Office of Biostatistics & Quantitative Health Sciences

JOHN A. BURNS SCHOOL OF MEDICINE

Research Using Large Healthcare Datasets Lecture 2

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Outline

Lecture 1 (10/17/2018)

- The goal of statistics
- Introduction to descriptive biostatistics
- Some research design and data presentation issues

Lecture 2 (12/19/2018)

Large databases

Lecture 3 (01/23/2019)

- Introduction to inferential statistics
- Some commonly used statistical approaches

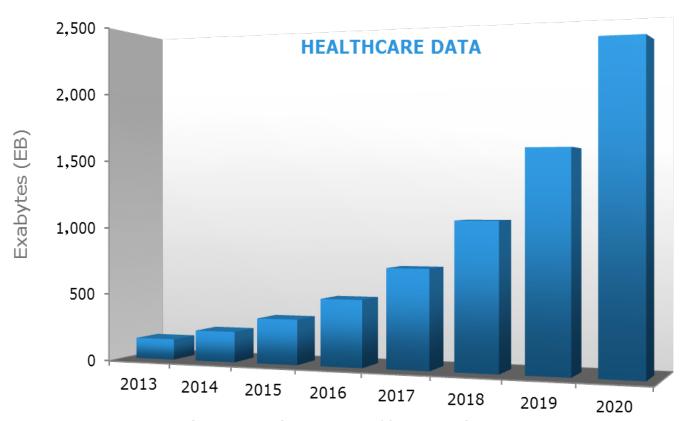


Lecture Objectives

- Clarify the goal of statistics
- Grasp descriptive statistics
- Be familiar with various data presentation approaches
- Introduce key concepts of inferential statistics
- Survey some commonly used statistical approaches
- Understand basic research design principles
- Understand the pros and cons of large databases
- Some examples of research using large databases
- Build a foundation which will facilitate the active participation in clinical and translational research



Growth of Healthcare Data

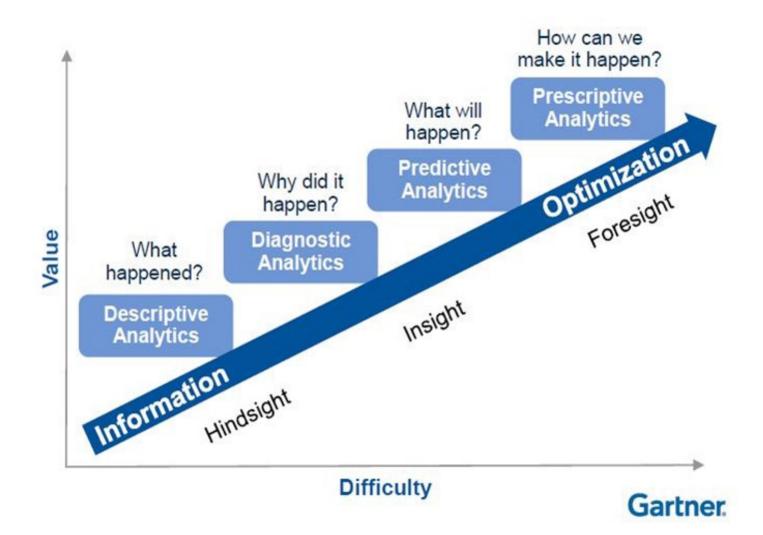


Enterprise Strategy Group 2011, Research report: North American Healthcare Provider Information includes hospitals & Ambulatory Health Care Provider Market Size & Forecast

