTS=SUMMARY(UNPACK)</td>
</tr>
<tr>
<td>Interval</td>
<td>Confidence intervals for means</td>
<td>PLOTS=INTERVAL</td>
</tr>
<tr>
<td>ProfilesOverPrd</td>
<td>Plot of response profiles over periods 1 and 2 for an AB/BA crossover design</td>
<td>VAR / CROSSOVER=</td>
</tr>
<tr>
<td></td>
<td></td>
<td>PLOTS=PROFILES(TYPE=PERIOD)</td>
</tr>
<tr>
<td>ProfilesOverTrt</td>
<td>Plot of response profiles over first and second treatments for an AB/BA crossover</td>
<td>VAR / CROSSOVER=</td>
</tr>
<tr>
<td></td>
<td>design</td>
<td>PLOTS=PROFILES</td>
</tr>
<tr>
<td>ProfilesPlot</td>
<td>Plot of response profiles over first and second response values for a paired</td>
<td>PAIRED statement</td>
</tr>
<tr>
<td></td>
<td>design</td>
<td>PLOTS=PROFILES</td>
</tr>
<tr>
<td>QQPlot</td>
<td>Normal quantile-quantile plots</td>
<td>PLOTS=QQ</td>
</tr>
<tr>
<td>SummaryPanel</td>
<td>Histograms with overlaid kernel densities (and also normal densities if DIST=NORMAL) and box plots (and also with confidence band for one-sample or paired design)</td>
<td>PLOTS=SUMMARY</td>
</tr>
</tbody>
</table>

### Interpreting Graphs

**Agreement Plots for Paired Designs**

For paired designs, the second response of each pair is plotted against the first response, with the mean shown as a large bold symbol. If the `WEIGHT` statement is used, then the mean is the weighted mean. A diagonal line with slope=0 and y-intercept=1 is overlaid. The location of the points with respect to the
diagonal line reveals the strength and direction of the difference or ratio. The tighter the clustering along the same direction as the line, the stronger the positive correlation of the two measurements for each subject. Clustering along a direction perpendicular to the line indicates negative correlation.

**Period Agreement Plots for Crossover Designs**

The response in the second period is plotted against the response in the first period, with plot symbols distinguishing the two treatment sequences and the two sequence means shown larger in bold. If the WEIGHT statement is used, then the means are weighted means. A diagonal line with slope=0 and y-intercept=1 is overlaid.

In the absence of a strong period effect, the points from each sequence will appear as mirror images about the diagonal line, farther apart with stronger treatment effects. Deviations from symmetry about the diagonal line indicate a period effect. The spread of points within each treatment sequence is an indicator of between-subject variability. The tighter the clustering along the same direction as the line (within each treatment sequence), the stronger the positive correlation of the two measurements for each subject. Clustering along a direction perpendicular to the line indicates negative correlation.

The period agreement plot is usually less informative than the treatment agreement plot. The exception is when the period effect is stronger than the treatment effect.

**Treatment Agreement Plots for Crossover Designs**

The response associated with the second treatment is plotted against the response associated with the first treatment, with plot symbols distinguishing the two treatment sequences and the two sequence means shown larger in bold. If the WEIGHT statement is used, then the means are weighted means. A diagonal line with slope=0 and y-intercept=1 is overlaid.

The location of the points with respect to the diagonal line reveals the strength and direction of the treatment effect. Substantial location differences between the two sequences indicates a strong period effect. The spread of points within each treatment sequence is an indicator of between-subject variability. The tighter the clustering along the same direction as the line (within each treatment sequence), the stronger the positive correlation of the two measurements for each subject. Clustering along a direction perpendicular to the line indicates negative correlation.

**Box Plots**

The box is drawn from the 25th percentile (lower quartile) to the 75th percentile (upper quartile). The vertical line inside the box shows the location of the median. If DIST=NORMAL, then a diamond symbol shows the location of the mean. The whiskers extend to the minimum and maximum observations, and circles beyond the whiskers identify outliers.

For one-sample and paired designs, a confidence interval for the mean is shown as a band in the background. If the analysis is an equivalence analysis (with the TOST option in the PROC TTEST statement), then the interval is a $100(1 - 2\alpha)\%$ confidence interval shown along with the equivalence bounds. The inclusion of this interval completely within the bounds is indicative of a significant $p$-value. If the analysis is not an equivalence analysis, then the confidence level is $100(1 - \alpha)\%$. If the SHOWH0 global plot option is used,
then the null value for the test is shown. If the WEIGHT statement is used, then weights are incorporated in the confidence intervals.

**Histograms**

The WEIGHT statement is ignored in the computation of the normal and kernel densities.

**Confidence Intervals**

If the analysis is an equivalence analysis (with the TOST option in the PROC TTEST statement), then unless the TYPE=PERGROUP option is used, the interval is a $100(1 - 2\alpha)\%$ mean confidence interval shown along with the equivalence bounds. The inclusion of this interval completely within the bounds is indicative of a significant $p$-value.

If the analysis is not an equivalence analysis, or if the TYPE=PERGROUP option is used, then the confidence level is $100(1 - \alpha)\%$. If the SHOWH0 global plot option is used, then the null value for the test is shown.

