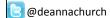
Data Management and Accessibility

Deanna M. Church Senior Director of Genomics and Content Personalis, Inc





Short Course in Medical Genetics 2014

Disclosures

Deanna Church works for Personalis, Inc. A company that provides whole genome and whole exome sequencing and analysis services.

The nature of data management has changed

Was...



- Motes
- Pictures of gels
- Autoradiograms

Now...



- Notes
- Data images
- Sequence files

http://www.sxc.hu/photo/1072645

http://www.hcii.cmu.edu/M-HCL/2006/MEDRADProject/images/notebook.jpg

Protocols

Preparation of Electrocompetent E. coli (i.e. DH5a)

revised 2/24/96

1 L 2XYT media (no antibiotics!), store at RT 1 L chilled autoclaved dH₂0, stored in 4° cold room.

Before starting procedure, prepare/chill the following:

1 L chilled autoclaved dH₂0, stored in 4° cold room. 100ml chilled 10% glycerol / dH₂0 solution, store at 4°C.

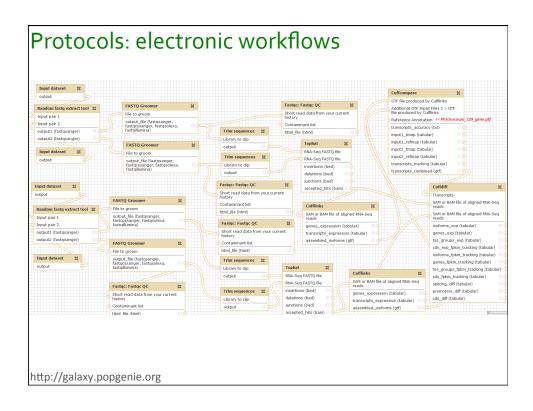
Corning 250ml pointed bottles (w/ orange caps) with white adapters chilled at 4°C. (used in Beckman tabletop GPR with GH-3.7 rotor, Church 64)
Rinse bottles first with 95% EtOH, then really well with dH₂0. No soap!

- Step 1. Preferably, select single colony of E. coli from fresh LB plate for inoculating a 10 ml 2XYT overnight (O/N) starter culture. Alternatively, streak out frozen glycerol stock of bacterial cells onto LB plate, grow plate O/N, and then select single colony for starter culture. Grow 10 ml starter culture O/N in 37°C shaker (250rpm).
- Step 2. Inoculate 1L of 2XYT media and place culture in 37° shaker. Grow cells and measure OD₆₀₀ every 45min-1hr. When the OD₆₀₀ equals 0.6-0.9 (log phase growth), remove the cells from the shaker and place on ice.

NOTE: It very important to keep the cells at 4°C (or on ice) for the remainder of the procedure.

- Step 3. Split the 1L culture into four equal parts by pouring ~250ml of culture into each chilled 250ml Corning pointed bottle.
- Step 4. Spin (#1) in GPR centrifuge at 4000rpm, 25min at 4°C. (if you chose to use the J6/ JS-4.2 rotor (E. Davidson Lab), use 1L bottles, fill half full, spin 4000rpm, 20min, at 4°C.)

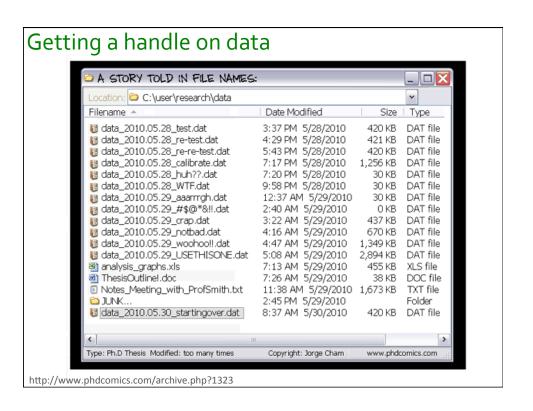
http://www.its.caltech.edu/~bjorker/protocols.html

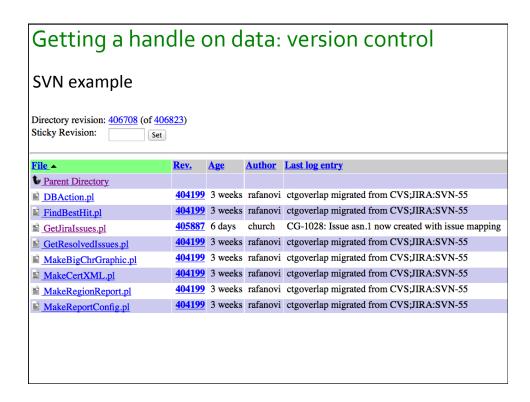


Electronic protocols: workflows

- Often require a Unix environment
- Often run on the command line
- Not always easy to make work







Getting a handle on data: version control

SVN example

Links to HEAD: (view) (download) (as text) (annotate)

Sticky Revision: Set

Revision 405887 - (view) (download) (as text) (annotate) - [select for diffs]

Modified Tue Jul 9 13:19:25 2013 EDT (6 days, 23 hours ago) by church

File length: 46179 byte(s) Diff to previous 405859

CG-1028: Issue asn.1 now created with issue mapping

Revision 405859 - (view) (download) (as text) (annotate) - [select for diffs]