Source: EMC Digital Universe with Research & Analysis by IDC

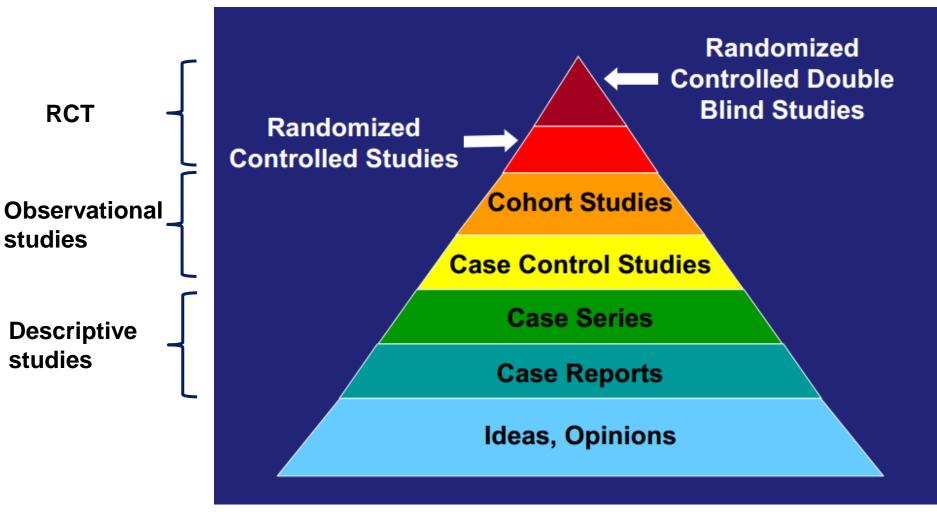


Analytics Are No Longer a Nice to Have





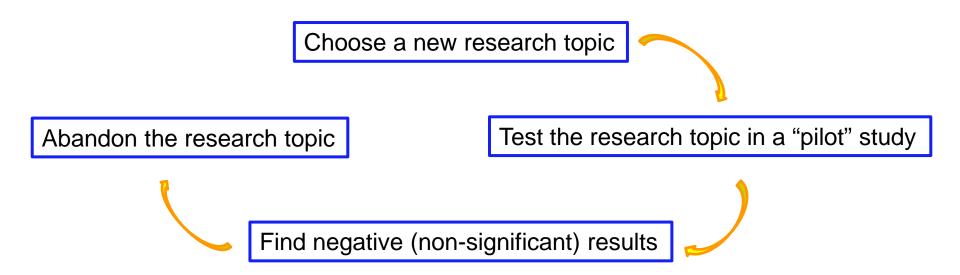
Clinical Research & Scientific Evidence



Barnett Kramer (NIH)



Nonproductive Research Strategy



A "pilot" study?

- Little or no funding?
- A vague and poorly developed research proposal?
- No time for detailed thought about subsequent studies?



Lancaster, GA. Pilot and feasibility studies come of age! *Pilot and Feasibility Studies* 2015; 1:1.



Pilot and Feasibility Studies

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Aims and scope

Aims and scope

Pilot and Feasibility Studies encompasses all aspects of the design, conduct and reporting of pilot and feasibility studies in biomedicine. The journal publishes research articles that are intended to directly influence future clinical trials or large scale observational studies, as well as protocols, commentaries and methodology articles. The journal also ensures that the results of all well-conducted, peer-reviewed, pilot and feasibility studies are published, regardless of outcome or significance of findings.

Pilot and feasibility studies are increasingly conducted prior to a full randomized controlled trial. However, these studies often lack clear objectives, many remain unpublished, and there is confusion over the meanings of the words "pilot" and "feasibility". Pilot and Feasibility Studies provides a forum for discussion around this key aspect of the scientific process, and seeks to ensure that these studies are published, so as to complete the publication thread for clinical research.

Standard Research Process

Identifying a research question and a hypothesis

Designing study and developing research protocol

Gathering preliminary data and revising the protocol

Conducting the study

Analyzing data analysis and interpreting results

Drawing conclusions and disseminating the results



Database Research Process

Identifying a research question and a hypothesis

Designing study and developing research protocol

Gathering preliminary data and revising the protocol

Conducting the study

Analyzing data analysis and interpreting results

Drawing and disseminating the conclusions



Primary versus Secondary Data

- Primary data: data collected specifically for the purpose of answering a research question and to meet specific research objectives
- Secondary data: data that have already been collected for some other purposes

Secondary data analysis: The practice of re-analyzing an existing data set for a purpose other than the purpose for which it was originally created.