If the SIDES=L or SIDES=U option is used in the PROC TTEST statement, then the unbounded side of the one-sided interval is represented with an arrowhead. Note that the actual location of the arrowhead is irrelevant.

If the WEIGHT statement is used, then weights are incorporated in the confidence intervals.

**Profiles for Paired Designs**

For paired designs, a line is drawn for each observation from left to right connecting the first response to the second response. The mean first response and mean second response are connected with a bold line. If the WEIGHT statement is used, then the means are weighted means. The more extreme the slope, the stronger the effect. A wide spread of profiles indicates high between-subject variability. Consistent positive slopes indicate strong positive correlation. Widely varying slopes indicate lack of correlation, while consistent negative slopes indicate strong negative correlation.

**Profiles over Period for Crossover Designs**

For each observation, the response for the first period is connected to the response for second period, regardless of the treatment applied in each period. The means for each treatment sequence are shown in bold. If the WEIGHT statement is used, then the means are weighted means.

In the absence of a strong period effect, the profiles for each sequence will appear as mirror images about an imaginary horizontal line in the center. Deviations from symmetry about this imaginary horizontal line indicate a period effect. A wide spread of profiles within sequence indicates high between-subject variability.

The TYPE=PERIOD plot is usually less informative than the TYPE=TREATMENT plot. The exception is when the period effect is stronger than the treatment effect.
Profiles over Treatment for Crossover Designs

For each observation, the response for the first treatment is connected to the response for the second treatment, regardless of the periods in which they occur. The means for each treatment sequence are shown in bold. If the WEIGHT statement is used, then the means are weighted means.

In general, the more extreme the slope, the stronger the treatment effect. Slope differences between the two treatment sequences measure the period effect. A wide spread of profiles within sequence indicates high between-subject variability.

Q-Q Plots

Q-Q plots are useful for diagnosing violations of the normality and homoscedasticity assumptions. If the data in a Q-Q plot come from a normal distribution, the points will cluster tightly around the reference line. You can use the UNIVARIATE procedure with the NORMAL option to numerically check the normality assumption.

Examples: TTEST Procedure

Example 95.1: Using Summary Statistics to Compare Group Means

This example, taken from Huntsberger and Billingsley (1989), compares two grazing methods using 32 steers. Half of the steers are allowed to graze continuously while the other half are subjected to controlled grazing time. The researchers want to know if these two grazing methods affect weight gain differently. The data are read by the following DATA step:

```
data graze;
  length GrazeType $ 10;
  input GrazeType $ WtGain @@;
datalines;
controlled 45  controlled 62
controlled 96  controlled 128
controlled 120 controlled 99
controlled 28  controlled 50
controlled 109 controlled 115
controlled 39  controlled 96
controlled 87  controlled 100
controlled 76  controlled 80
continuous 94  continuous 12
continuous 26  continuous 89
continuous 88  continuous 96
continuous 85  continuous 130
continuous 75  continuous 54
continuous 112 continuous 69
continuous 104 continuous 95
```
The variable GrazeType denotes the grazing method: “controlled” is controlled grazing and “continuous” is continuous grazing. The dollar sign ($) following GrazeType makes it a character variable, and the trailing at signs (@@) tell the procedure that there is more than one observation per line.

If you have summary data—that is, just means and standard deviations, as computed by PROC MEANS—then you can still use PROC TTEST to perform a simple \( t \) test analysis. This example demonstrates this mode of input for PROC TTEST. Note, however, that graphics are unavailable when summary statistics are used as input.

The MEANS procedure is invoked to create a data set of summary statistics with the following statements:

```plaintext
proc sort;
  by GrazeType;
proc means data=graze noprint;
  var WtGain;
  by GrazeType;
  output out=newgraze;
run;
```

The NOPRINT option eliminates all printed output from the MEANS procedure. The VAR statement tells PROC MEANS to compute summary statistics for the WtGain variable, and the BY statement requests a separate set of summary statistics for each level of GrazeType. The OUTPUT OUT= statement tells PROC MEANS to put the summary statistics into a data set called newgraze so that it can be used in subsequent procedures. This new data set is displayed in Output 95.1.1 by using PROC PRINT as follows:

```plaintext
proc print data=newgraze;
run;
```

The _STAT_ variable contains the names of the statistics, and the GrazeType variable indicates which group the statistic is from.

**Output 95.1.1** Output Data Set of Summary Statistics

<table>
<thead>
<tr>
<th>Obs</th>
<th>GrazeType</th>
<th><em>TYPE</em></th>
<th><em>FREQ</em></th>
<th><em>STAT</em></th>
<th>WtGain</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>continuous</td>
<td>0</td>
<td>16</td>
<td>N</td>
<td>16.000</td>
</tr>
<tr>
<td>2</td>
<td>continuous</td>
<td>0</td>
<td>16</td>
<td>MIN</td>
<td>12.000</td>
</tr>
<tr>
<td>3</td>
<td>continuous</td>
<td>0</td>
<td>16</td>
<td>MAX</td>
<td>130.000</td>
</tr>
<tr>
<td>4</td>
<td>continuous</td>
<td>0</td>
<td>16</td>
<td>MEAN</td>
<td>75.188</td>
</tr>
<tr>
<td>5</td>
<td>continuous</td>
<td>0</td>
<td>16</td>
<td>STD</td>
<td>33.812</td>
</tr>
<tr>
<td>6</td>
<td>controlled</td>
<td>0</td>
<td>16</td>
<td>N</td>
<td>16.000</td>
</tr>
<tr>
<td>7</td>
<td>controlled</td>
<td>0</td>
<td>16</td>
<td>MIN</td>
<td>28.000</td>
</tr>
<tr>
<td>8</td>
<td>controlled</td>
<td>0</td>
<td>16</td>
<td>MAX</td>
<td>128.000</td>
</tr>
<tr>
<td>9</td>
<td>controlled</td>
<td>0</td>
<td>16</td>
<td>MEAN</td>
<td>83.125</td>
</tr>
<tr>
<td>10</td>
<td>controlled</td>
<td>0</td>
<td>16</td>
<td>STD</td>
<td>30.535</td>
</tr>
</tbody>
</table>

