

## 0.1 STAT220: BASIC STATISTICS, WIN 2007

Instructor: Prof. Elizabeth Thompson  
eathomp@u.washington.edu

Teaching assistants: Linqiu Wu, Sections AA, AB

Lee Barr, Sections AC, AD

Le Bao, Sections AE, AF

- The web page:

[www.stat.washington.edu/thompson/S220\\_07](http://www.stat.washington.edu/thompson/S220_07)

Or, link from Statistics Dept home page is now working.

For course requirements,

book info (FPP), homework info,  
exams and quizzes info,  
and lecture notes.

In particular, check the web schedule

– it has links to the other things.

Also, office hours of TAs and Instructor (TBA).

## 0.2 WHAT IS STATISTICS?

- Quantitative facts, numerical descriptions (data)
- Set of tools for the collection and analysis of data
- ... in order to make decisions or draw conclusions from the data

## 0.3 WHERE DO WE SEE DATA ANALYSES ?

- News reports, Weather forecastes
- School records, grades, course evaluations
- Consumer reports, Election polls
- Environmental standards, air pollution, endangered species
- Medical and dental records, diagnostic procedures.
- Stock market, business plans, marketing surveys.
- ... and many more

#### 0.4 WHAT DOES DOING STATISTICS INCLUDE?

Blank page for your notes

- Study design and data collection:  
Experiments, Studies and Surveys  
FPP Chapters 1, 2, 19
- Data description and exploration  
Graphical and numerical summaries.  
FPP Chapters 3,4
- Modeling Data : for example, the normal curve  
FPP Chapter 5
- Forecasting and prediction  
The relationship between 2 or more variables  
Correlation and association: FPP Chapters 7,8,9  
Regression: FPP Chapters 10,11, 12
- Understanding variation and randomness  
Chance and randomness: FPP Chapters 16,17,18  
Accuracy and measurement error: FPP 20,23,24
- Drawing inferences and making decisions  
Estimates and confidence intervals: FPP Ch. 21  
Testing hypotheses: FPP Chapter 26, 27

## 1.1 COLLECTION OF DATA

There are three basic study designs

### 1. Controlled experiments (FPP Chapter 1)

Investigator gives treatment to subjects in the treatment group.

### 2. Observational studies (FPP Chapter 2)

Investigator does not control who is in the treatment group.

### 3. Sample surveys (FPP Chapter 19)

A type of observational study. We study a sample of individuals from a population

- Subjects: Study units, Experimental units
- Population: The set of individuals of interest
- Sample: Chosen subset of the population
- Variable: Characteristic or property of a subject

Blank page for your notes

## 1.2 EXAMPLE: SALK VACCINE TRIALS

Blank page for your notes

- From 1916, polio killed hundreds of thousands US children
- In 1954, Jonas Salk's vaccine seemed promising
- Need comparison: treatment and controls
  - Vaccinated, about 500,000 children
  - Unvaccinated, about 1,000,000 children
  - Refused vaccination, about 500,000 children
- Randomized controlled trial:
  - Investigator decides who is to be vaccinated/not.
  - Use random assignment to treatment or control group— those whose parents refused vaccination are not good controls (confounding factors)
- Use of placebo, avoids placebo effect
  - Subjects should not know whether they are treatment or control —use of saline solution for children in control group.
- Double blind assessment
  - Neither subjects nor diagnosing physicians know who is treatment and who is control.

### 1.3 TREATMENTS, RESPONSE, and FACTORS

Blank page for your notes

- Treatment or control is applied to the experimental unit (or subject).
- The response is the outcome: the data we analyze.
- The treatment may involve several factors. For example, cancer treatment may involve surgery, radiation therapy and chemotherapy.
  - Surgery: yes or no (2 levels)
  - Radiation treatment: high dose, low dose, or none (3 levels)
  - Chemotherapy: protocol-1, protocol-2, or none (3 levels)
- Need to try (all?) combinations to assess treatments
  - Some combinations may not be feasible/ethical
  - Issues of time, cost (numbers of subjects)
- If do do all combinations, this is a complete factorial design

#### 1.4 RANDOMIZED CONTROLLED TRIALS

Blank page for your notes

- Investigator assigns subject to treatment/control.  
This avoids confounding factors How should he/she assign?
- We want treatment group to be similar to control group  
but any directed attempt to make them similar may lead to bias.
- Only random assignment of eligible subjects is safe  
then we can assess results objectively  
Subjective confounding factors will not cause bias.
- What about obvious confounding factors.  
For example, gender, in study of hormone drug reactions.  
Randomization will take care of it, on average.
- But also we can stratify the subjects by gender  
—Essentially do two experiments.
- Randomize or stratify ? – BOTH  
Within each stratum (gender), randomize.  
Issues of cost?

## 1.5 OBSERVATIONAL STUDIES

Blank page for your notes

- The subjects assign themselves
  - Study of cancer: smokers and non-smokers
  - Study of income at 40: choice of major at UW
  - Study of drug: who keeps to protocol?
- WYSIWYG: Investigators just watch the outcomes!  
Association is not causation.
- How did subjects come to be in treatment/control?
  - identify likely confounding factors.
- Control for known confounding factors – stratify!  
that is, analyze in smaller more homogeneous groups
- Example: Bias in graduate admissions at Berkeley (1973)??
  - FPP 17-19
  - Choice of major is confounded with gender.
- This is an example of Simpson's paradox.
- Other considerations:
  - Can we observe? outcomes, behaviors, but not beliefs or attitudes (contrast with survey).
  - Cost? – time is money.
  - Observer presence may affect outcome?

## 1.6 SELECTING A SAMPLE

- We have a population of interest.  
We want to know some parameters of the population.  
But we cannot look at the whole population.
- We select a sample (subset) from the population  
We compute a statistic based on the sample, to estimate the parameter.
- We can choose individuals randomly, or judge what factors may be important.
- If we choose a random sample, we do so carefully to avoid systematic bias
- Unintended selection bias. Non-response bias.
- Large samples do not protect against bias.
- If we first consider some important factors, then sample randomly within categories: OK!  
This is stratified random sampling
- If we use these factors to select the sample  
this is quota sampling  
and is subject to unintended biases,  
due to confounding factors associated with selection factors