Sources & Types of Secondary Data

- Government-funded datasets
- Records/files
- Internet
- Archives
- Previous studies including journal supplements

Some are restricted and others are publicly available



Some Large Clinical Databases

American Hospital Association Annual Survey	Annual survey data from US hospitals with focus on organiztaional characteristics and health care utilization			Medical Expenditure Panel Survey (MEPS)	families,	ide panel surveys of indiv health care providers, ar rs covering a variety of to			
AMA Physician Masterfile	Database of US physicians with information on education, training, and professional certification			Medical School Graduation Questionnaire and Matriculating Student	Nationwide surveys of medical students about career choices, medical school admissions process, and educational				
Area Resource File (ARF)	professionals, and census da			Questionnaire	Claims	Survey of Adolescent	National longitudinal survey of causes and outcomes of health-related behaviors in		
Atherosclerosis Risk in Communities (ARIC)	subjects and o		Popul	Medicare Claims Data	demogra wide var settings	National Program of	Data assem	and young adults bled from state cancer registries formation about patient	
Behavioral Risk Factor Surveillance System (BRFSS)	Serial cross-se health risk beh	Health and Retirement Study (HRS) & Asset	Longi		Panel su use, hea demogra	(NPCR)	counts by g	cs as well as cancer rates and eographic area database of treatment for	
California Health Interview Survey (CHIS)	Sprial cross-se	and Health Dynamics among the Oldest Old	and e		expendit Longitud adults in	Myocardial Infarctions (NRMI)	patients hos infarction	spitalized with myocardial	
Canadian Institute for Health Information databases	children in Cal	HCUP State Inpatient Databases (SID) HCUP State Ambulatory	from i	States (MIDUS)	sociodei characte	National Surgical Quality Improvement Program (NSQIP)	procedures data on pre	going data collection on major surgical cedures from U.S. hospitals including a on preoperative patient characteristics, apperative variables, and morbidity and tality	
Cardiovascular Health Study (CHS)	issues	HCUP State Emergency Department Databases	partic Clinic from 6	(Long-Term Care)	resident	National Survey of Ambulatory Surgery	Nationally-re ambulatory	epresentative survey of surgery in hospital-based and g ambulatory surgery centers	
Community Tracking Study (CTS) Consumer Assessment of Healthcare Providers and Systems (CAHPS)	Family of surv	HCUP Nationwide Inpatient Sample (NIS) HCUP Nationwide	Natio	Medical Care Survey	sectiona emerger	Surveillance, Epidemiology and End Results Program (SEER), and SEER- Medicare data		ncer registry database, available but linkages to Medicare data	
SGIM Society of General Internal Medicine		Sample (NEDS) Health Information	depar Biann knowl	ment visits ual telephone survey of the public's edge, attitudes, and behavior ling cancer		U.S. Census data Bureau; cens		provided by the U.S. Census insus data can be linked to other in example to obtain median each U.S. zip code	
					U.S. Renal Data System (USRDS)		ationwide data on patients with enal disease, collected on an		

aspects of cancer care

Creating Value for Patients

ongoing basis

Some Education & Social Sciences Datasets

- Common Core of Data (CCD)
- Current Population Survey (CPS)
- Early Childhood Longitudinal Study (ECLS): Birth (ECLS-B) and Kindergarten (ECLS-K) Cohort
- General Social Survey (GSS)
- Head Start Family and Child Experiences Survey (FACES)
- Monitoring the Future (MTF)
- National Assessment of Educational Progress (NAEP)
- National Education Longitudinal Study (NELS)
- National Household Education Surveys (NHES)
- National Longitudinal Study of Adolescent Health (Add Health)
- National Longitudinal Survey of Youth (NLSY)
- National Survey of American Families (NSAF)
- National Survey of Child and Adolescent Well-Being (NSCAW)
- National Survey of Families and Households (NSFH)
- NICHD Study of Early Child Care and Youth Development (SECCYD)
- Programme for International Student Assessment (PISA)
- Progress in International Reading Literacy Study (PIRLS)
- Trends in International Mathematics and Science Study (TIMSS)
- U.S. Panel Study of Income Dynamics (PSID): Child Development Supplement (CDS)



Useful Websites for Secondary Databases

- Office of Minority Health: <u>https://www.minorityhealth.hhs.gov/Default.aspx</u>
- Society of General Internal Medicine: <u>http://www.sgim.org/communities/research/dataset-compendium/proprietary-datasets</u>
- UCSF Large Dataset Inventory: <u>http://ctsi.ucsf.edu/research/celdac</u>
- Institute for Social Research University of Michigan: http://www.icpsr.umich.edu/icpsrweb/ICPSR/index.jsp
- JABSOM CIM: <u>http://oitwp02.jabsom.hawaii.edu/cim/research/healthc</u> are-databases/



Behavioral Risk Factor Surveillance System (BRFSS)