The following statements invoke PROC TTEST with the newgraze data set, as denoted by the DATA= option:

```plaintext
continuous 53 continuous 21
;
run;
```
Example 95.1: Using Summary Statistics to Compare Group Means

```
proc ttest data=newgraze;
  class GrazeType;
  var WtGain;
run;
```

The `CLASS` statement contains the variable that distinguishes between the groups being compared, in this case `GrazeType`. The summary statistics and confidence intervals are displayed first, as shown in Output 95.1.2.

### Output 95.1.2 Summary Statistics and Confidence Limits

```
<table>
<thead>
<tr>
<th>GrazeType</th>
<th>N</th>
<th>Mean</th>
<th>Std Dev</th>
<th>Std Err</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>continuous</td>
<td>16</td>
<td>75.19</td>
<td>33.81</td>
<td>8.45</td>
<td>12.00</td>
<td>130.0</td>
</tr>
<tr>
<td>controlled</td>
<td>16</td>
<td>83.13</td>
<td>30.54</td>
<td>7.63</td>
<td>28.00</td>
<td>128.0</td>
</tr>
</tbody>
</table>
```

Diff (1-2)  
-7.94
32.22
11.39

```
<table>
<thead>
<tr>
<th>GrazeType</th>
<th>Method</th>
<th>Mean</th>
<th>95% CL Mean</th>
<th>Std Dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>continuous</td>
<td></td>
<td>75.19</td>
<td>57.17</td>
<td>33.82</td>
</tr>
<tr>
<td>controlled</td>
<td></td>
<td>83.13</td>
<td>66.54</td>
<td>30.54</td>
</tr>
<tr>
<td>Diff (1-2)</td>
<td>Pooled</td>
<td>-7.94</td>
<td>-31.21</td>
<td>32.22</td>
</tr>
<tr>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>-7.94</td>
<td>-31.21</td>
<td>32.22</td>
</tr>
</tbody>
</table>
```

In Output 95.1.2, The `GrazeType` column specifies the group for which the statistics are computed. For each class, the sample size, mean, standard deviation and standard error, and maximum and minimum values are displayed. The confidence bounds for the mean are also displayed; however, since summary statistics are used as input, the confidence bounds for the standard deviation of the groups are not calculated.

Output 95.1.3 shows the results of tests for equal group means and equal variances.

### Output 95.1.3 t Tests

```
| Method      | Variances | DF  | t Value | Pr > |t| |
|-------------|-----------|-----|---------|------|---|
| Pooled      | Equal     | 30  | -0.70   | 0.49 |12|
| Satterthwaite | Unequal  | 29.694 | -0.70 | 0.49 |13|
```
A group test statistic for the equality of means is reported for both equal and unequal variances. Both tests indicate a lack of evidence for a significant difference between grazing methods ($t = -0.70$ and $p = 0.4912$ for the pooled test, $t = -0.70$ and $p = 0.4913$ for the Satterthwaite test). The equality of variances test does not indicate a significant difference in the two variances ($F' = 1.23$, $p = 0.6981$). Note that this test assumes that the observations in both data sets are normally distributed; this assumption can be checked in PROC UNIVARIATE by using the NORMAL option with the raw data.

Although the ability to use summary statistics as input is useful if you lack access to the original data, some of the output that would otherwise be produced in an analysis on the original data is unavailable. There are also limitations on the designs and distributional assumptions that can be used with summary statistics as input. For more information, see the section “Input Data Set of Statistics” on page 8015.

**Example 95.2: One-Sample Comparison with the FREQ Statement**

This example examines children’s reading skills. The data consist of Degree of Reading Power (DRP) test scores from 44 third-grade children and are taken from Moore (1995, p. 337). Their scores are given in the following DATA step:

```plaintext
data read;
  input score count @@;
datalines;
40 2  47 2  52 2  26 1  19 2
25 2  35 4  39 1  26 1  48 1
14 2  22 1  42 1  34 2  33 2
18 1  15 1  29 1  41 2  44 1
51 1  43 1  27 2  46 2  28 1
49 1  31 1  28 1  54 1  45 1
;
run;
```

The following statements invoke the TTEST procedure to test if the mean test score is equal to 30.

```plaintext
ods graphics on;

proc ttest data=read h0=30;
  var score;
  freq count;
run;

ods graphics off;
```
The count variable contains the frequency of occurrence of each test score; this is specified in the FREQ statement. The output, shown in Output 95.2.1, contains the results.

