Data Linkages

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- Use an in-house SAS program
 - Created by Glenn Wright
 <u>https://www.lexjansen.com/wuss/2011/data/Papers_Wright_G_7</u>
 <u>6128.pdf</u>
- Probabilistic matching algorithm
- Point-based/similarity scores
 - Point system for rewarding how close a match is
- Also includes nicknames for first name matching



Step 1: Standardize and clean data

mnamemom = scan(fnamemom,2,' '); fnamemom = scan(fnamemom,1,' '); fnamemom=compress(fnamemom, 'abcdefghijklmnopqrstuvwxyzABCDEFGHIJKLMNOPQRSTUVWXYZ', 'k'); lnamemom=compress(lnamemom, 'abcdefghijklmnopqrstuvwxyzABCDEFGHIJKLMNOPQRSTUVWXYZ', 'k'); lnamedad=compress(lnamedad, 'abcdefghijklmnopqrstuvwxyzABCDEFGHIJKLMNOPQRSTUVWXYZ', 'k');

addmom=tranwrd(addmom,'STREET','ST'); addmom=tranwrd(addmom,'AVENUE','AVE'); addmom=tranwrd(addmom,'DRIVE','DR'); addmom=tranwrd(addmom,'ROAD','RD'); addmom=tranwrd(addmom,'COURT','CT'); addmom=tranwrd(addmom,'UNIT',''); addmom=tranwrd(addmom,'APT',''); addmom=tranwrd(addmom,'POBOX',''); addmom=tranwrd(addmom,'LANE','LN'); addmom=tranwrd(addmom,'SOUTH','S'); addmom=tranwrd(addmom,'SOUTH','S'); addmom=tranwrd(addmom,'NORTH','N'); addmom=tranwrd(addmom,'NORTH','N'); addmom=tranwrd(addmom,'NORTH','N'); addmom=tranwrd(addmom,'WEST','E');

addmom=tranwrd(addmom,'1ST','FIRST'); addmom=tranwrd(addmom,'2ND','SECOND'); addmom=tranwrd(addmom,'3RD','THIRD'); addmom=tranwrd(addmom,'4TH','FOURTH'); addmom=tranwrd(addmom,'5TH','FIFTH'); addmom=tranwrd(addmom,'6TH','SEVENTH'); addmom=tranwrd(addmom,'8TH','EIGTH'); addmom=tranwrd(addmom,'9TH','NINTH');

BDADDRESSMATCH=compress(addmom, '0123456789abcdefghijklmnopqrstuvwxyzABCDEFGHIJKLMNOPQRSTUVWXYZ', 'k').

BDADDRESSMATCH=COMPRESS (BDADDRESSMATCH) ;

Step 2: Assign weights by frequencies

* Macro to create weights for how frequent values of various variables are - an exact or near match of a rare name (or bday, or zip code address) is more of a sign of an actual match than a match of something more common - code copied from Glenn Wright;

Smacro create_weights_fmt (data=, var=, fmtname=,log=, asian_adjust=FALSE);

```
%* create variable freq which is amount each value of a variable
shows up as a percent of all valid values
insert value of variable into new variable start -
do proceedure differently for logarithmic weights
and for straight-up counts;
proc sql;
create table temp as
select &var. as start, count(&var.)/&n. as freq
from &data.
where &var. is not missing
group by &var.;
quit;
```

%* Create a format - variable value maps to the left side ("start"), result of equasion maps to ("label"); data temp2; set temp end=last;

```
fmtname = "$&fmtname";
```

label = put(-log2(freq), z8.3);

```
if last then do;
    start = '';
    hlo = 'o';
    label = put(log2(&n.),z8.3);
end;
run;
```

Example: Sarah has less weight than Pearl



Step 3: Build SAS macro/code similarity scores

```
1%macro sql_blocking(var1,var2,var3);
```

```
DOB unless year is the same - this part will only get triggered if year is different */
when (month(a.date_of_birth)=1 & month(b.date_of_birth)= 1 & day(a.date_of_birth)= 1 & day(b.date_of_birth)= 1)
    then -8
/* month and day agree, year disagrees */
```

```
when the day agree, your distinctes / when (month(a.date_of_birth)=month(b.date_of_birth) & day(a.date_of_birth)=day(b.date_of_birth)) then 5
/* year agrees, month and day different */
when year(a.date_of_birth)=year(b.date_of_birth) then -7
/* Complete disagreement */
else -11 end as score dob,
```

from	&var2. as a INNER JOIN &var2. as b	
on	a.&var1. is not missing	Cartesian product of A, B, and C:
and	b.&var1. is not missing	
and	a.id < b.id	$A \leftrightarrow B$
and	b.id > &maxid.	$A \leftrightarrow C$
and	a.&var1. = b.&var1.	$B \leftrightarrow C$
where	calculated score >= 21	



⊡proc sql;

```
create table linked_pairs as
%sql_blocking(ssn,setx11,1)
UNION
%sql_blocking(date_of_birth,setx11,1)
UNION
%sql_blocking(first_name,setx11,1)
UNION
%sql_blocking(last_name,setx11,1)
;
quit;
```



• Total scores are summations of individual similarity scores

calculated score_ssn + calculated score_sex + calculated score_fname + calculated score_pris +
 calculated score_lname + calculated score_dob + calculated score_race + calculated score_middle
 + calculated score_geo as score

Histogram of total scores





Final Step: Deduplication

- Use an in-house SAS program to deduplicate the linked pairs output from our matching algorithm
 - Also created by Glenn Wright <u>Microsoft Word -</u> <u>WUSS2010_final_hash_082410.doc (lexjansen.com)</u>
- SAS hash objects to transform the file of linked pairs into a file of clustered records with new common identifiers.
 - In linked pairs file, we have redundant links (e.g., A and B AND and B and C AND A and C are the same person.