- A large, cross-sectional annual survey data from 1984 (~500,000/year)
- Emphasize health-related risk behaviors, chronic health conditions, and use of preventive services
- Conducted at the state level and complied for public use
- Hawaii data identifiable

Behavioral Risk Factor Surveillance System









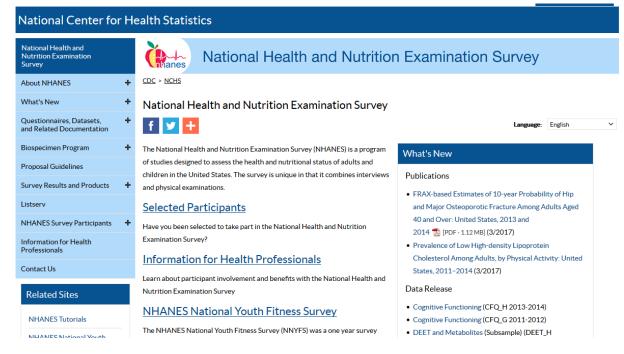


The Behavioral Risk Factor Surveillance System (BRFSS) is the nation's premier system of health-related telephone surveys that collect state data about U.S. residents regarding their health-related risk behaviors, chronic health conditions, and use of preventive services. Established in 1984 with 15 states, BRFSS now collects data in all 50 states as well as the District of Columbia and three U.S. territories. BRFSS completes more than 400,000 adult interviews each year, making it the largest continuously conducted health survey system in the world. See

https://www.cdc.gov/brfss/

National Health and Nutrition Examination Survey (NHANES)

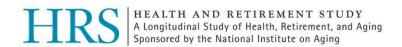
- A national biennial cross-sectional survey (~5,000 persons / year)
- A rich assortment of data based on
 - Interviews: demographics, socioeconomic status, dietary, and health behavior questions
 - Direct examination: medical, dental, physiological measurements, and lab tests





Health and Retirement Survey (HRS)

- A representative, longitudinal national study of aging, beginning at age 50
- Multiple cohorts of spouses, maintaining ~26,000 active participants
- Biennial examinations on all survivors, started in 1992



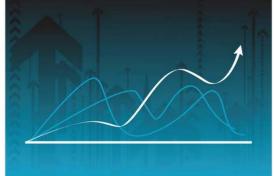
A public resource for data on aging in America since 1990

For more than 20 years, the Health and Retirement Study (HRS) has been a leading source for information on the health and economic well-being of adults over age 50 in the United States. We do this by asking people across the country to participate in the study and tell us about their lives.

As the world around us changes, the HRS follows the impact of these changes on our older population. This information has been used by scientists to publish over 3,000 books, articles and papers.

The HRS is a longitudinal project sponsored by the National Institute on Aging (NIA U01AG009740) and the Social Security Administration. The study director is Dr. David R. Weir of the Survey Research Center at the University of Michigan's Institute for Social Research.







http://hrsonline.isr.umich.edu/

Hawaii Hospitalization Database

- 26 hospitals and medical centers, with over 100,000 hospitalizations per year
- Over 90 variables (e.g., age, payer, hospital and location of residence, admission/discharge dates, LOS, total charges, disposition)
- 13,000 ICD-9 codes (primary and 19 secondary diagnoses or procedures)
- 32 different ethnic categories, including over 20 Asians/PIs, besides
 NH
- Gathered in Hawaii: before 2017 by HHIC; after 2017 by Laulima Data Alliance
- State and National Healthcare Cost and Utilization Project (HCUP) data







Pros of Large Clinical Databases

- All subjects, large population, real world patients and settings, ideal for comparative-effectiveness and costeffectiveness research
- Often contain longitudinal data across multiple years
- Relatively low-cost and quick alternative to generating own data
- Convenient for exploratory analyses and developing research hypothesis
- Large sample sizes, allowing subgroup and multivariable analysis
- Education and training use



Cons of Large Clinical Databases

- Data may not be of research quality
- No control over what data is collected
- Lack of compatibility between data and research questions
- Usually very large and complex
- Require not only analytical and technical expertise, but familiarity and experience with specific databases





QHS 650: Secondary Data Analysis QHS 651: Secondary Data Analysis Practicum

Spring 2019 (2-Credits Each)
Tuesday Afternoons (1:00-4:30 pm) at Kaka'ako Campus
in Health Sciences Library Computer Lab (MEB 107D)

Why Take This Course?

