BIOSTATISTICS FINAL EXAM

Imports the dataset to SAS.

a. Descriptive stat

```
proc means
  data = bone;
  class group;
  var t;
run;
```

The MEANS Procedure								
Analysis Variable : T T								
Group	N Obs	N	Mean	Std Dev	Minimum	Maximum		
ALL	9	9	162.0000000	147.4364270	1.0000000	466.0000000		
AML-High Risk	5	5	2319.00	215.9212820	2133.00	2640.00		
AML-Low Risk	9	9	240.6666667	251.5173950	10.0000000	704.0000000		

INTERPRETATION:

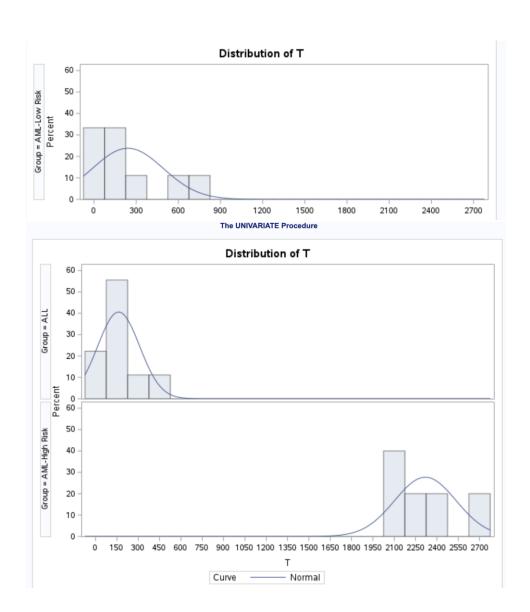
ALL: There are 9 observations, with a mean disease-free survival time of 162. The values range from 1 to 466, with a standard deviation of approximately 147.44.

AML-High Risk: In this group, there are 5 observations. The mean disease-free survival time' value is significantly higher (2319), with values ranging from 2133 to 2640. The standard deviation is about 215.92.

AML-Low Risk: This group has 9 observations. The mean disease-free survival time' value is 240.67, with a wider range of values (from 10 to 704) compared to the other groups. The standard deviation is approximately 251.52.

(b) Check the normality assumption. proc univariate data= bone normal; class group;

var T;
histogram T/normal;
run;



INTERPRETATION:

Based on the graphs, they do not follow a normal distribution.

(c) Is disease-free survival time different among the three groups? State the null and alternative hypotheses. Which test do you use? Choose the most appropriate test and justify it. Conduct the test at the 0.05 level of significance. If needed, conduct post-hoc tests. What do you conclude?

Null Hypothesis: There is no difference in the disease-free survival time between the 3 groups: ALL (acute lymphoblastic leukemia), AML-Low Risk (acute myelocytic leukemia, low risk), and AML-High Risk.

Alternative hypothesis: There is difference in disease-free survival time between at least one of the 3 groups: ALL (acute lymphoblastic leukemia), AML-Low Risk (acute myelocytic leukemia, low risk), and AML-High Risk.

We use anova test in this case. ANOVA is used when comparing the means of three or more groups for a single variable. It helps determine if there's a statistically significant difference in disease-free survival times among the groups (ALL, AML-Low Risk, AML-High Risk). ANOVA allows for this simultaneous comparison while considering the assumption of equal variances among the groups.

proc anova data=bone;
class group;
model T =group;
means group/hovtest;
run;

				Depender	nt Vari	able: T T				
Sour	ource DF		FS	Sum of Squares		Mean Square F		F۷	/alue	Pr > F
Mode	el		2	1757593	38.87	8787969.43		20	2.84	<.0001
Error	•	2	0	86647	6.00	43323.80				
Corrected Total		al 2	2	18442414.8						
R-		R-Sq	uare	Coeff Va	r Ro	oot MSE	T Me	ean		
		0.95	3017	31.4561	1 2	08.1437	661.69	957		
	Source	DF	A	nova SS	Mear	n Square	F Val	lue	Pr>	F

/* With a p value of <.0001 and an alpha of 0.05, We reject the null hypothesis. This suggests that there's evidence that at least one of the groups (ALL, AML-Low Risk, AML-High Risk) differs in disease-free survival time. */

/* posthoc */

The results of the ANOVA test indicated that at least one of the groups (ALL, AML-Low Risk, and AML-High Risk) had a different disease-free survival time. Similar to Tukey's HSD, the post hoc test is performed to identify which particular groups show these significant differences.

This test helps to clarify which pairs of groups differ in disease-free survival time, enhancing our understanding beyond the general difference highlighted by ANOVA.

