Qualitative analysis of introductory biology students' recollections after a completing a CURE or non-CURE laboratory course indicates that CUREs Framework constructs are salient to students

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CONTEXT & RESEARCH QUESTION

RESEARCH QUESTION: We investigated how participating in a CURE curriculum, or its traditional non-CURE counterpart, impacted students' perceptions of their introductory biology laboratory experience at the University of Washington (UW). Prior studies have defined key features of CUREs as: 1) use of scientific practices, 2) discovery and relevance, 3) collaboration, and 4) iteration {Auchincloss, 2014}. We analyzed students' post-course reflections to assess the extent to which CURE and non-CURE students consider these to be prominent features of their labs. Two primary goals of our study were to determine how our CURE curriculum influenced students' affect towards and understanding of scientific research, and to analyze how students' perceptions of their CURE or non-CURE laboratory experiences aligned with the field's established framework

Figure 1 - Interview Participants: 40 volunteers were randomly selected from 4 introductory biology course-offerings spanning 3 quarters (Fall 2018, Winter 2019, and Spring 2019). All enrolled students were invited to participate in focus group interviews after completing the respective course(s). Sixteen focus groups interviews were conducted; data shown reflect 8 interviews analyzed thus far.

WINTER 2019

FALL 2018

10 students interviewed

4 students' data shown

Bio 2 Non-CURE ~550 students enrolled 9 students interviewed **Bio 1 Non-CURE** 5 students' data shown ~1200 students enrolled

> **Bio 1 CURE** ~800 students enrolled 9 students interviewed 4 students' data shown

SPRING 2019

Bio 2 CURE ~350 students enrolled 12 students interviewed 6 students' data shown

Laboratory Modules in Non-CURE and CURE Bio1 & Bio2

Figure 2 – Venn Diagrams of Laboratory Modules: Laboratory modules included in the Non-CURE (orange shading) and CURE (blue shading) versions of Bio1 and Bio2, respectively, are shown. Overlapping modules were retained to meet key learning objectives not fulfilled in the CURE project, which focused on evolution of antibiotic resistance (CURE Bio1) and its molecular basis (CURE Bio2).

- Biodiversity

& Ecosystem

Function (I + II)

Non-CURE Bio 1

- Hypothesis Testing & Experimental Design - Genetic Variation: Meiosis - Intro to Statistics & Data Analysis - Antibiotic Resistance Data Analysis - Phylogenies I: Inference - Phylogenies II: Reading Trees

CURE Bio 1

- Isolate antibiotic-resistant mutants - Discuss Selection, Fitness, Cost of Resistance - Intro to Data Analysis & RStudio - Intro to Collateral Effects & Statistics - Assay Fitness & Resistance levels - Processing our Fitness & Resistance data - Graphing and Interpreting our data

Non-CURE Bio 2

- Research Data Analysis

- Membrane and Cell Transport - Exploring the Central Dogma - Meiosis & Gametogenesis

Enzyme activity - Cellular Respiration - Gene Regulation - DNA Extraction & Fertilization Chick Embryo Development

CURE Bio 2

- Validate & sequence PCR products - DNA sequence analysis - Protein structure analysis

- Connect Genotype, Phenotype, Fitness - Poster Session

Errs/UnexpctRslts

Evo/NatSelec

ExpsBioSys

LrnAbtRsch

ConLbSkl

PrcSkills

LrnExpDsgr

FamLectCont

ThinkSci

FigrCncptThmslf

METHODS

RESEARCH DESIGN: We conducted focus group interviews with students who had recently completed either the CURE or non-CURE version of Part 1 or Part 2 of the introductory biology series at UW. All enrollees were invited to participate, and 40 volunteers were randomly selected (n = 16 groups; mean group-size: 2.5 students, range: 2-4 students). We employed a semistructured interview protocol with 17 questions.