**Output 95.2.1**  TTEST Results

<table>
<thead>
<tr>
<th>N</th>
<th>Mean</th>
<th>Std Dev</th>
<th>Std Err</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>44</td>
<td>34.8636</td>
<td>11.2303</td>
<td>1.6930</td>
<td>14.0000</td>
<td>54.0000</td>
</tr>
</tbody>
</table>

Mean 95% CL Mean Std Dev 95% CL Std Dev

| 34.8636 | 31.4493 | 38.2780 | 11.2303 | 9.2788  | 14.2291 |

DF  t Value  Pr > |t|

| 43  | 2.87  | 0.0063 |

The SAS log states that 30 observations and two variables have been read. However, the sample size given in the TTEST output is N=44. This is due to specifying the count variable in the FREQ statement. The test is significant ($t = 2.87, p = 0.0063$) at the 5% level, so you can conclude that the mean test score is different from 30.

The summary panel in **Output 95.2.2** shows a histogram with overlaid normal and kernel densities, a box plot, and the 95% confidence interval for the mean.
The Q-Q plot in Output 95.2.3 assesses the normality assumption.
The tight clustering of the points around the diagonal line is consistent with the normality assumption. You could use the UNIVARIATE procedure with the NORMAL option to numerically check the normality assumption.

Example 95.3: Paired Comparisons

When it is not feasible to assume that two groups of data are independent, and a natural pairing of the data exists, it is advantageous to use an analysis that takes the correlation into account. Using this correlation results in higher power to detect existing differences between the means. The differences between paired observations are assumed to be normally distributed. Some examples of this natural pairing are as follows:

- pre- and post-test scores for a student receiving tutoring
- fuel efficiency readings of two fuel types observed on the same automobile
- sunburn scores for two sunblock lotions, one applied to the individual’s right arm, one to the left arm
• political attitude scores of husbands and wives

In this example, taken from the *SUGI Supplemental Library User’s Guide, Version 5 Edition*, a stimulus is being examined to determine its effect on systolic blood pressure. Twelve men participate in the study. Each man’s systolic blood pressure is measured both before and after the stimulus is applied. The following statements input the data:

```plaintext
data pressure;
  input SBPbefore SBPafter @@;
datalines;
  120 128 124 131 130 131 118 127
  140 132 128 125 140 141 135 137
  126 118 130 132 126 129 127 135
;run;
```

The variables SBPbefore and SBPafter denote the systolic blood pressure before and after the stimulus, respectively.

The statements to perform the test follow:

```plaintext
ods graphics on;
proc ttest;
  paired SBPbefore*SBPafter;
run;
ods graphics off;
```

The PAIRED statement is used to test whether the mean change in systolic blood pressure is significantly different from zero. The tabular output is displayed in Output 95.3.1.

**Output 95.3.1** TTEST Results

<table>
<thead>
<tr>
<th>The TTEST Procedure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Difference: SBPbefore - SBPafter</td>
</tr>
<tr>
<td>N</td>
</tr>
<tr>
<td>12</td>
</tr>
<tr>
<td>Mean</td>
</tr>
<tr>
<td>-1.8333</td>
</tr>
<tr>
<td>DF</td>
</tr>
<tr>
<td>11</td>
</tr>
</tbody>
</table>

The variables SBPbefore and SBPafter are the paired variables with a sample size of 12. The summary statistics of the difference are displayed (mean, standard deviation, and standard error) along with their
confidence limits. The minimum and maximum differences are also displayed. The \( t \) test is not significant \((t = -1.09, p = 0.2992)\), indicating that the stimuli did not significantly affect systolic blood pressure.

The summary panel in Output 95.3.2 shows a histogram, normal and kernel densities, box plot, and \(100(1 - \alpha)\% = 95\%\) confidence interval of the \(\text{SBP}_{\text{before}} - \text{SBP}_{\text{after}}\) difference.

**Output 95.3.2** Summary Panel

![Distribution of Difference: SBPbefore - SBPafter](image)

The agreement plot in Output 95.3.3 reveals that only three men have higher blood pressure before the stimulus than after.
But the differences for these men are relatively large, keeping the mean difference only slightly negative.

The profiles plot in Output 95.3.4 is a different view of the same information contained in Output 95.3.3, plotting the blood pressure from before to after the stimulus.
**Output 95.3.4** Profiles over Treatments

The Q-Q plot in **Output 95.3.5** assesses the normality assumption.
Chapter 95: The TTEST Procedure

Output 95.3.5  Q-Q Plot

The Q-Q plot shows no obvious deviations from normality. You can check the assumption of normality more rigorously by using PROC UNIVARIATE with the NORMAL option.

Example 95.4: AB/BA Crossover Design

Senn (2002, Chapter 3) discusses a study comparing the effectiveness of two bronchodilators, formoterol ("for") and salbutamol ("sal"), in the treatment of childhood asthma. A total of 13 children are recruited for an AB/BA crossover design. A random sample of 7 of the children are assigned to the treatment sequence for/sal, receiving a dose of formoterol upon an initial visit ("period 1") and then a dose of salbutamol upon a later visit ("period 2"). The other 6 children are assigned to the sequence sal/for, receiving the treatments in the reverse order but otherwise in a similar manner. Periods 1 and 2 are sufficiently spaced so that no carryover effects are suspected. After a child inhales a dose of a bronchodilator, peak expiratory flow (PEF) is measured. Higher PEF indicates greater effectiveness. The data are assumed to be approximately normally distributed.

