

---

CMED 305

---

# Cross-Sectional Studies

---

---

**Nurah Alamro, MD. MPH. DrPH.**

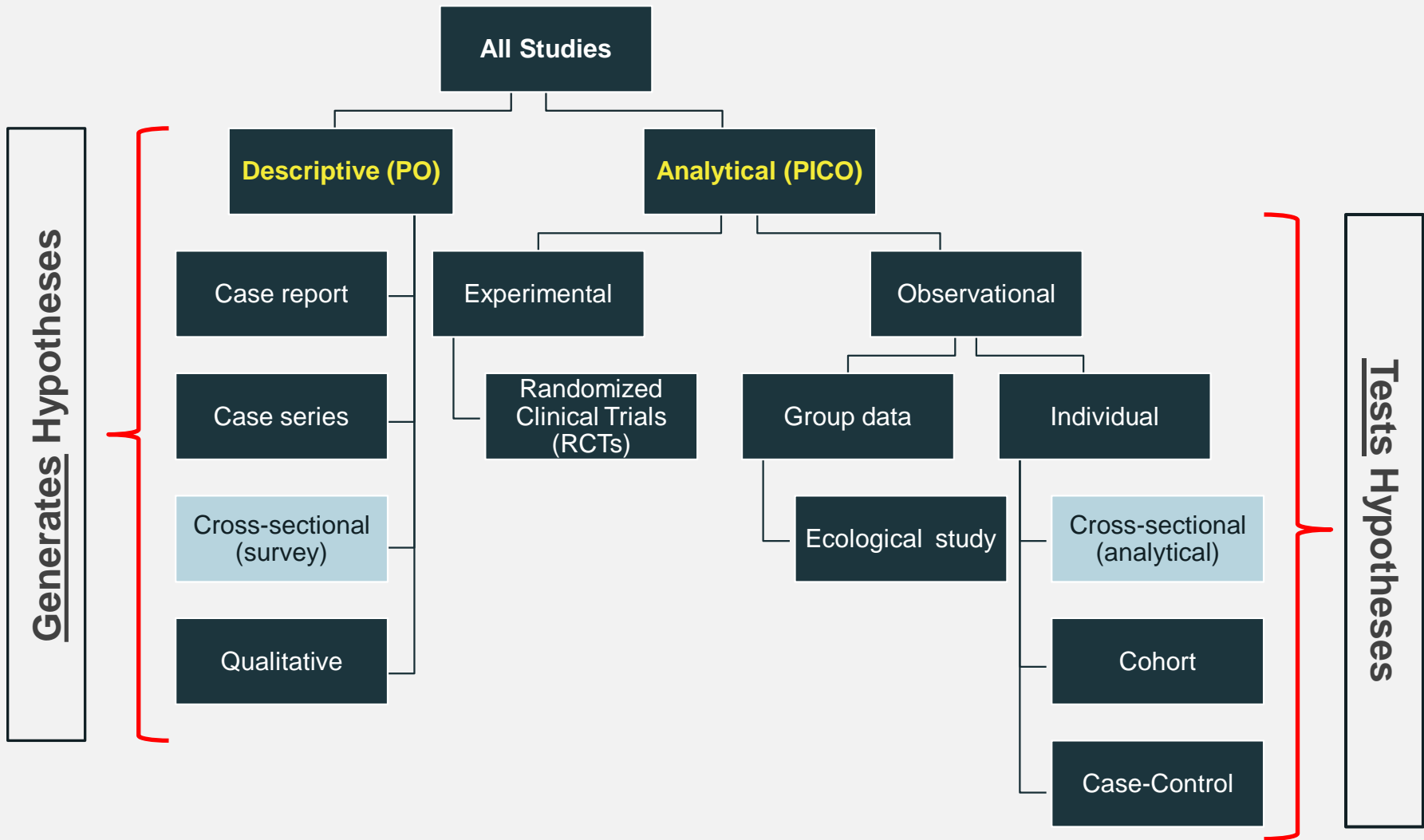
Assistant Professor - Community Medicine Unit,  
Family & Community Medicine Department

[nmalamro@ksu.edu.sa](mailto:nmalamro@ksu.edu.sa)

**Learning Objectives:** By end of this session students will be able to:

1. Describe types of cross-sectional studies
2. Identify steps for conducting cross-sectional studies
3. Identify issues in the design of cross-sectional studies
4. Describe the strengths and weaknesses of cross-sectional studies

# 1 Types of cross-sectional studies



A cross-sectional study is a study that either quantifies an outcome of interest **AND/OR** examines the relationship between disease (or other health related state) and other variables of interest as they exist in a defined population at a single point in time or over a short period of time.