PROC GLM DATA=bone; CLASS Group; MODEL T = Group; LSMEANS Group / DIFF ADJUST=TUKEY; RUN;

Group		T LSMEAN		LSMEAN Number			
ALL		162.00000				1	
AML-High Risk		2319.00000			2		
AML	Low	Risk	240	240.66667		3	
		•				ct Group	
		> t for	· H0: L	eans for SMean(i	i)=LS	Mean(j)	
		> t for	· H0: L	.SMean(i	i)=LS	Mean(j)	
	Pr	> t for	· H0: L	.SMean(i	ble: 1	Mean(j)	
	Pr:	> t for	· H0: L	SMean(i	ble: 1	Mean(j)	

Interpretation:

Comparing 'ALL' with 'AML-High Risk' and 'AML-Low Risk' showed significant differences in disease-free survival time (p < 0.0001).

There was also a significant difference between 'AML-High Risk' and 'AML-Low Risk' (p < 0.0001).

However, there doesn't appear to be a significant difference between 'ALL' and 'AML-Low Risk' (p = 0.7063), based on these pairwise comparisons.

data clinic;

input Treatment \$ Response \$ Count; datalines;

Placebo Satisfactory 2

Placebo Unsatisfactory 3

Test Satisfactory 4
Test Unsatisfactory 1
;
proc print data= clinic;
run;

a. Create a 2x2 table and print it using SAS.

b. Is there any association between treatment and response? State the null and alternative hypotheses. Which test do you use? Choose the most appropriate test and justify it. Conduct the test at the 0.05 level of significance. What do you conclude?

Null hypothesis: There is no association between treatment and response.

Alternative hypothesis: Not Null hypothesis

proc freq data= clinic;
table Treatment*response/chisq;
Weight count;
run;

Frequency	Table of Treatment by Response						
Percent Row Pct		Response					
Col Pct	Treatment	Satisfac	Unsatisf	Total			
	Placebo	2	3	5			
		20.00	30.00	50.00			
		40.00	60.00				
		33.33	75.00				
	Test	4	1	5			
		40.00	10.00	50.00			
		80.00	20.00				
		66.67	25.00				
	Total	6	4	10			
		60.00	40.00	100.00			

a. This is 2x2 table.

b.

Statistics for Table of Treatment by Response

Statistic	DF	Value	Prob
Chi-Square	1	1.6667	0.1967
Likelihood Ratio Chi-Square	1	1.7261	0.1889
Continuity Adj. Chi-Square	1	0.4167	0.5186
Mantel-Haenszel Chi-Square	1	1.5000	0.2207
Phi Coefficient		-0.4082	
Contingency Coefficient		0.3780	
Cramer's V		-0.4082	
WARNING: 100% of the calls be		nacted car	inte loce

WARNING: 100% of the cells have expected counts less than 5. Chi-Square may not be a valid test.

Fisher's Exact Test					
Cell (1,1) Frequency (F)	2				
Left-sided Pr <= F	0.2619				
Right-sided Pr >= F	0.9762				
Table Probability (P)	0.2381				
Two-sided Pr <= P	0.5238				

Sample Size = 10

INTERPRETATION:

The Chi-square test is used to check the relationship between treatment and response. It is used to assess the association between treatment and response because both variables are categorical. This test determines if there's a significant relationship between categorical variables.

Since more than 20% of the cells had expected counts below 5, indicating low counts, the Chi-square test might be unreliable. Therefore, Fisher's exact test is preferred. With a p-value of 0.5238 at a significance level of 0.05, we fail to reject the null hypothesis. Therefore, based on this analysis, there isn't enough evidence to claim an association between the treatment administered and the observed response.

```
dbms = xlsx
out = hypo;
```

run;

a. Compute descriptive statistics for TWA_HR (time-weighted average of heart rate during surgery), age, gender (female), sleeptime (time spent at Sao2<90%), AHI (Apenea/Hyponea index).

```
proc means
  data = hypo maxdec=2;
  var TWA_HR age female sleeptime AHI;
run;
```

The MEANS Procedure							
Variable	Label	N	Mean	Std Dev	Minimum	Maximum	
TWA HR	TWA HR	203	76.10	10.16	54.22	109.09	
Age _	Age	281	47.29	11.12	16.40	73.80	
Female	Female	281	0.72	0.45	0.00	1.00	
Sleeptime	Sleeptime	281	18.71	23.73	0.00	99.60	
AHI .	AHI .	278	2.87	1.08	1.00	4.00	

Interpretation:

TWA_HR: The average time-weighted heart rate during surgery was approximately 76 beats per minute, ranging from 54 to 109 beats per minute.