- Secondary data analysis takes advantage of data originally collected for other purposes in order to answer additional research questions.
- Many secondary data sources are readily available but underutilized. As such, they provide a rich opportunity for research and subsequent publications.
- QHS 650 provides a didactic overview of issues in secondary data analysis. This
 course will allow students new to using secondary data to become comfortable
 with accessing the data, forming hypotheses, and designing study proposals. In
 addition, the course introduces both basic and advanced statistical approaches for
 analyzing secondary data.
- The practicum course (QHS 651) is intended for students who plan to analyze secondary datasets in their own research. The course adds the hands-on experience needed to conduct individual research using secondary databases.

Course Objectives

- · Learn how to locate and download public-release data;
- Learn the steps involved in conducting research using secondary data;
- Provide hands-on training for students to perform proper secondary data analysis using R (only for QHS 651).

Prerequisites: QHS 650: None; QHS 651: QHS 601 or equivalent, and QHS 651 (which can be taken concurrently with QHS 650)

Instructors: Eunjung Lim, Ph.D., & James Davis, Ph.D., Office of Biostatistics & Quantitative Health Sciences, Department of Complementary & Integrative Medicine, John A. Burns School of Medicine, University of Hawaii

Class Time: 1:00-2:40 pm for QHS 650; 2:50-4:30 pm for QHS 651

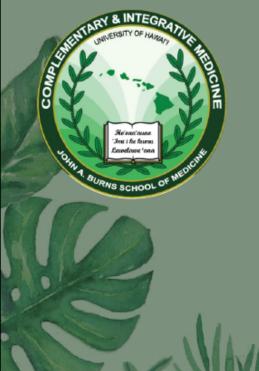


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The Clinical and Translational Research (CTR) graduate program will prepare graduates with skills for successful careers in clinical and translational research and research support.

Clinical Research (CR) Track

Develop knowledge and skills to investigate clinical research topics though coursework and research projects focused on research design, methodologies, quantitative methods, scientific writing, ethical issues, and the capacity in obtaining research funding.

Quantitative Health Sciences (QHS) Track

Courses and research projects focus on biostatistical and bioinformatic methods development and application to improve population and individual health. Students will acquire big data skills and master the scientific principles and methodologies that underlie basic science, clinical, and translation research.

Career

Research, research support, data analyst positions at:

- -Academia
- -Hospitals
- -Government agencies
- -Healthcare organizations
- -Pharmaceutical companies

Program Curricula

- -2-year 34 total credit hours graduate program
- -Plan A (Thesis):
- 24 credits of didactic courses
- -Plan B (Capstone Project):28 credits of didactic courses

How to Apply

Visit http://manoa.hawaii.edu/graduate/ content/clinical-research to either fill out an application or download a PDF form Application Deadline: May 30

For more information

Phone: (808) 692-1840

Email: GradCTR@hawaii.edu

Web: http://msctr.jabsom.hawaii.

MSCTR Graduate Program Website

https://msctr.jabsom.hawaii.edu/



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Department of Complementary & Integrative Medicine
John A. Burns School of Medicine

Clinical and Translational Research Graduate Program



The Clinical and Translational Research (CTR) graduate program will prepare graduates with skills for successful careers in clinical and translational research and research support. The CTR program leading to a MS degree is currently offered with two tracks, both available in either Plan A (thesis option) or Plan B (capstone project option):

- 1. Clinical Research (CR)
- 2. Quantitative Health Sciences (QHS)

The CR track focuses on the study of methods suitable to investigate clinical research topics. Students enrolled in the CR track are required to complete a combination of course work and clinical research. The competency domains include clinical and translational research, quantitative health skill, professionalism, communication, and interdisciplinary collaboration. Students will also develop the ability to identify and resolve ethical issues in clinical research, to ensure the safeguarding of human subjects, and to understand the workings of Institutional Review Boards and relevant organizational requirements. In addition, students will develop and/or increase their capacity in obtaining research

Admission

Program Requirements

MS Curriculum

MS Key Course Descriptions

MS Student Learning Outcomes

Tuition & Financial Aid

Regular Graduate Faculty

Contact

Graduate Student Handbook

Lecture Objectives

- Clarify the goal of statistics
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