# Types of Cross-Sectional Studies

Descriptive

Analytical

Study **prevalence** of health related events at a point in time/snapshot

(Diseases, risk factors, coverage of interventions, health service utilization, knowledge, attitude and practice)

Assess **association** between exposure and outcome.

Exposure and disease status are assessed **simultaneously** among individuals at the same point in time

**Compare prevalence** of disease in persons with and without the exposure of interest

---

---

In practice, cross-sectional studies will include an element of both types of design.

---

---

# When to Conduct a Cross- Sectional Study

---

- To estimate prevalence of a health condition or prevalence of a behavior or risk factor
- To learn about characteristics such as knowledge, attitude and practices of individuals in a population
- To monitor trends over time with serial cross-sectional studies (National example of cross-sectional studies of great importance is the National Health and Nutrition Surveys (**NHANES**)).




# { 2 How to conduct a cross-sectional study? }

## Steps in conducting a cross-sectional study


1- Define a **population** of interest  
(reference or source population)



2- Recruit a representative **sample**  
(adequate size, random selection)



3- Measure the **variables** of  
interest (exposure/outcome) at the  
same point in time



4- Analyze the **data**

The participants in a cross-sectional study are selected based on **the inclusion and exclusion criteria** set for the study.

**Identify Subjects from population**

**Collect data on exposure and outcome (e.g. disease)**

Exposed  
and have a  
disease

Not Exposed  
and have a  
disease

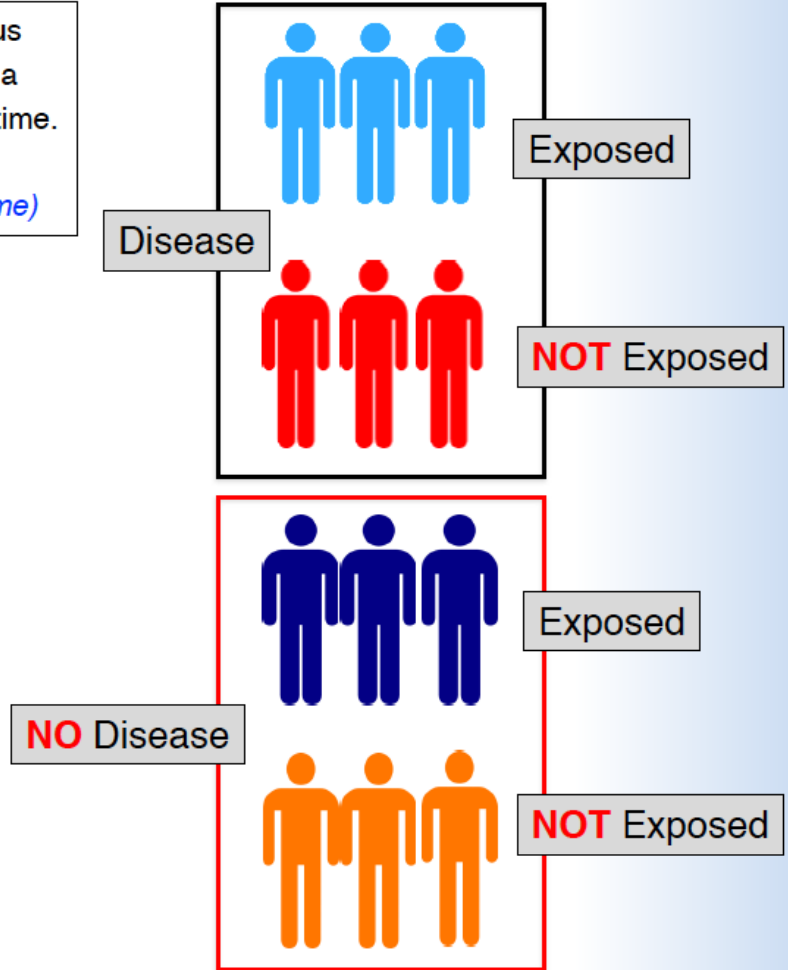
Exposed, and  
Do not have  
a disease

Not Exposed,  
and Do not  
have a  
disease

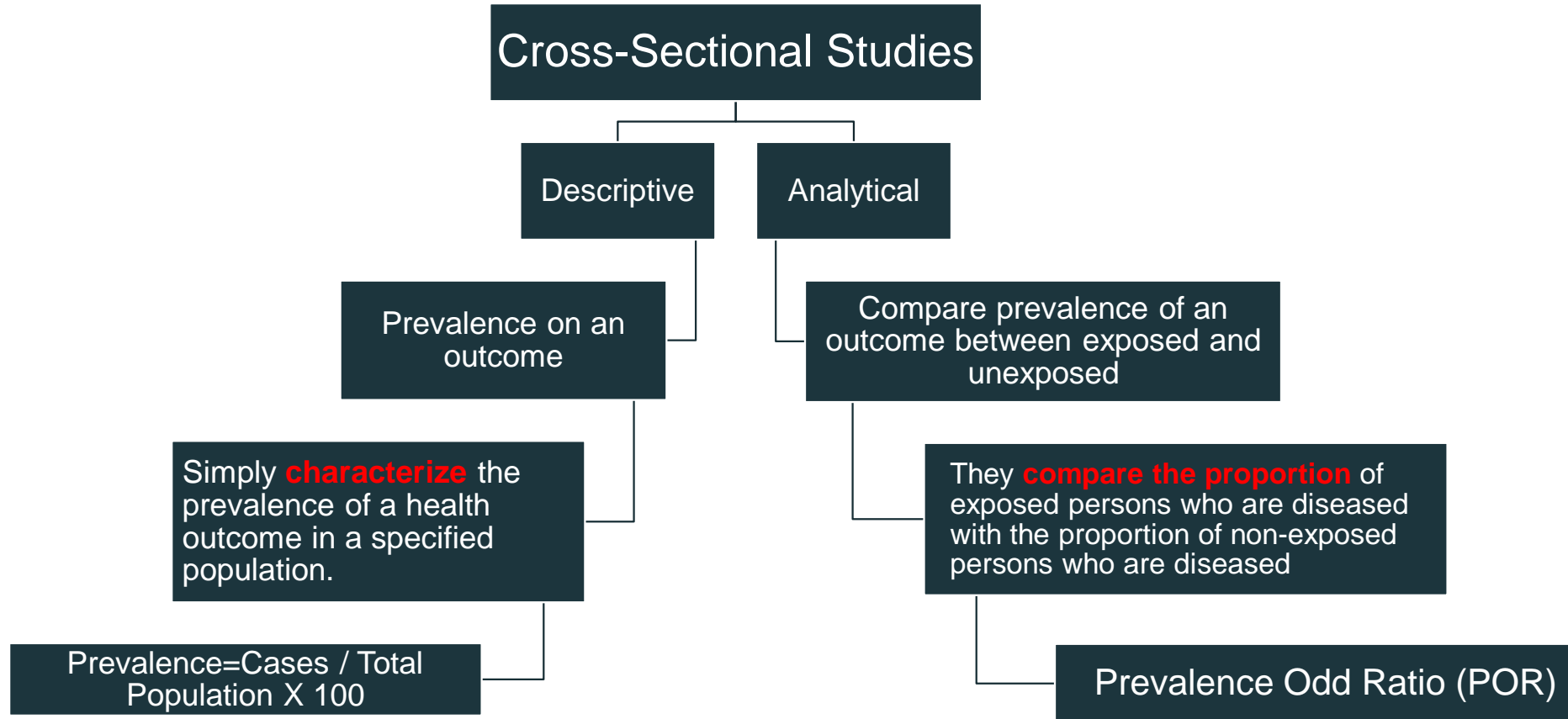
Measure disease and exposure status **simultaneously** among individuals in a well-defined population at a point in time.  
*(Snapshot of the health status of populations at a certain point in time)*



Study Population



# Measurement & Analysis in Cross-Sectional studies





# Vaping and Advertisement

You identify a random sample of young adults aged 18 – 25 at city of Riyadh.