The data set is generated with the following statements:
data asthma;
  input Drug1 $ Drug2 $ PEF1 PEF2 @@;
datalines;
  for sal 310 270 for sal 310 260 for sal 370 300
  for sal 410 390 for sal 250 210 for sal 380 350
  for sal 330 365
  sal for 370 385 sal for 310 400 sal for 380 410
  sal for 290 320 sal for 260 340 sal for 90 220
;run;

You can display the data by using the following statements, which produce Output 95.4.1:

proc print data=asthma;
run;

Output 95.4.1  Asthma Study Data

<table>
<thead>
<tr>
<th>Obs</th>
<th>Drug1</th>
<th>Drug2</th>
<th>PEF1</th>
<th>PEF2</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>for</td>
<td>sal</td>
<td>310</td>
<td>270</td>
</tr>
<tr>
<td>2</td>
<td>for</td>
<td>sal</td>
<td>310</td>
<td>260</td>
</tr>
<tr>
<td>3</td>
<td>for</td>
<td>sal</td>
<td>370</td>
<td>300</td>
</tr>
<tr>
<td>4</td>
<td>for</td>
<td>sal</td>
<td>410</td>
<td>390</td>
</tr>
<tr>
<td>5</td>
<td>for</td>
<td>sal</td>
<td>250</td>
<td>210</td>
</tr>
<tr>
<td>6</td>
<td>for</td>
<td>sal</td>
<td>380</td>
<td>350</td>
</tr>
<tr>
<td>7</td>
<td>for</td>
<td>sal</td>
<td>330</td>
<td>365</td>
</tr>
<tr>
<td>8</td>
<td>sal</td>
<td>for</td>
<td>370</td>
<td>385</td>
</tr>
<tr>
<td>9</td>
<td>sal</td>
<td>for</td>
<td>310</td>
<td>400</td>
</tr>
<tr>
<td>10</td>
<td>sal</td>
<td>for</td>
<td>380</td>
<td>410</td>
</tr>
<tr>
<td>11</td>
<td>sal</td>
<td>for</td>
<td>290</td>
<td>320</td>
</tr>
<tr>
<td>12</td>
<td>sal</td>
<td>for</td>
<td>260</td>
<td>340</td>
</tr>
<tr>
<td>13</td>
<td>sal</td>
<td>for</td>
<td>90</td>
<td>220</td>
</tr>
</tbody>
</table>

The variables PEF1 and PEF2 represent the responses for the first and second periods, respectively. The variables Drug1 and Drug2 represent the treatment in each period.

You can analyze this crossover design by using the CROSSOVER= option after a slash (/) in the VAR statement:

ods graphics on;
proc ttest data=asthma plots=interval;
  var PEF1 PEF2 / crossover= (Drug1 Drug2);
run;
ods graphics off;

With the default PROC TTEST options TEST=DIFF and DIST=NORMAL and the lack of the IGNOREPERIOD option in the VAR statement, both the treatment difference and the period difference are assessed. The PROC TTEST default options H0=0, SIDES=2, and ALPHA=0.05 specify a two-sided analysis with 95% confidence limits comparing treatment and period differences to a default difference of zero. The default CI=EQUAL option in the PROC TTEST statement requests equal-tailed confidence intervals for
standard deviations. The PLOTS=INTERVAL option produces TYPE=TREATMENT confidence intervals, in addition to the default plots AGREEMENT(TYPE=TREATMENT), BOX, HISTOGRAM, PROFILES(TYPE=TREATMENT), and QQ.

Output 95.4.2 summarizes the response and treatment variables for each period.

**Output 95.4.2** Crossover Variable Information

<table>
<thead>
<tr>
<th>Period</th>
<th>Response</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>PEF1</td>
<td>Drug1</td>
</tr>
<tr>
<td>2</td>
<td>PEF2</td>
<td>Drug2</td>
</tr>
</tbody>
</table>

Output 95.4.3 displays basic summary statistics (sample size, mean, standard deviation, standard error, minimum, and maximum) for each of the four cells in the design, the treatment difference within each treatment sequence, the overall treatment difference, and the overall period difference.

