GER1000 Quantitative Reasoning Help Sheet for Final Examination

Disclaimer: I hereby affirm that any errors found in this sheet is solely due to my own human error and are not committed on purpose in order to "snake".

Joshua of House Ursaia

Controlled Experiments	
Experiment that compares between th controlled group.	e response of a treatment group and a
Randomized Control	Double-Blinding
Experiment that assigns subjects into the control and treatment group randomly. ■ Large number of subjects → likely that the two groups are similar in all aspect.	Experiment where the subjects do not know whether they are in the treatment or control groups; neither do those who evaluate the responses. Guards against bias either in the responses or evaluations. Minimizes confounding

Observational :	Studies
Experiment wh	ere the investigators do not assign the subjects into the
treatment or co	ontrol group.
Consequence	Effects of treatment may be confounded with the effects of factors that got the subjects into the treatment and control groups in the first place.

Rate		
$rate(A \mid B)$	Rate of A among people with B	
$rate(A \mid not B)$	Rate of A among people without B	
Basic Rule on Rates		
$rate(B)$ is always between $rate(B \mid A)$ and $rate(B \mid not A)$.		
$rate(A) \rightarrow 100\%$	$rate(B) \rightarrow rate(B \mid A)$	
$rate(B \mid A)$	$rate(B) = rate(B \mid A)$	
$= rate(B \mid not A)$		
rate(A) = 50%	$rate(B)$ is exactly halfway between $rate(B \mid A)$	
	and $rate(B \mid not A)$.	

Yule-Simpson Paradox

Suppose that the population consists of several subgroups, and in each subgroup, $rate(B \mid A) > rate(B \mid not A)$.

When the **subgroups are combined**, it may happen that $rate(B \mid A) \le rate(B \mid not A)$.

Association	
A and B are positively associated	rate(A B) > rate(A not B)
	rate(B A) > rate(B not A)
A and B are negatively associated	rate(A B) < rate(A not B)
	rate(B A) < rate(B not A)
No Association	rate(A B) = rate(A not B)
	rate(B A) = rate(B not A)

Confounders	
A confounder is a th	hird variable associated with both exposure and disease.
Methods to control for confounding factors	
Slicing Method	Compare smaller groups which are relatively
	homogeneous with respect to the confounding factor.
Regression	Explained in detail later.

Types of Relationships	
Deterministic	The value of the dependent variable can be determined from the value of the independent variable.
Statistical	The average pattern of one variable can be described with the value of the other variable.

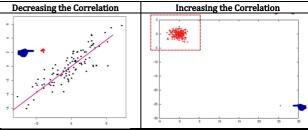
Linear Regre	ssion
Linear Regre	ession is used to investigate association between two continuous
variables.	
Linear	The line of best fit to data, where the sum of squares of the
Regression	distance of each data point to the line is minimum.
Line	
Linear Regre	ession Coefficient, $r (-1 \le r \le 1)$
Sign	$r > 0 \rightarrow$ Positive association
	$r < 0 \rightarrow$ Negative association
	$r = 0 \rightarrow$ No linear association (may still have non-linear
	association)
Magnitude	The closer the value to 1 $or - 1$, the stronger the linear
	association
Not	1. Interchange of the two variables
affected by	2. Adding a number to all values of a variable
	3. Multiplying a positive number to all values of a variable
Points to tak	e note:
 In gene 	eral, the slope of the linear regression line ≠ the correlation
coeffic	ient.
 A high 	linear correlation coefficient does not necessarily mean that the

relationship is linear.

 Extrapolation: Predicting the value of the dependent variable beyond the observed range of the independent variable is dangerous.

Impact of Outliers on Correlation

It is dangerous to exclude outliers from the analysis without understanding the causes of their occurrence.



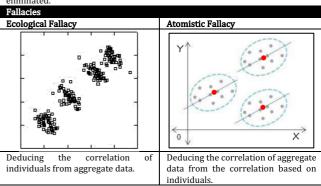
Attenuation Effect

Range restriction in one variable could cause the correlation coefficient obtained to "understate" the strength of association between two variables

Check for an "oval shape" in your scatter diagram. If you see it, you are less likely to have the attenuation effect.

Ecological Correlation

Correlation based on **aggregated data**, such as group average or rates. When the associations for both individuals and aggregates are in the same direction, the ecological correlation will typically "**overstate**" the strength of association in individuals, because the variability among individuals have been eliminated.



Estimate = Parameter + Bias + Random Error	
A numerical fact about a population.	
Depends on the method of sampling.	
Depends on the size of sampling.	

Sampling Frame

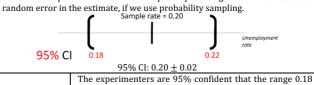
A list of sampling units intended to identify all units of a population.

 Has to cover exactly or bigger than target population and has to be upto-date.

Biases and their Causes		
Selection Bias	Systematic tendency to exclude one kind of person or another from the sample, caused by: Imperfect sampling frame (which excludes certain desired units) Non-probability sampling methods	
Non-Response Bias	Differences between non-respondents and respondents	
Response Bias	Phrasing of the questions, tone, or attitude of the interviewers.	
Other Types	Subjects may have a tendency to understate responses about undesirable social habits (ashamed).	

