Assessing Data Quality

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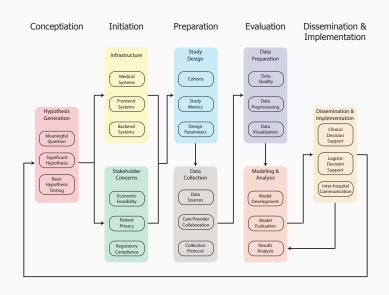
Center for Health Data Innovations

## **Assessing Data Quality**

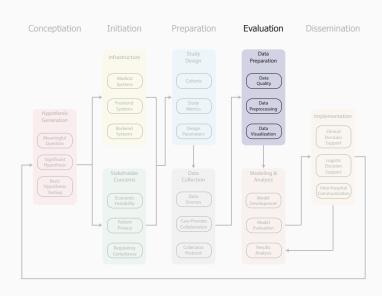
#### After this lecture students will be able to

- Assess the quality of data
- Trace the steps where data quality can be affected
- Define measures to ensure quality assurance and quality control of data
- Describe the components of an ETL pipeline
- Examine data for problems and discuss possible causes
- Design a process for imputation of missing data

# **Bioinformatics Pipeline**



## **Data Quality**



## **Overview**

Analysis is only ever as good as the data it's built upon.

- What is data quality? What makes data high quality vs low quality?
- Where along the process can you affect data quality?
- How can you design a study to collect high quality data (Quality assurance)?
- How can you identify and correct errors during and after data collection (Quality control)?

Overview

Analysis is only ever as good as the data it's built upon

Overview

- . What is data quality? What makes data high quality vs low quality? . Where along the process can you affect data quality?
- . How can you design a study to collect high quality data (Quality
- . How can you identify and correct errors during and after data
- collection (Quality control)?

 Data quality consists of the objective faccuracy, validity (not outside range of possibilities. all data is for the same pt, formatting requirements, DICOM dates), reliability (dx matches problem list matches coding), legibility (units, shorthand)] and subjective [completeness]

#### Steps

- Definition/Design lack of clear definitions for data items/collection, incompatible units, precision, scope, depth
- Collection not enough documentation (drug given/dosage altered but no start and end date), non-adherence to data definitions (collecting data outside of protocol time), human variance/error (bp cuff, RR, incorrect units), Orders are placed (procedures, medications) which are not connected to a rationale or sufficient reason
- Processing interpretation ('initial' lab, diagnosis date), coding error (mis-entering information such as order of birthdate, or height as 9 cm instead of 90 cm), random (mistyping, illegible handwriting), software errors, Assigning codes to problems treated vs problems tested and ruled out, which complaints do you code/document (doctors as coders)

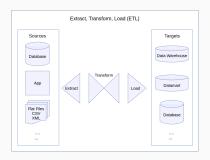
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- How can you identify and correct errors during and after data
- collection (Quality control)?
- quality assurance training of personnel (mock exams and reporting), site visits, reduce open-ended questions
- quality control data monitoring (compare to independent source), hand verification, entering data in twice (by different sources), consistency checks

# Extract, Transform and Load (ETL)

- Extract
  - Extract data from source(s)
- Transform
  - Modify the data for the purposes of analysis or further querying
- Load
  - Load the data into the target database



# **Quality Assurance - DICOM**

- DICOM Digital Imaging and Communications in Medicine is the international standard for medical images and related information. It defines the formats for medical images that can be exchanged with the data and quality necessary for clinical use
- DICOM groups information into data sets, e.g., an x-ray would contain the patient ID within the file, so that the image can never be separated from this information by mistake.
- DICOM Value Representations

https://www.dicomstandard.org/about/

# Quality Assurance - DICOM

name	VR	value
Group Length	UL	532
Image Type	CS	DERIVED
SOP Class UID	UI	1.2.840.10008.5.1.4.1.1.2
SOP Instance UID	UI	1.2.840.114356.2008.11.30.12.34.2.329.999
Study Date	DA	20081230
Content Date	DA	20081230
Study Time	TM	122731
Content Time	TM	12299.0000
Modality	CS	CT
Institution Name	LO	Manhasset Diagnostic Imaging
Station Name	SH	
Study Description	LO	MOSES CT Outside Reference Images
Procedure Code Sequence	SQ	[{(0008, 0100): (0008, 0100) Code Value
Code Value	SH	MOSESOUTREFCT
Coding Scheme Designator	SH	GEIIS
Coding Scheme Version	SH	0
Code Meaning	LO	MOSES CT Outside Reference Images
Series Description	LO	Reformatted
Referenced SOP Class UID	UI	1.2.840.113619.2.51762891606.1649.1005918257.250
Referenced SOP Instance UID	UI	1.2.840.114356.2008.11.30.12.34.2.329.1301