**Output 95.4.3** Statistics

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Treatment</th>
<th>Period</th>
<th>N</th>
<th>Mean</th>
<th>Std Dev</th>
<th>Std Err</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>for</td>
<td>1</td>
<td>7</td>
<td>337.1</td>
<td>53.7631</td>
<td>20.3206</td>
</tr>
<tr>
<td>2</td>
<td>for</td>
<td>2</td>
<td>6</td>
<td>345.8</td>
<td>70.8814</td>
<td>28.9372</td>
</tr>
<tr>
<td>2</td>
<td>sal</td>
<td>1</td>
<td>6</td>
<td>283.3</td>
<td>105.4</td>
<td>43.0245</td>
</tr>
<tr>
<td>1</td>
<td>sal</td>
<td>2</td>
<td>7</td>
<td>306.4</td>
<td>64.7247</td>
<td>24.4636</td>
</tr>
<tr>
<td>1</td>
<td>Diff (1-2)</td>
<td>7</td>
<td>30.7143</td>
<td>32.9682</td>
<td>12.4608</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Diff (1-2)</td>
<td>6</td>
<td>62.5000</td>
<td>44.6934</td>
<td>18.2460</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td></td>
<td>46.6071</td>
<td>19.3702</td>
<td>10.7766</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td></td>
<td>-15.8929</td>
<td>19.3702</td>
<td>10.7766</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Treatment</th>
<th>Period</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>for</td>
<td>1</td>
<td>250.0</td>
<td>410.0</td>
</tr>
<tr>
<td>2</td>
<td>for</td>
<td>2</td>
<td>220.0</td>
<td>410.0</td>
</tr>
<tr>
<td>2</td>
<td>sal</td>
<td>1</td>
<td>90.0000</td>
<td>380.0</td>
</tr>
<tr>
<td>1</td>
<td>sal</td>
<td>2</td>
<td>210.0</td>
<td>390.0</td>
</tr>
<tr>
<td>1</td>
<td>Diff (1-2)</td>
<td></td>
<td>-35.0000</td>
<td>70.0000</td>
</tr>
<tr>
<td>2</td>
<td>Diff (1-2)</td>
<td></td>
<td>15.0000</td>
<td>130.0</td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

The treatment difference “Diff (1-2)” corresponds to the “for” treatment minus the “sal” treatment, because “for” appears before “sal” in the output, according to the ORDER=MIXED default PROC TTEST option. Its mean estimate is 46.6071, favoring formoterol over salbutamol.

The standard deviation (Std Dev) reported for a “difference” is actually the pooled standard deviation across both treatment sequence (for/sal and sal/for), assuming equal variances. The standard error (Std Err) is the standard deviation of the mean estimate.
The top half of the table in Output 95.4.4 shows 95% two-sided confidence limits for the means for the same criteria addressed in the table in Output 95.4.3.

**Output 95.4.4** Confidence Limits

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Treatment</th>
<th>Period</th>
<th>Method</th>
<th>Mean</th>
<th>95% CL Mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>for</td>
<td>1</td>
<td></td>
<td>337.1</td>
<td>287.4</td>
</tr>
<tr>
<td>2</td>
<td>for</td>
<td>2</td>
<td></td>
<td>345.8</td>
<td>271.4</td>
</tr>
<tr>
<td>2</td>
<td>sal</td>
<td>1</td>
<td></td>
<td>283.3</td>
<td>172.7</td>
</tr>
<tr>
<td>1</td>
<td>sal</td>
<td>2</td>
<td></td>
<td>306.4</td>
<td>246.6</td>
</tr>
<tr>
<td>1</td>
<td>Diff (1-2)</td>
<td></td>
<td>Pooled</td>
<td>30.7143</td>
<td>0.2238</td>
</tr>
<tr>
<td>2</td>
<td>Diff (1-2)</td>
<td></td>
<td>Pooled</td>
<td>62.5000</td>
<td>15.5972</td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>46.6071</td>
<td>22.8881</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>-15.8929</td>
<td>-39.6119</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>-15.8929</td>
<td>-40.8415</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Treatment</th>
<th>Period</th>
<th>Method</th>
<th>Std Dev</th>
<th>95% CL Std Dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>for</td>
<td>1</td>
<td></td>
<td>53.7631</td>
<td>34.6446</td>
</tr>
<tr>
<td>2</td>
<td>for</td>
<td>2</td>
<td></td>
<td>70.8814</td>
<td>44.2447</td>
</tr>
<tr>
<td>2</td>
<td>sal</td>
<td>1</td>
<td></td>
<td>105.4</td>
<td>65.7841</td>
</tr>
<tr>
<td>1</td>
<td>sal</td>
<td>2</td>
<td></td>
<td>64.7247</td>
<td>41.7082</td>
</tr>
<tr>
<td>1</td>
<td>Diff (1-2)</td>
<td></td>
<td>Pooled</td>
<td>32.9682</td>
<td>21.2445</td>
</tr>
<tr>
<td>2</td>
<td>Diff (1-2)</td>
<td></td>
<td>Pooled</td>
<td>44.6934</td>
<td>27.8980</td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>19.3702</td>
<td>13.7217</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>Diff (1-2)</td>
<td>Satterthwaite</td>
<td>19.3702</td>
<td>13.7217</td>
<td></td>
</tr>
</tbody>
</table>

For the mean differences, both pooled (assuming equal variances for both treatment sequences) and Satterthwaite (assuming unequal variances) intervals are shown. For example, the pooled confidence limits for the overall treatment mean difference (for sal) assuming equal variances are 22.8881 and 70.3262.

The bottom half of Output 95.4.4 shows 95% equal-tailed confidence limits for the standard deviations within each cell and for the treatment difference within each sequence. It also shows confidence limits for the pooled common standard deviation assuming equal variances. Note that the pooled standard deviation of 19.3702 and associated confidence limits 13.7217 and 32.8882 apply to both difference tests (treatment and period), since each of those tests involves the same pooled standard deviation.

Output 95.4.5 shows the results of t tests of treatment and period differences.