Age: The average age in the dataset was around 47 years, with ages ranging from 16 to 73 years.

Gender (Female): The mean value of 0.72 suggests that, on average, about 72% of the dataset consists of females.

Sleeptime: On average, patients spent approximately 18.71% of their time at a Sao2 level below 90%, with a wide variation from 0% to almost 100%.

AHI: The average Apnea/Hypopnea Index was 2.87, ranging from 0 to 16.40, indicating the frequency of apnea and hypopnea events during sleep.

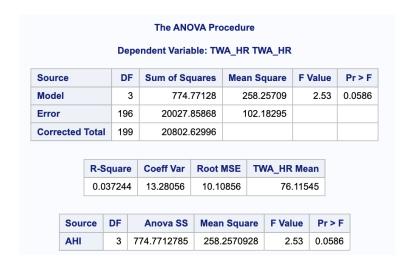
b. Is TWA_HR different among AHI? State null and alternative hypotheses. Which test do you use? Conduct the test at the 0.05 level of significance. Interpret the results.

Null Hypothesis: TTWA_HR is different among AHI

Alternative Hypothesis: Not null

Anova test is used here.

proc anova data=hypo; class AHI; model TWA_HR = AHI; means AHI/hovtest; run;

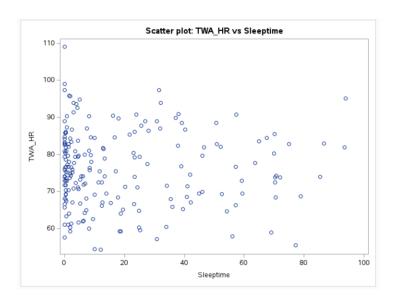


With a p value of 0.0586 and an alpha of 0.05, We do not reject the null hypothesis. There isn't enough evidence to conclude that TTWA_HR significantly differs among different AHI levels.

c.Produce a scatter plot of TWA_HR versus Sleeptime. Produce another scatter plot of TWA versus minimum nocturnal Sao2. Conduct a correlation analysis for these three variables and interpret the results.

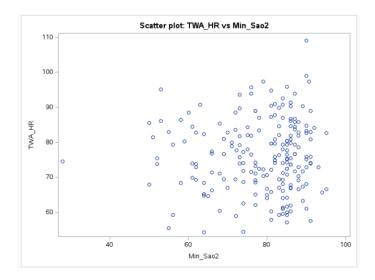
Scatter plot of TWA_HR versus Sleep time:

```
title "Scatter plot: TWA_HR vs Sleeptime";
proc sgplot data= hypo;
scatter y=TWA_HR x=Sleeptime;
run;
title "";
```



Scatter plot of TWA versus minimum nocturnal Sao2:

```
title "Scatter plot: TWA_HR vs Min_Sao2";
proc sgplot data= hypo;
scatter y=TWA_HR x=Min_Sao2;
run;
title "";
```



Correlation analysis for these three variables and interpret the results.

Null Hypothesis: The population correlation coefficient (ρ) is 0 and there is no correlation between TWA HR and AHI.

Alternative Hypothesis: Not null hypothesis.

Null Hypothesis: The population correlation coefficient (ρ) is 0 and there is no correlation between TWA_HR and Min_Sao2.

Alternative Hypothesis: Not null hypothesis.

Null Hypothesis: The population correlation coefficient (ρ) is 0 and there is no correlation between AHI and Min Sao2.

Alternative Hypothesis: Not null hypothesis.

proc corr data=hypo plots=matrix(histogram);
var TWA_HR AHI Min_Sao2;
run;

Pearson Correlation Coefficients Prob > r under H0: Rho=0 Number of Observations							
	TWA_HR	АНІ	Min_Sao2				
TWA_HR	1.00000	-0.07266	0.03874				
TWA_HR		0.3066	0.5832				
_	203	200	203				
AHI	-0.07266	1.00000	-0.50245				
AHI	0.3066		<.0001				
	200	278	278				
Min_Sao2	0.03874	-0.50245	1.00000				
Min_Sao2	0.5832	<.0001					
	203	278	281				

With a p value of 0.3066 and alpha of 0.05, we fail to reject the null hypothesis. So, there's insufficient evidence to claim a correlation between TWA_HR and AHI.

With a p value of <0.0001 and alpha of 0.05, we reject the null hypothesis. So, there is no correlation AHI and Min_Sao2.

With a p value of 0.5832 and alpha of 0.05, we fail to reject the null hypothesis. So, t here's insufficient evidence to claim a correlation between TWA_HR and Min_Sao2.