# **Quality Assurance - DICOM**

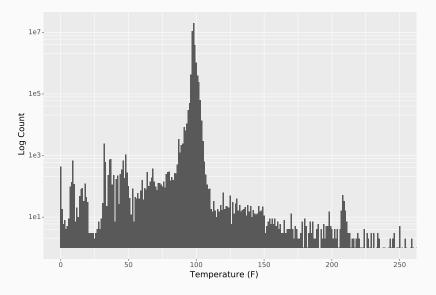
name	VR	value
Study Date	DA	20081230
Content Date	DA	20081230
Study Time	TM	122731
Content Time	TM	12299.0000

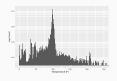
- DA A string of characters of the format YYYYMMDD
- **TM** A string of characters of the format HHMMSS.FFFFFF.
  - One or more of the components MM, SS, or FFFFFF may be unspecified as long as every component to the right of an unspecified component is also unspecified

Whose fault is this?

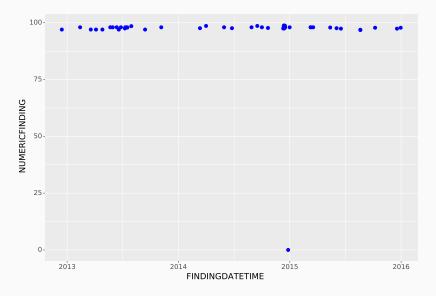
# **Quality Control - Temperature**

How can I find the temperatures recorded from every patient in the hospital





- How do you interpret the temps around 0 (probably had to enter something), how about around 37F (centigrade), how about 212, how about 95 (MICE)
- Let's take a look at an individual patient's data, who had a temp of 0
- maybe the data is being pulled from 5 different hospitals and it's the ETL which is causing the errors, because it doesn't know Celsius from F
- let's pick a patient whose temperature is zero and see what the rest of their temperature values look like (and then look at her results for that date)



# **Associated Values**

FINDINGDATETIME	FINDINGDESC	NUMERICFINDING
2014-12-26	PULSE OXIMETRY	97.00
2014-12-26	WEIGHT/SCALE (ounces)	2800.16
2014-12-26	HEIGHT (inches)	62.00
2014-12-26	Diastolic Blood Pressure	82.00
2014-12-26	Systolic Blood Pressure	139.00
2014-12-26	HEIGHT (CM)	157.48
2014-12-26	PULSE	75.00
2014-12-26	BODY MASS INDEX	32.13
2014-12-26	O2 SAT%	97.00
2014-12-26	TEMPERATURE (F)	0.00
2014-12-26	Systolic Blood Pressure	139.00
2014-12-26	WEIGHT (KG)	79.38
2014-12-26	Diastolic Blood Pressure	82.00

# Imputation and Extrapolation

# Can we develop a systematic way to deal with missing data

• What are the different ways that data could be missing

Can we develop a systematic way to deal with missing data

\* What are the different ways that data could be mining

Imputation and Extrapolation

## Imputation and Extrapolation

- data could be MCAR, MAR, MNAR or because we are slicing the data into chunks smaller than the sampling rate
- Missing Data Procedure
  - Variable correctness var correctly derived/appropriate to include, e.g., complete or near-complete missingness or same value in all rows.
  - Time freq Ensure that time blocks used in time series data are appropriate to the task
  - Determine how frequently every variable is measured
  - Use the frequency range from the previous step for each variable to do ffill
  - Encounters without data or good data cannot add value and should be dropped.
  - Drop beginning blocks if empty, drop end blocks if dead or discharged
  - Imputation MICE, NN. Cases where imputation should not be done are when the
    missingness itself is significant or if the imputation cannot be done by adding another
    class. An example of the latter is would be if an x-ray is performed. X-rays not being
    performed are another class that can be added to the column.
  - Anything which is not imputed is masked (-9999, not 0)

### Sources

- WHO data quality
- Healthcare Data Warehousing and Quality Assurance
- (2002). Defining and improving data quality in medical registries JAMIA, 9(6), 600-611.