Confidence Intervals

Confidence Interval is the range of values in which we are reasonably certain our unknown parameter lies in. It is helpful in providing information about the random error in the estimate, if we use probability sampling.



Interpretations

The experimenters are 95% confident that the range 0.18 and 0.22 contains the population parameter.

95% of the researchers who repeat the experiment will have intervals that contain the population parameter.

Link to Random Error For an experiment with a **larger sample size** \rightarrow likely to have **smaller random error** \rightarrow at the same 95% confidence interval, the experiment will have a **smaller range**.

Types of Probability Sampling

Simple Random Sampling

Draw units at random from a sampling frame without replacement, with every unit having an equal chance of being selected.

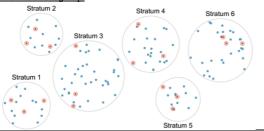


Systematic Sampling

Select a random starting point, r. Include every k^{th} unit after r into the sample.

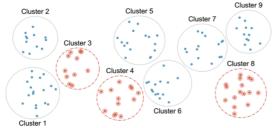
Stratified Sampling

Divide the population into homogeneous subgroups (strata). Take a <u>random sample from each subgroup.</u>



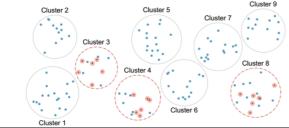
Cluster Sampling

Divide population into naturally occurring subgroups (clusters). Take random sample of clusters.



Multi-Stage Sampling

Sampling is carried out in stages with smaller and smaller sampling units at each stage. A probability sampling plan is implemented at each stage.



Types of Non-Probability Sampling

Volunteer / Self-Selected Sampling

Only those people interested in the study would respond.

→ Hard to talk about the degree of non-response bias.

Convenience / Haphazard Sampling

Using the most convenient group available or deciding on the sample on the spot.

Judgement Sampling

Using human judgement to choose "representative" units.

Quota Sampling

Interviewers are free to select anyone they like until they meet a fixed quota of units to interview for a fixed number for certain categories (eg. sex, age)

Risk and Risk Ratio	
Risk (Disease Male)	Number of Males with Disease
	Number of Males
Risk Ratio of Disease	Risk (Disease Male)
between Males to	Risk (Disease Female)
Females	` ' '

Odds and Odds Ratio	
Odds (Disease)	Risk
among Males	1 - Risk
	Number of Males with Disease
	Number of Males without Disease
Odds Ratio (Disease)	Odds (Disease) among Females
between Females and	Odds (Disease) among Males
Males (Females is the "first" group, Males is the "baseline" group)	You can also use the cross-product ratio , by setting up a 2x2 contingency table: • Event of interest on the first column • The first group (Females) are on the first row.
Interpretation of Odds Ratio	
OR = 1	No difference in disease risk in the two groups, and
	RR = 1
OR < 1	Higher risk in the first group, and RR > 1
OR > 1	Lower risk in the first group, and RR < 1

0K < 1	riighei risk iii the first group, and kk > 1
OR > 1	Lower risk in the first group, and RR < 1
Types of Observational Studies	
In observational studies, there is Association between the exposure and	
disease if Risk Ratio $\neq 1$ or Odds Ratio $\neq 1$.	

ease if $Risk Rullo \neq 1$	or Odds Ratto 7 1.	
	Can estimate Population Odds Ratio?	Can estimate Population Risk Ratio?
hort Study	Yes	Yes

<u>Starts from the exposure</u> by investigating 100 smokers and 100 nonsmokers

 <u>Looks into the future</u> and observes how many people from each group eventually have cancer.

ase-Control Study

• <u>Starts from the outcome</u>, or the disease statistics by investigating 100 people with cancer and 100 people without cancer.

<u>Looks into the background</u> and observes how many people from each group were smokers.

Cross Sectional Study

•	Starts from a population	sample by investigating	ng 1000 random people.

Looks at their current disease and exposure status.

Probability Rules			
Complement Rule	P(Complement) = 1 - P(A)		
"At Least One"	$P(At \ least \ one) = 1 - P(None)$		
Addition Rule	P(A or B) = P(A) + P(B) - P(A and B)		
Mutually Exclusive Events	P(A or B) = P(A) + P(B)		
Multiplication Rule	$P(A \text{ and } B) = P(A) \times P(B \mid A)$		
Independent Events	$P(A \text{ and } B) = P(A) \times P(B)$		
Average Value of an Action			
Assumption: Outcome A and B are mutually exclusive.			
Value of Action = $V(A) \times P(A) + V(B) \times P(B)$			

Hypothesis Testing		
p-value	The sum of probabilities of events similar to and more	
	extreme than our observation.	
Null Hypothesis	Corresponds to the idea that an observation is due to	
	chance.	
Alternative	Corresponds to the idea that an observation is due to	
Hypothesis	chance.	
If the p-value is very small → our observed result of the experiment is likely		
not due to chance → reject the Null Hypothesis.		

Testing for Rare Events		
Base Rate	P(Disease)	
Sensitivity	P(Positive Disease)	
Specificity	P(Negative No Disease)	