– This program cleans this up and gives us a patient level registry!

Issues

- The program takes around 3 days to run.
- We broke it!
 - Tried to do an inner join of ~8 million records and ran out of memory
 - Tried to overcome this issue by matching new data (~2 million) to old registry (~6 million)
 - For context, this worked and created a linked pairs file with ~36 million records.
- We broke it again!
 - Tried to run the deduplication program and ran out of a memory again
 - Because it uses hash objects, it is much more difficult to troubleshoot compared to the matching program.







Methodology issues

- The registry was recreated every month from scratch
 - not very computationally effective, especially in SAS
- The *link* ID that was created was flawed
 - Different individuals had the same *linkID*
 - Same individual had different *linkID*
- De-duplication method was more complicated than it needed to be
- Inconsistent formatting issues as well



Towards Developing a new HCV Registry

- Import and clean up Legacy data up to March 2022
 - Standardize dates, data_sources, last and first names, lab names, lhj, dob....
 - Removed complete duplicates and observations without complete last name, sex, dob and lhj (~2% of the data)
 - 5,561,964 Observations (18GB)
- Currently there is no ELR data for Los Angeles or San Diego
 - but we keep trying 💽
- There is no longer a link ID, well sort of anyway,
 - If there was a CalREDIE personID associated with any of the observations within a linkID then the personID was assigned for that person. If personID was still missing, linkID was temporarily substituted, with the distinction of having leading zeroes







System Workflow





System Architecture

HCV Registry Linkage.1

- Extensive matching is no longer necessary since the personID is set, but matching to old data is still necessary
- Connect to SQL Server via RODBC connection in R to perform deduplication and matching and send new events back to SQL Server
 - Standardize dates, data_sources, last and first names, lab names, lhj, dob....
 - Added report year and age group
 - Using the *recordlinkage* package in R
 - First run based on last name, sex, dob and lhj
- Populate patientID from CalREDIE where missed in first substitution effort



HCV Registry Linkage.2

• IN RSTUDIO SERVER:

- Import last 3 years, from date of production, using the CalREDIE DDP in two flat files currently processed in SAS >>> SQL:
 - DDP_UDF_Extract_ChronicHepC
 - DDP_System_Lab_Extract_ChronicHepC
- Import new Non-participating Jurisdiction data currently processed in SAS >>> SQL
- Import hcv_incident



HCV Registry Linkage (R space)



RECORD LINKAGE

- Record linkage definition: determine if pairs of data records describe the same entity, join two heterogeneous relations and remove duplicates from a single relation
- Use *recordlinkage* package in R
 - Methods based on a stochastic approach are implemented as well as classification algorithms from the machine learning domain.
 - Further documentation found here: <u>https://cran.r-</u> project.org/web/packages/RecordLinkage/RecordLinkage.pdf.



Record Linkage in action

```
hcv_pairs <-
 cr %>%
 select(person_ID, firstname, lastname, dob, sex, lhj) %>%
 RLBigDataLinkage(incident %>%
                    select(person_ID, firstname, lastname, dob, sex, lhj),
                 blockfld = c("dob"),
                 strcmp = c("lastname","firstname"),
                                                          the blockfld option specifies a set of
                 exclude = c("person_ID", "sex", "lhj")
                                                          columns in which two records must agree to
hcv_gotpairs <-
                                                           be included in the output
getPairs(
 epiClassify(epiWeights(hcv_pairs), 0.95),
 #rl_epiclass.
                                                           strcmp are which columns are the
        filter.link = c("link", "possible"),
                                                          comparisons being ran on.
        single.rows = T) \%
 as_tibble()
                                                          epiClassify Classifies record pairs as link,
hcv_gotpairs_wide <- hcv_gotpairs %>%
                                                           non-link or possible link based on weights
 select(-id.1, -id.2, -is_match, -Class) %>%
 rename(person_ID = incidentid.1,
                                                          computed by epiWeights which are weights
        RECIP_ID = incidentid.2,
        sex = sex.1,
                                                          for Record Linkage based on an EM
        RECIP_SEX = sex.2
                                                          algorithm
        ) %>%
  left_join(df_cases %>%
           by = "incidentid"
                                                          RLBigDataLinkage Represents a record
            <u>%>%</u>
                                                          linkage setup with two datasets which are to
 left_join(incident,
           by = "RECIP_ID"
                                                          be linked together.
```

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Deduplication

 After matching is done and patient_IDs have been assigned, the dataset is deduplicated and merged with the existing incident level data using a simple SQL code chunk such as"

SELECT *

FROM hcv_incident

UNION

SELECT DISTINCT patient_ID, dob, age, agemnth, sex, ssn, race_ethnicity.....

FROM temp;



PERSON LEVEL DATASET

- From *hcv_incident* a person-level registry file (*hcv_person*) contains aggregated data:
 - Patient_ID, demographics
 - First, last, classified & confirmed: dates, lhj, pregnant status, prison status, homeless status...
 - Genotype classification and current case status