ANALYSES: Interviews were audio recorded, transcribed, and analyzed by a team of 6 qualitative data analysts using grounded theory to define emergent codes and themes. At least 2 coders analyzed each of the interview transcripts; all 6 analysts discussed their individuallygenerated codes to develop consensus codes. Tests of inter-rater reliability in applying consensus codes are in progress.

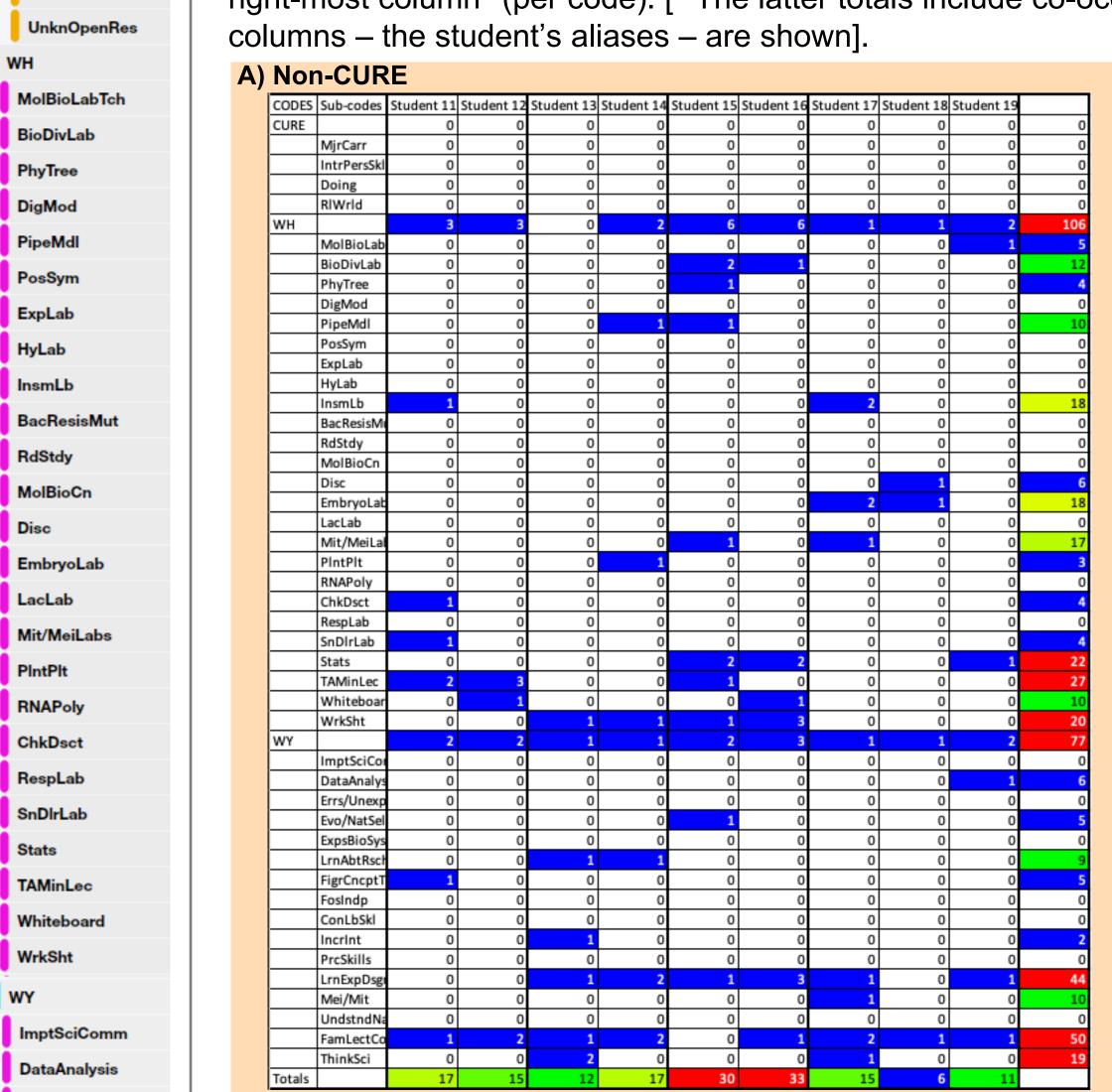
RESULTS

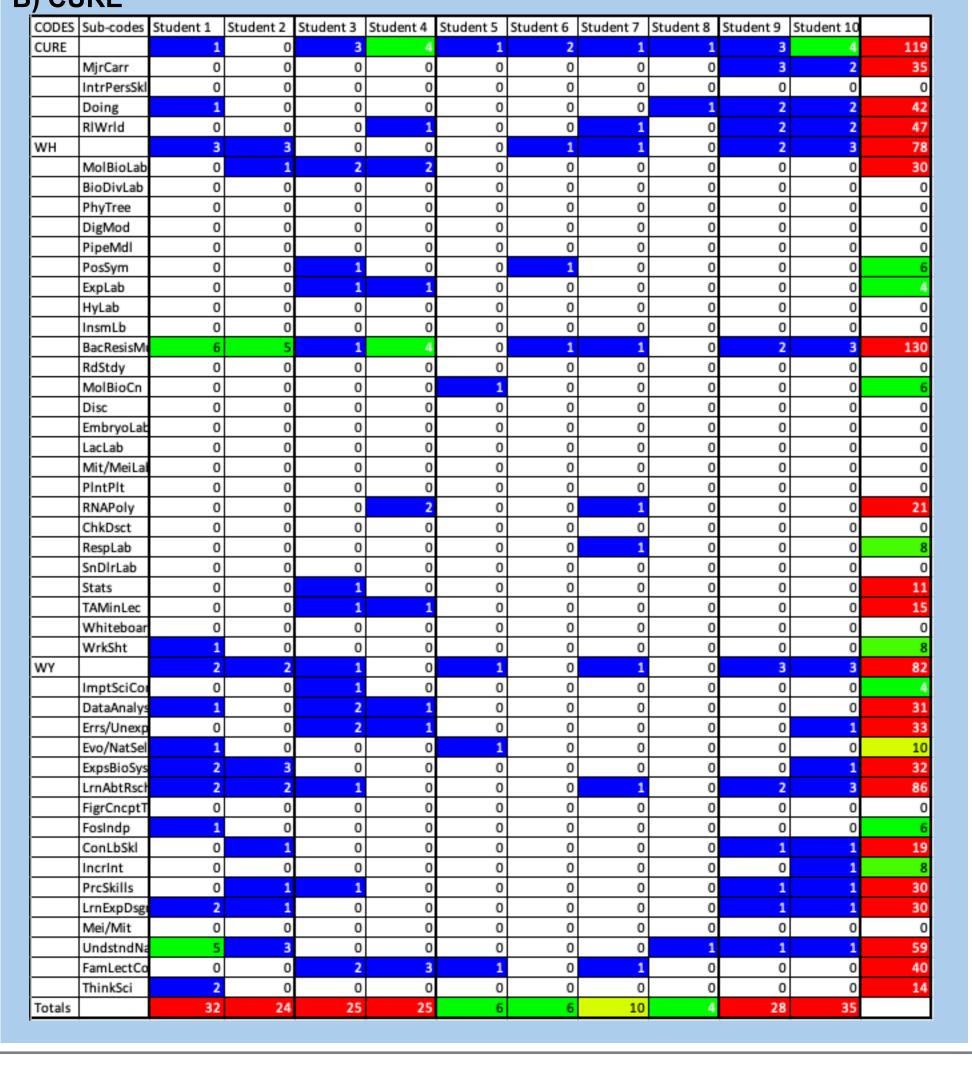
Here, we present preliminary qualitative analyses of students' responses to Question #3: "What did you do in labs for this course and why did you do it?", after participating in a CURE or Non-CURE series of laboratory modules (described in Figure 2).