**Exposure:** Ads about vaping

**Outcome:** Vaping

|        | Vaping | Not Vaping | Total |
|--------|--------|------------|-------|
| Ads    | 50     | 200        | 250   |
| No Ads | 50     | 700        | 750   |
| Total  | 100    | 900        | 1000  |

|        | Vaping | Not Vaping | Total |
|--------|--------|------------|-------|
| Ads    | 50     | 200        | 250   |
| No Ads | 50     | 700        | 750   |
| Total  | 100    | 900        | 1000  |

**Descriptive Cross-Sectional:**

What is the prevalence of vaping?

= Number of people who vape / Total population X 100

= 100 / 1000 X 100

= 10%

**Analytical Cross-Sectional:**

Does the prevalence of vaping vary by the status of exposure to advertisement?

I.e. What are the **odds** of vaping given exposure to advertisement versus not exposed to advertisement?





|        | Vaping      | Not Vaping   | Total |
|--------|-------------|--------------|-------|
| Ads    | 50 <b>a</b> | 200 <b>b</b> | 250   |
| No Ads | 50 <b>c</b> | 700 <b>d</b> | 750   |
| Total  | 100         | 900          | 1000  |

### Analytical Cross-Sectional:

Does the prevalence of vaping vary by the status of exposure to advertisement?

i.e. What are the **odds** of vaping given exposure to advertisement versus not exposed to advertisement?

$$\begin{aligned}
 \text{OR} &= \frac{\text{odds an exposed person develop the outcome (a/b)}}{\text{odds an unexposed person develop the outcome (c/d)}} \\
 &= \mathbf{ad / bc} \\
 &= (50 \times 700) / (200 \times 50) = 3.5
 \end{aligned}$$

**What does 3.5 mean??**

**The odds of vaping is 3.5 times higher after seeing a vaping advertisement as opposed to not seeing one.**

{ 3

**Issues in the design of cross-sectional studies**

}

# Choosing a representative sample

---

- A cross-sectional study should be representative of the population if generalizations from the findings are to have any validity.
- For example, a study of the prevalence of diabetes among women aged 40-60 years in Town A should comprise a random sample of all women aged 40-60 years in that town.