**Output 95.4.5** t Tests

| Treatment    | Period | Method     | Variances | DF | t Value | Pr > |t| |
|--------------|--------|------------|-----------|----|---------|------|---|
| Diff (1-2)   |        | Pooled     | Equal     | 11 | 4.32    | 0.0012 |
| Diff (1-2)   |        | Satterthwaite | Unequal  | 9.1017 | 4.22    | 0.0022 |
| Diff (1-2)   |        | Pooled     | Equal     | 11 | -1.47   | 0.1683 |
| Diff (1-2)   |        | Satterthwaite | Unequal  | 9.1017 | -1.44   | 0.1838 |
Both pooled and Satterthwaite versions of the test of treatment difference are highly significant ($p = 0.0012$ and $p = 0.0022$), and both versions of the test of period difference are insignificant ($p = 0.1683$ and $p = 0.1838$).

The folded $F$ test of equal variances in each treatment sequence is shown in Output 95.4.6.

**Output 95.4.6 Equality of Variances Test**

<table>
<thead>
<tr>
<th>Method</th>
<th>Num DF</th>
<th>Den DF</th>
<th>F Value</th>
<th>Pr &gt; F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Folded F</td>
<td>5</td>
<td>6</td>
<td>1.84</td>
<td>0.4797</td>
</tr>
</tbody>
</table>

The insignificant result ($p = 0.48$) implies a lack of evidence for unequal variances. However, it does not demonstrate equal variances, and it is not very robust to deviations from normality.

Output 95.4.7 shows the distribution of the response variables PEF1 and PEF2 within each of the four cells (combinations of two treatments and two periods) of the AB/BA crossover design, in terms of histograms and normal and kernel density estimates.
Output 95.4.7 Comparative Histograms

The distributions for the first treatment sequence (for/sal) appear to be somewhat symmetric, and the distributions for the sal/for sequence appear to be skewed to the left.

Output 95.4.8 shows a similar distributional summary but in terms of box plots.
Output 95.4.8 Comparative Box Plots

The relative locations of means and medians in each box plot corroborate the fact that the distributions for the sal/for sequence are skewed to the left. The distributions for the for/sal sequence appear to be skewed slightly to the right. The box plot for the salbutamol treatment in the first period shows an outlier (the circle on the far left side of the plot).

The treatment agreement plot in Output 95.4.9 reveals that only a single observation has a higher peak expiratory flow for salbutamol.
The mean for the sal/for treatment sequence is farther from the diagonal equivalence line, revealing that the treatment difference is more pronounced for the 6 observations in the sal/for sequence than for the 7 observations in the for/sal sequence. This fact is also seen numerically in Output 95.4.3 and Output 95.4.4, which show within-sequence treatment differences of 30.7 for for/sal and 62.5 for sal/for.

The profiles over treatment plot in Output 95.4.10 is a different view of the same information contained in Output 95.4.9, plotting the profiles from formoterol to salbutamol treatments. The lone observation for which the peak expiratory flow is higher for salbutamol appears as the only line with negative slope.
The Q-Q plots in Output 95.4.11 assess normality assumption within each of the four cells of the design.
The two Q-Q plots for the sal/for sequence (lower left and upper right) suggest some possible normality violations in the tails, but the sample size is too small to make any strong conclusions. You could use the UNIVARIATE procedure with the NORMAL option to numerically check the normality assumptions.

Finally, Output 95.4.12 shows both pooled and Satterthwaite two-sided 95% confidence intervals for the treatment difference.
Chapter 95: The TTEST Procedure

Output 95.4.12  Confidence Intervals for Treatment Difference

The pooled interval is slightly smaller than the Satterthwaite interval. (This is not always the case.)

Example 95.5: Equivalence Testing with Lognormal Data

Wellek (2003, p. 212) discusses an average bioequivalence study comparing the AUC (area under serum-concentration curve) measurements for two different drugs, denoted “Test” and “Reference,” over a period of 20 hours. This example looks at a portion of Wellek’s data, conducting an equivalence analysis with a paired design that uses AUC values on the original scale (assumed to be lognormally distributed). Each subject in the study received the Test drug upon one visit and then the Reference drug upon a later visit, sufficiently spaced so that no carryover effects would occur.

The goal is to test whether the geometric mean AUC ratio between Test and Reference is between 0.8 and 1.25, corresponding to the traditional FDA (80%, 125%) equivalence criterion. See the section “Arithmetic and Geometric Means” on page 8016 for a discussion of the use of geometric means for lognormal data.

The following SAS statements generate the data set:

```sas
data auc;
  input TestAUC RefAUC @@;
datalines;
103.4 90.11 59.92 77.71 68.17 77.71 94.54 97.51
69.48 58.21 72.17 101.3 74.37 79.84 84.44 96.06
96.74 89.30 94.26 97.22 48.52 61.62 95.68 85.80
; run;
```

You can display the data by using the following statements, which produce Output 95.5.1:

```sas
proc print data=auc;
run;
```
Output 95.5.1  AUC Data for Test and Reference Drugs