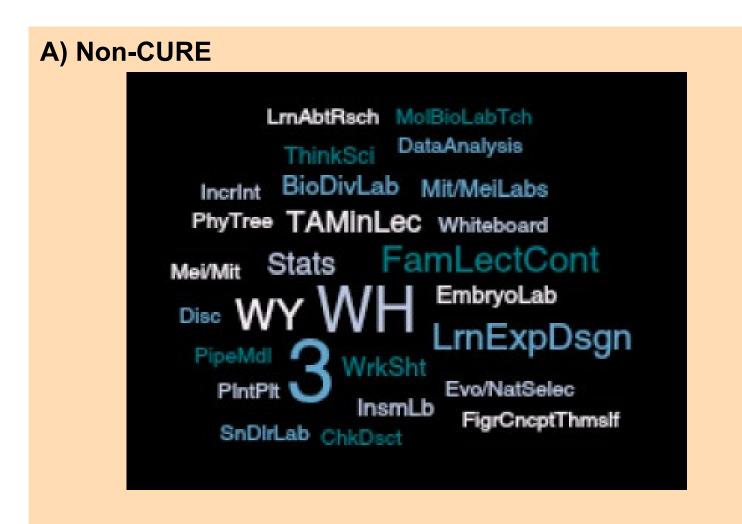
Figure 3 - Tree of qualitative codes applied to analyze Non-CURE and CURE students' responses to Question #3: Emergent codes were generated by 6 qualitative data analysts (raters) who read and memo-ed transcripts, and discussed the meaning of participants' statements to arrive at consensus codes, shown below-left. The raters were blind to whether participants were in a Non-CURE or CURE course.

- Parent-level codes are denoted with teal-blue bars; child-codes with pink, and sub-child-codes with orange. Parent-level codes were "CURE", What ("WH"), and Why ("WY").
- Child-codes under "CURE" include alignment with undergraduate major and/or career objective (MjrCarr), Interpersonal Skills (IntrPersSkll), doing research (Doing), real-world applications (RIWrld), Contributing to a professional research-lab's research (ContrReal Rsch), and obtaining unknown/open-ended results (UnkwnOpenRes).
- Child-codes under "What" (WH) mainly refer to topics of specific lab modules (Figure 2), with the exception of activities, such as Discussion (Disc), experience with lab-work and techniques (ExpLab), reading studies (RdStdy), mini-lectures by the teaching assistant (TAMinLec), teamwork on dry-erase boards (Whiteboard), and worksheets (WrkSht); and one experimental outcome: RNA polymerase change due to selection of resistance-conferring mutation(s) (RNAPoly).
- Child-codes under "Why" (WY) range in focus from providing direct support for lecture objectives [e.g., learn Mitosis/Meiosis (Me/Mit), familiarize with lecture content (FamLectCont)], to transferrable skills, science process skills, and broader relevance of the labs' objectives.

Figure 4 – Code co-occurrence plots for participants who completed a Non-CURE or CURE series of lab modules: Codes that 2 independent raters assigned to describe statements made by participants who completed a Non-CURE (Panel A) or CURE (Panel B) CURE series of lab modules are shown, below-right. The code-tree (described in Figure 3) appears in the left-most 2 columns, and student-MjrCarr aliases for the participants in each treatment-group are noted at the top of each column. The frequency with which codes were IntrPersSkll assigned is color-coded in a heat-map: low, intermediate and high frequencies are indicated with blue, green, and red shading, RIWrld respectively; absence of color indicates no code was assigned. Totals are reported in the bottom-most row (per participant), and the ContrRealRsch right-most column* (per code). [* The latter totals include co-occurrence with all other codes in the table, although only a subset of columns - the student's aliases - are shown]. A) Non-CURE







B) CURE ExpLab ExpsBioSys Errs/UnexpctRsIts ThinkSci ProSkills FamLectCont TAMinLec RIWrld UndstndNatSelec LrnAbtRsch

Figure 5 – Word-cloud of codes applied to statements from interview participants who completed a Non-CURE or CURE series of lab modules: Word-clouds visualizations of codes that 2 independent raters assigned to describe statements made by participants who completed a Non-CURE (Panel A) or CURE (Panel B) series of lab modules are shown above. These word-clouds provide an alternative view of the code-frequency data depicted in Figure 4.