# Sample Size

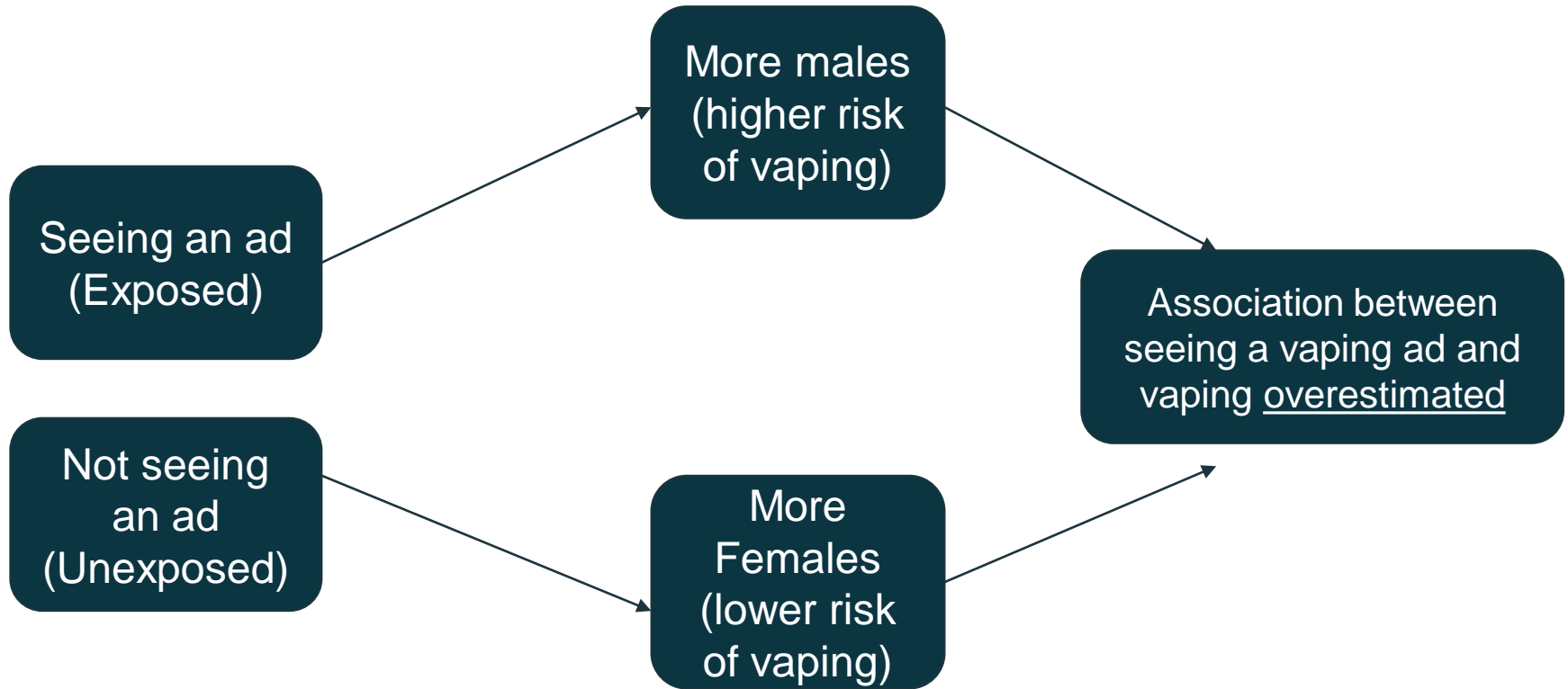
---

- The sample size should be sufficiently large enough to estimate the prevalence of the conditions of interest with adequate precision.
- Sample size calculations can be carried out using sample size tables or statistical packages such as Epi Info.

# Biases in Cross-Sectional Studies

---

1. **Selection Bias**: when the study participants are systematically different in their characteristics compared with eligible participants who were not selected for the study. **Common type**: Non-response bias.
2. **Recall bias**
3. **Confounding!**



**Confounding**

{ 4

**Strengths & Weaknesses**

}



## Strengths

- Relatively quick and easy to conduct
- Data on all variables is only collected once.
- Able to measure prevalence for all factors under investigation.
- Multiple outcomes and exposures can be studied.
- Good for descriptive analyses and for generating hypotheses.

## Weakness

- Difficult to determine whether the outcome followed exposure in time or exposure resulted from the outcome.
- Difficult to determine whether the outcome followed exposure in time or exposure resulted from the outcome.
- Associations identified may be difficult to interpret.
- Susceptible to bias due to low response and misclassification due to recall bias.

# Thank you!

**Office Hours (by appointment via email):**

Mondays & Wednesdays

11 AM – 1 PM

College of Medicine West Building

Level 1 - Office 4011034

---

[nmalamro@ksu.edu.sa](mailto:nmalamro@ksu.edu.sa)

---

## References:

- Celentano, David D., and Scd Mhs. Gordis Epidemiology. Elsevier, 2018.
- Hulley, Stephen B., ed. Designing clinical research. Lippincott Williams & Wilkins, 2007.
- Haynes, R. Brian. Clinical epidemiology: how to do clinical practice research. Lippincott williams & wilkins, 2012.
- Carlson, Melissa DA, and R. Sean Morrison. "Study design, precision, and validity in observational studies." Journal of palliative medicine 12.1 (2009): 77-82.
- The Centre for Evidence-Based Medicine develops, promotes and disseminates better evidence for healthcare. Study Design. NA. Accessed September 13, 2019: <https://www.cebm.net/2014/04/study-designs/>