<table>
<thead>
<tr>
<th>Obs</th>
<th>Test AUC</th>
<th>RefAUC</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>103.40</td>
<td>90.11</td>
</tr>
<tr>
<td>2</td>
<td>59.92</td>
<td>77.71</td>
</tr>
<tr>
<td>3</td>
<td>68.17</td>
<td>77.71</td>
</tr>
<tr>
<td>4</td>
<td>94.54</td>
<td>97.51</td>
</tr>
<tr>
<td>5</td>
<td>69.48</td>
<td>58.21</td>
</tr>
<tr>
<td>6</td>
<td>72.17</td>
<td>101.30</td>
</tr>
<tr>
<td>7</td>
<td>74.37</td>
<td>79.84</td>
</tr>
<tr>
<td>8</td>
<td>84.44</td>
<td>96.06</td>
</tr>
<tr>
<td>9</td>
<td>96.74</td>
<td>89.30</td>
</tr>
<tr>
<td>10</td>
<td>94.26</td>
<td>97.22</td>
</tr>
<tr>
<td>11</td>
<td>48.52</td>
<td>61.62</td>
</tr>
<tr>
<td>12</td>
<td>95.68</td>
<td>85.80</td>
</tr>
</tbody>
</table>

The TestAUC and RefAUC variables represent the AUC measurements for each subject under the Test and Reference drugs, respectively. Use the following SAS statements to perform the equivalence analysis:

```sas
ods graphics on;
proc ttest data=auc dist=lognormal tost(0.8, 1.25);
   paired TestAUC*RefAUC;
run;
ods graphics off;
```

The DIST=LOGNORMAL option specifies the lognormal distributional assumption and requests an analysis in terms of geometric mean and coefficient of variation. The TOST option specifies the equivalence bounds 0.8 and 1.25.

Output 95.5.2 shows basic summary statistics for the ratio of TestAUC to RefAUC.

Output 95.5.2  Summary Statistics

<table>
<thead>
<tr>
<th></th>
<th>Geometric Mean</th>
<th>Coefficient of Variation</th>
<th>Minimum</th>
<th>Maximum</th>
</tr>
</thead>
<tbody>
<tr>
<td>12</td>
<td>0.9412</td>
<td>0.1676</td>
<td>0.7124</td>
<td>1.1936</td>
</tr>
</tbody>
</table>

The geometric mean ratio of 0.9412 is the sample mean of the log-transformed data exponentiated to bring it back to the original scale. So the plasma concentration over the 20-hour period is slightly lower for the Test drug than for the Reference drug. The CV of 0.1676 is the ratio of the standard deviation to the (arithmetic) mean.
Output 95.5.3 shows the 100(1 − α)% = 95% confidence limits for the geometric mean ratio (0.8467 and 1.0462) and CV (0.1183 and 0.2884).

**Output 95.5.3 Confidence Limits**

<table>
<thead>
<tr>
<th>Geometric Mean</th>
<th>95% CL Mean</th>
<th>Coefficient of Variation</th>
<th>95% CL CV</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.9412</td>
<td>0.8467</td>
<td>1.0462</td>
<td>0.1676</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>0.1183</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>0.2884</td>
</tr>
</tbody>
</table>

Output 95.5.4 shows the 100(1 − 2α)% = 90% confidence limits for the geometric mean ratio, 0.8634 and 1.0260.

**Output 95.5.4 Equivalence Limits**

<table>
<thead>
<tr>
<th>Geometric Mean</th>
<th>Lower Bound</th>
<th>90% CL Mean</th>
<th>Upper Bound Assessment</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.9412</td>
<td>0.8 &lt;</td>
<td>0.8634</td>
<td>1.0260 &lt; 1.25 Equivalent</td>
</tr>
</tbody>
</table>

The assessment of “Equivalent” reflects the fact that these limits are contained within the equivalence bounds 0.8 and 1.25. This result occurs if and only if the p-value of the test is less than the α value specified in the ALPHA= option in the PROC TTEST statement, and it is the reason that 100(1 − 2α)% confidence limits are shown in addition to the usual 100(1 − α)% limits.

Output 95.5.5 shows the p-values for the two one-sided tests against the upper and lower equivalence bounds.

**Output 95.5.5 TOST Equivalence Test**

<table>
<thead>
<tr>
<th>Test</th>
<th>Null</th>
<th>DF</th>
<th>t Value</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Upper</td>
<td>0.8</td>
<td>11</td>
<td>3.38</td>
<td>0.0031</td>
</tr>
<tr>
<td>Lower</td>
<td>1.25</td>
<td>11</td>
<td>-5.90</td>
<td>&lt;.0001</td>
</tr>
<tr>
<td>Overall</td>
<td></td>
<td></td>
<td></td>
<td>0.0031</td>
</tr>
</tbody>
</table>

The overall p-value of 0.0031, the larger of the two one-sided p-values, indicates significant evidence of equivalence between the Test and Reference drugs.

The summary panel in Output 95.5.6 shows a histogram, kernel density, box plot, and 100(1 − 2α)% = 90% confidence interval of the Test-to-Reference ratio of AUC, along with the equivalence bounds.
The confidence interval is closer to the lower equivalence bound than the upper bound and contained entirely within the bounds.

The agreement plot in Output 95.5.7 reveals that the only four subjects with higher AUC for the Test drug are at the far lower or far upper end of the AUC distribution. This might merit further investigation.
The profiles plot in Output 95.5.8 is a different view of the same information contained in Output 95.5.7, plotting the AUC from Test to Reference drug.


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