

<b>Logistics</b>	1
<b>Module 1 - Descriptive</b>	2
History of epi	2
Definition of epi	2
Core functions	2
Morbidity and mortality	2
Outbreak investigations and disease surveillance	3
Measures of validity and reliability	3
<b>Module 2 - Study Designs</b>	4
Types of populations	4
Validity in study design	4
Types of study design	4
<b>Module 3 - Analysis and Inference</b>	4
Measures	4
Ratios	5
Differences	5
Necessary versus sufficient	5
Epidemiological guidelines for suggesting causation	5
Chance	5
Bias	5
Confounding	5
Effect modification	6
<b>Module 4 - Applications of Epi</b>	6

## Logistics

12/7 office hours 12-1PM

12/12 office hours 12-1PM

Look over:

- HW assignments
- Midterm exams
- Discussion slides

N-Z Hodson 210 9AM-12PM 12/13

1 double sided 3x5 index card

50MC, 3-4 guest lecture questions

### Extra Tips

- Make sure to check whether a question is asking for survival or death
- For cumulative incidence, check for loss to followup

## Module 1 - Descriptive

### History of epi

- John Snow (cholera)
- Semmelweis (handwashing after autopsy)
- Jenner (smallpox vaccine)
- Graunt (life table)
- Lind (scurvy, controlled trial)
- Typhoid Mary [Q]

Commented [1]: Fa18 Q23

### Definition of epi

- study of distribution and determinants of health-related states or events in specified populations and the application of this knowledge to the control of health problems

### Core functions

- Assessment, policy development, assurance
- Prevention - Primary, Secondary, Tertiary [Q]

Commented [2]: HW1 Q1-3  
Sp18 Q1  
Fa18 Q1

### Morbidity and mortality

- Prevalence - burden of disease =  $\text{pop}(\text{cases}) / \text{pop}(\text{total})$  [Q]
- Incidence - risk of disease
  - Cumulative = new cases/total pop at risk [Q]
  - Incidence rate = new cases/total pop-time at risk [Q]
  - Assumes complete followup
- Using prevalence versus incidence [Q]
- Mortality rate
  - All-cause =  $P(\text{deaths}(\text{total}) | \text{pop}_{\text{midyear}})$  [Q]
  - Cause-specific =  $P(\text{deaths}(\text{disease}) | \text{pop}_{\text{midyear}})$  [Q]
- Case fatality rate - severity of disease =  $P(\text{deaths} | \text{disease})$  [Q]
- Proportionate mortality - proportion of deaths due to disease =  $P(\text{death}(\text{disease}) | \text{death})$  [Q]
- Age-adjusted mortality rate (direct adjustment) =  $\sum_{a \in \text{age}} \text{pop}(\text{standard}, \text{age}) * \frac{\text{\#deaths}_{\text{age}}}{\text{pop}(\text{group}, \text{age})} / \sum_{a \in \text{age}} \text{pop}(\text{standard}, \text{age})$  [Q]

Commented [3]: Sp18 Q16  
Fa18 Q18

Commented [4]: HW1 Q10  
Sp18 Q20  
Fa18 Q16

Commented [5]: HW1 Q9  
Sp18 Q19  
Fa18 Q4,17

Commented [6]: HW1 Q11,18  
Sp18 Q3  
F18 Q20

Commented [7]: HW1 Q7,15

Commented [8]: HW1 Q4  
Sp18 Q18

Commented [9]: HW1 Q8,12  
Sp18 Q4

Commented [10]: Sp18 Q5

Commented [11]: HW1 Q13-14,16  
Sp18 Q9,12  
Fa18 Q5,19

- Standardized mortality ratio (indirect adjustment) =  $\text{number of deaths in subpop} / (\text{mortality rate of standard} * \text{subpop})$  [Q] [Q]
- Observed survival
  - Life table [Q]
  - Kaplan-Meier method [Q]
  - Life table versus Kaplan-Meier [Q]

Commented [12]: HW1 Q17  
Sp18 Q22

Commented [13]: Age adjustment in general  
Fa18 Q25

Commented [14]: Sp18 Q24,25  
Fa18 Q14,15

Commented [15]: HW1 Q22

Commented [16]: HW1 Q24  
Fa18 Q9

Commented [17]: Fa18 Q8

Commented [18]: Sp18 Q17  
F18 Q6

Commented [19]: HW2 Q2  
Sp18 Q21  
Fa18 Q22

#### Outbreak investigations and disease surveillance

- Endemic versus epidemic versus pandemic [Q]
- Herd immunity - resistance of a group of people to a disease based on proportion of people immune to disease, applies to person-to-person transmission [Q]
- Types of outbreaks [Q]
  - Common source
    - Point source - food outbreak
    - Continuous - broken fountain
    - Intermittent - common flu
  - Propagated - HFM
  - Mixed
  - Using epi curve
- Attack rate
- Identifying probable cause of outbreak [Q]
- Incubation period, via cross-tabulation [Q]
- Types of surveillance
  - Sentinel - pick certain sites
  - Syndromic - between symptoms and diagnosis
  - Passive - mandatory reported diseases
  - Active
  - Identify type [Q]
- Sources of data - medical records, emergency departments, 911 calls, or nonclinical data from schools, labs, surveys, etc.