RESULTS SUMMARY

INTERPRETATIONS: Our preliminary analyses of students' responses to "What did you do in your labs and why?" in 8 transcripts indicate that CURE students mention parameters not found in non-CURE students' discourse. For example, whereas non-CURE students emphasized content-delivery methods, mastery of discrete skills, and reinforcement of lecture concepts; CURE students cited encountering unexpected results (opportunities for iteration) and learning about professional research by conducting their own research (scientific practices), which addressed questions that could help inform public health measures and clinical practices (discovery and relevance).

CONTRIBUTION: In this study, we found that students' salient reflections on their introductory biology laboratory experience align with outcomes predicted by the CUREs framework to a greater extent if they participated in a CURE than if they participated in a non-CURE laboratory experience. Therefore, our data suggests that students recognize the benefits of CUREs, and that students' perceptions of those benefits are consistent with CURE experts' views. Thus, our findings may help encourage faculty to implement CURE curricula by reinforcing – in the words of students – that the outcomes merit investing extra effort in leading CUREs.

FUTURE DIRECTIONS

Next steps for this project include:

- Analyze 8 additional interview transcripts (21 participants.)
- 2. Complete qualitative analysis for other interview questions within the 16 transcripts, such as:
- "What did you think about how science is done before taking this course's labs? If your understanding has changed in any way, how did it change?"
- "Did you encounter any unexpected results? If so, what was that like?"
- 3. Troubleshoot R source-code to:
- calculate Cohen's kappa (for inter-rater reliability),
- perform statistical analyses of significance, and
- graph percentage of interviewees in Non-CURE and CURE lab series who made a statement that elicited each code.
- 4. Seeking collaborators:
- To analyze Spring '21 Scientist Spotlights data, &/or R code.

ACKNOWLEDGEMENTS

We thank Scott Freeman, Sara Brownell, Alison Crowe, Mary Pat Wenderoth, Katie J. Dickinson, Ben Kerr, Ben Wiggins, and the Kerr and BERG Labs for helpful discussions. We thank UW Intro Bio CURE alumni (especially focus group interview volunteers and Peer Facilitators).

This work was supported by an HHMI (STEM-Dawgs) grant to Scott Freeman. JM would like to thank members of the UW Department of Biology for past and future collaborations, and colleagues in the Department of Biological Sciences at California State University, Sacramento; in particular, Kelly K. McDonald.



REFERENCES

1) Auchincloss, L. C., Laursen, S. L., Branchaw, J. L., Eagan, K. Graham, M., Hanauer, D. I., Lawrie, G., McLinn, C. M., Pelaez, N. Rowland, S., Towns, M., Trautmann, N. M., Varma-Nelson, P., Weston, T. J., & Dolan, E. L. (2014). Assessment of course-based undergraduate research experiences: a meeting report. CBE life sciences education, 13(1), 29-40. https://doi.org/10.1187/cbe.14-01Comments during Zoom poster session from visitors:

Met Supriya, from SEB's group at ASU (PD hired for quanitative; gave me GREAT advice on when to try quant or stay 100% qualitative – Follow up via email

More notes on "June 2020" page of Ram Dass notebook, from: Supriya, and from Jon Herron

Chat:

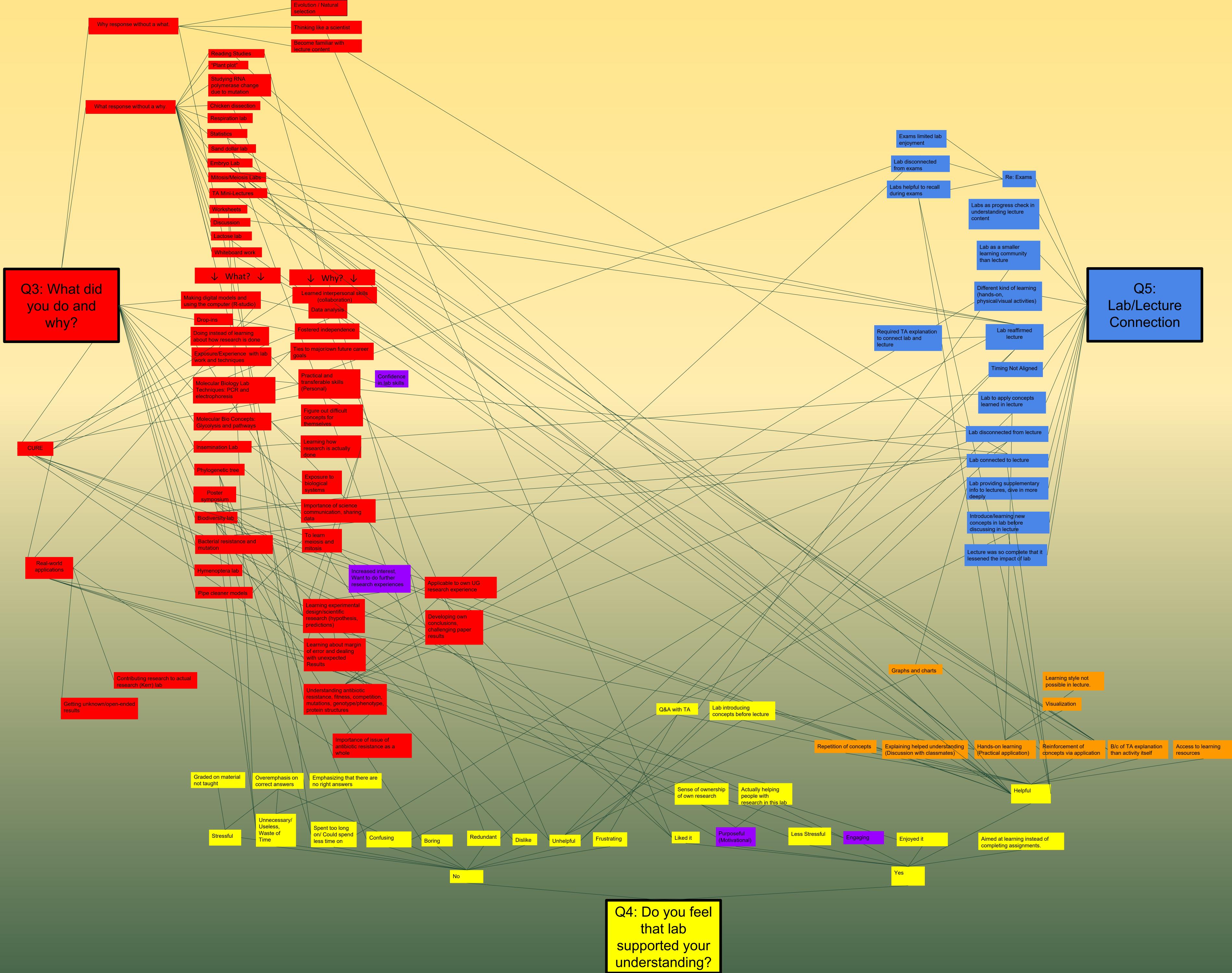
2:43:06 From Becca Price (she/hers): Joya - I've looked through your poster, and I like the plan to start lumping into bigger themes. Happy to talk in more detail later on. So nice to see you!

12:43:19 From Becca Price (she/hers): I've got a ton of posters to visit today, so I'm going to move on.

13:00:30 From Bailey Rowe (she/her): I'm going to run Joya. Your poster was great!! Super interested in how you do quantitative stuff going forward.