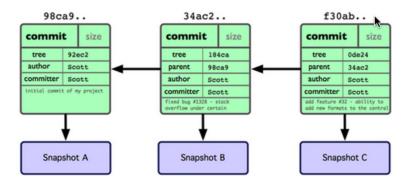
Modified Tue Jul 9 10:54:36 2013 EDT (7 days, 1 hour ago) by church

File length: 45341 byte(s) Diff to previous 405839

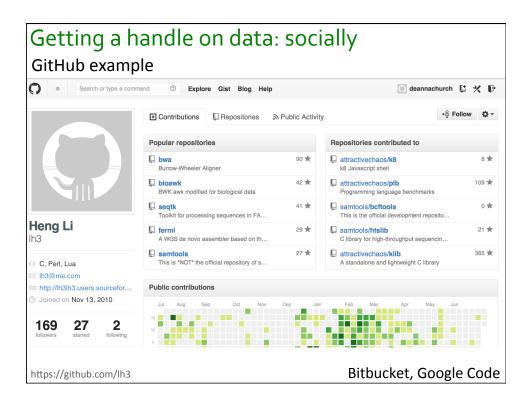
fixed CG-1488

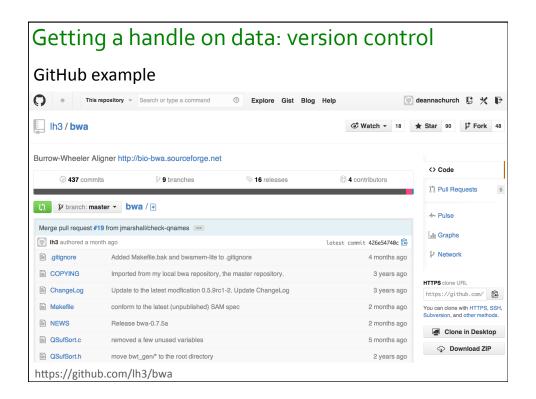
Found bug related to missed mappings

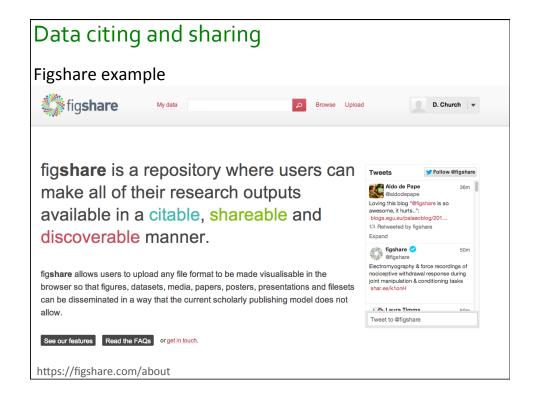
Getting a handle on data: version control Git

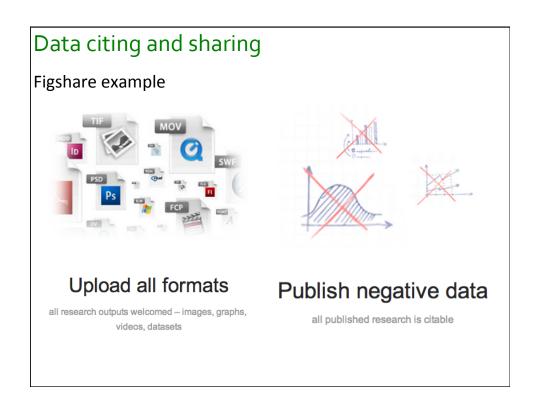


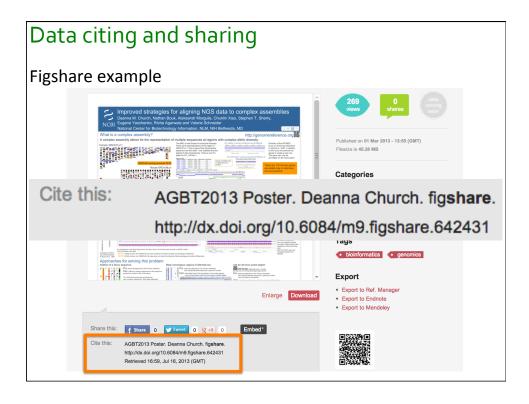
http://nyuccl.org/pages/GitTutorial/

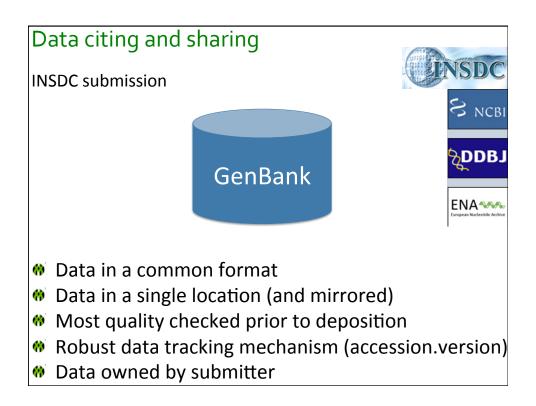












What is next?

- Talk to colleagues about their best practices
 - Solutions may exist in your institution
- Learn to program
 - Software carpentry (http://software-carpentry.org)
 - Codecademy (http://www.codecademy.com)
- Watch this video

http://www.youtube.com/watch?v=N2zK3sAtr-4