Commented [20]: Sp18 Q6,7  
Fa18 Q11

Commented [21]: HW2 Q1  
Sp18 Q8,15  
Fa18 Q10

Commented [22]: HW2 Q18  
Sp18 Q10  
Fa18 Q13,24

Commented [23]: \*\*\*

#### Measures of validity and reliability

- Validity - expresses degree to which a specified measurement measures what it purports to measure, sensitive and specificity are fixed characteristics of the test, predictive values depend on the prevalence of the disease, often for new tools [Q]
  - Sensitivity =  $P(\text{test positive} | \text{disease})$
  - Specificity =  $P(\text{test negative} | \text{no disease})$  [Q]
  - Net sensitivity simultaneous = inclusion exclusion
  - Net specificity simultaneous = product rule
  - Net sensitivity sequential = product rule
  - Net specificity = inclusion exclusion

Commented [24]: HW2 Q8

Commented [25]: Sp18 Q11,13  
Fa18 Q7

- Simultaneous and sequential testing - simultaneous captures positive for either, so high sensitivity, sequential captures test negative for either, so high specificity [Q]
- PPV =  $P(\text{disease} \mid \text{positive test})$ , increases with prevalence
- NPV =  $P(\text{no disease} \mid \text{negative test})$  [Q]
- NNH
- NNT [Q]
- ITT [Q]
- Reliability - degree to which the results can be replicated for a specified measurement procedure, aka repeatability, precision, comparing one tool to itself
  - intra and inter observer agreement
  - Kappa statistic - won't have to calculate, only interpret [Q]

Commented [26]: HW2 Q3,4,5  
Sp18 Q14  
Fa18 Q2,12

Commented [27]: Sp18 Q2  
Fa18 Q3

Commented [28]: HW2 Q23

Commented [29]: HW2 Q19

Commented [30]: HW2 Q6,7  
Sp18 Q23

## Module 2 - Study Designs

### Types of populations

- Target
- Source
- Study
- [Q]

Commented [31]: HW2 Q22

### Validity in study design

- Internal validity - need to consider methodological problems, bias, confounding, random error, must be established before generalizing to external bias
- External validity

Commented [32]: HW2 Q16

### Types of study design

- Clinical Trial
- Cohort
- Case control
- Cross-sectional
- Ecologic
- Identify type

Commented [33]: HW2 Q11

Commented [34]: HW2 Q10,13,15

Commented [35]: HW2 9, 12, 20, 21, 24, 25

Commented [36]: HW2 Q14

## Module 3 - Analysis and Inference

### Measures

- Calculate and interpret each measure
- Use in appropriate study design

Commented [37]: HW3 Q5,6,7,14,1,19

Commented [38]: HW3 Q9,10

## Ratios

- Relative risk (trial, cohort)
- Prevalence ratio (cross-sectional)
- Odds ratio (case control)
  - Matched and unmatched
- RR is roughly OR when disease is rare

Commented [39]: HW3 Q20,21

Commented [40]: HW3 Q12

## Differences

- Risk difference
- Prevalence difference
- Attributable risk

## Necessary versus sufficient

- Necessary - disease won't develop w/out exposure
- Sufficient - exposure guarantees disease

Commented [41]: HW3 Q4,23

## Epidemiological guidelines for suggesting causation

- Temporality (only one that is necessary)
- Strength of association
- Dose-response gradient
- Consistency/replication
- Coherence (biological plausibility)
- Experiment (cessation of exposure)
- Analogy
- Consideration of alternate explanations
- Specificity of association

Commented [42]: HW3 Q11,13,24,25

## Chance

- p value, sample size, precision of confidence interval determined by sample size, type I, II errors

Commented [43]: HW3 Q17,18,22

## Bias

- systematic/non-random error in design, conduct or analysis of study
- Selection bias - difference in characteristics between those who take part in a study versus those that do not
- Information bias - flaw in data collection or measurement that results in different quality/accuracy between comparison groups
- Identify type

Commented [44]: HW3 Q2

Commented [45]: HW3 Q3,16

Commented [46]: HW2 Q17

## Confounding

- Associated with exposure

- Cause of outcome
- Not in the causal pathway
- Control in study design
  - Restriction on confounding variable
  - Matching in a case-control study
  - Randomization in a clinical trial
- Control in the data analysis
  - Stratification
  - Adjustment

Commented [47]: HW3 Q8

Commented [48]: HW3 Q1

#### Effect modification

- Make a list of potential confounders and effect modifiers
- Calculate crude measure of association for exposure and outcome of interest, calculate measures for stratified

#### Module 4 - Applications of Epi