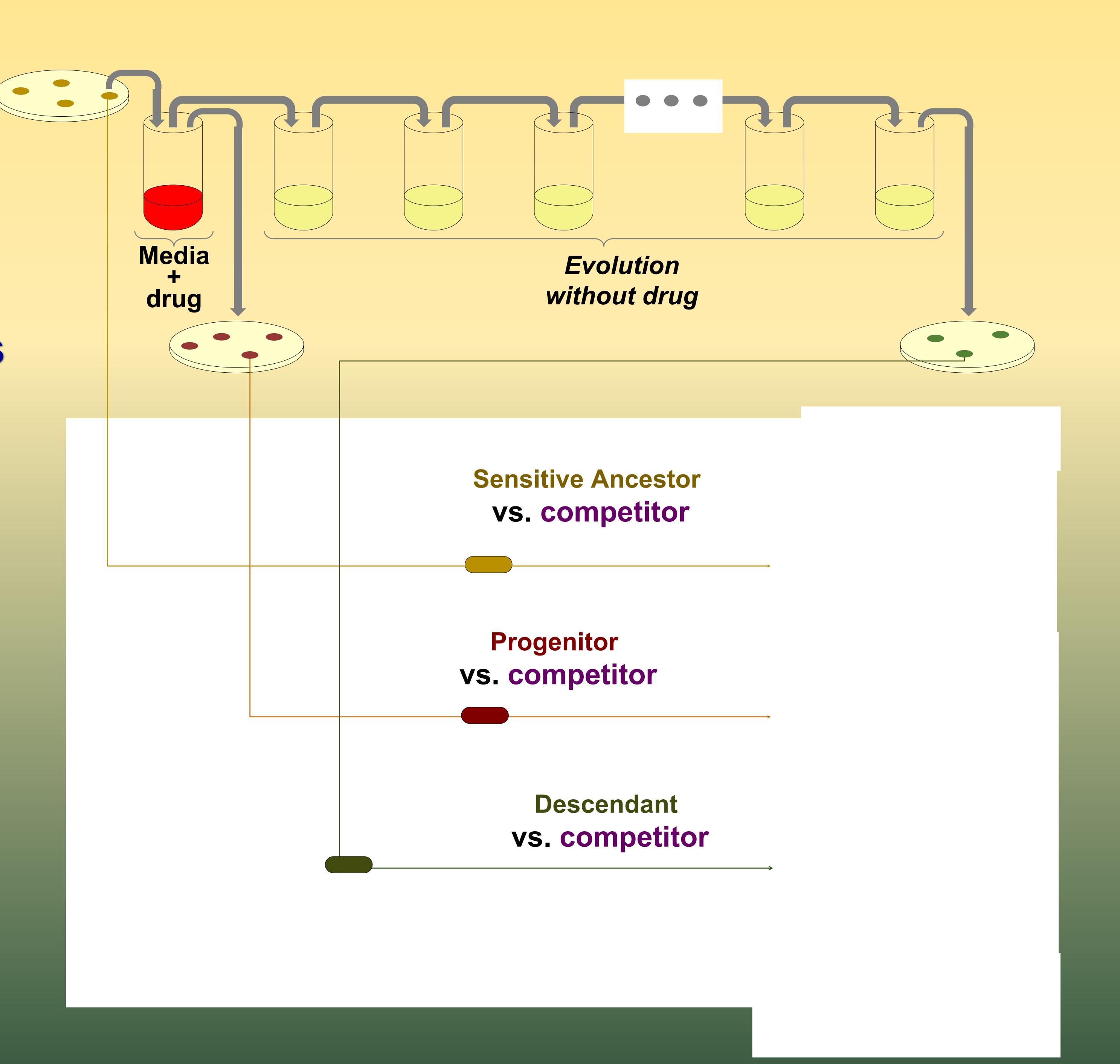
14:02:51 From Kelly McDonald : Nice work, Joya. I have to run to a Buddy meeting.

14:03:18 From Heather Fletcher : Thanks, Joya! :)



BIOL 1: Experimental

evolution & fitness competition assay





Molecular basis for fitness differences

