

# Lessons from a (Mostly) Successful Flip

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Active Learning Seminar

October 13, 2016



## Teaching Team



Susie Honig

- Anne Warner (TA)
- Chandra Goetsch (TA)
- Hailun Wan (LA)
- Rebecca Rook (LA)
- Kyler Asato (LA)
- Raymond Tan (MSI)

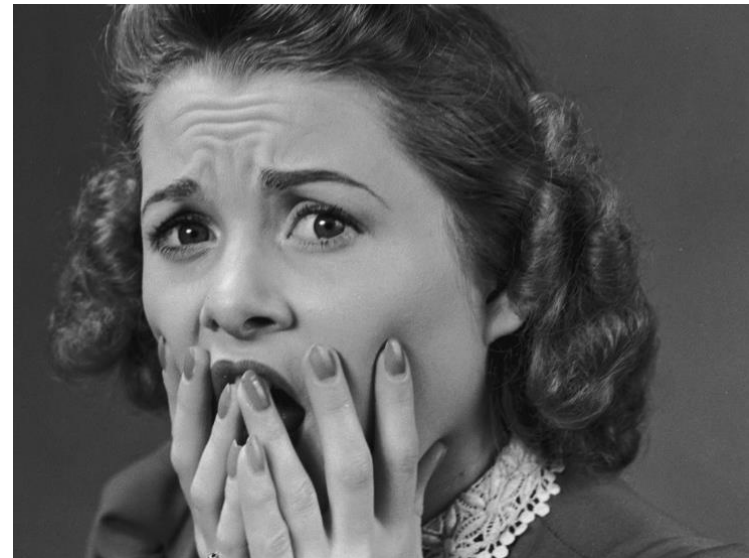
## AL/HHMI Team

- Lisa Hunter (ISEE)
- Manny Ares
- Ian Marcus
- Tammy Bye

***TEAM  
EFFORT!***

*With your neighbor*

*What are your biggest worries about flipping a class/incorporating active learning?*



# *What were our major concerns at the beginning?*

- Will students do the work needed outside of class?
- How much content are we going to have to cut?
- Will students engage or will it be like a whole quarter of flopping at open mic night?
- Will our evaluations suffer?
- How much extra time/work is this going to take?
- Will they really do better?



1

Planning the flip

2

Flipping: What did it look like?

3

Were we successful? How do we know?

4

What would we change?

5

Taking it to the masses!

# *What did we consider active learning for our purposes?*

Active Learning. The process of having students engage in some activity that forces them to reflect upon ideas and how they are using those ideas. Requiring students to regularly assess their own degree of understanding and skill at handling concepts or problems in a particular discipline. The attainment of knowledge by participating or contributing. The process of keeping students mentally, and often physically, active in their learning through activities that involve them in gathering information, thinking, and problem solving.

*- Collins and O'Brian, 2006*

# *What did we consider active learning for our purposes?*

- Active lecture (clickers, pair shares)
- Working on problems alone or in small groups
- Class discussions
- Worksheets
- Working on questions/problems at poster stations
- Class demonstrations
- Fish bowl exercises
- Peer Teaching
  
- Group Projects (quarter long)

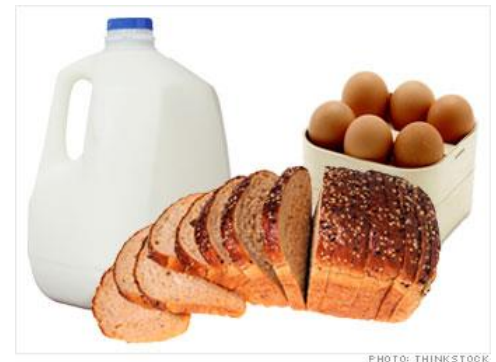
# *Planning to Flip*

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- Started with previous syllabus + info from “campfires”
- Quasi-Backwards design
- Went class by class and designed original activities, used online activities, relied on some “staples”

## *AL “Staples”*

- Active lecture
- Postering
- Worksheets paired with discussion






Instructor Notes  
Tuesday April 5 2016

**Class Schedule**

- 15 minutes: Students fill in "mouth to colon" drawing
- 30 minutes: Evidence and models activity
- 10 minutes: The 'real' models
- 10 minutes: Jigsaw
- 30 minutes: Leptin graphing and diabetes active lecture
- 10 minutes: Synthesis (What did we learn and assign HW)

**Class schedule (usually thrown out the window)**



**Before-Class Assignments**

- Read Chapter 51
- Online Lecture: The digestive system parts I and II: Crash Course/Khan Academy

**What students should have done before class**




**Learning Outcomes**

*Digestive System*

- Be able to describe the major organs of the digestive system including what type of fuels are digested and absorbed and where this occurs.
- Understand the difference between digestion and absorption
  - Students should be able to describe that the breakdown from polymers to monomers is digestion and they should predict the method of transport of different molecules based on polarity.
- Understand that digestion is extracellular
- Be able to describe mechanisms of transport of different molecules to cross epithelial cell
- Understand that mechanism of absorption is based on molecular characteristics.

**Learning outcomes based on what had been done in previous classes**



*Hormonal Control*

- Be able to describe hormonal regulation of digestion at the level of Figure 15.17 in Sadava.
- Draw a graph of blood sugar, insulin, and glucagon over time and before and after a meal.
- Describe the difference between Type I and Type II diabetes
- Draw a graph of leptin vs. body mass and predict the correct graph for humans or mice that lack the leptin gene

**Materials**

- Student Guide handouts for Evidence-based models
- Poster paper
- Markers

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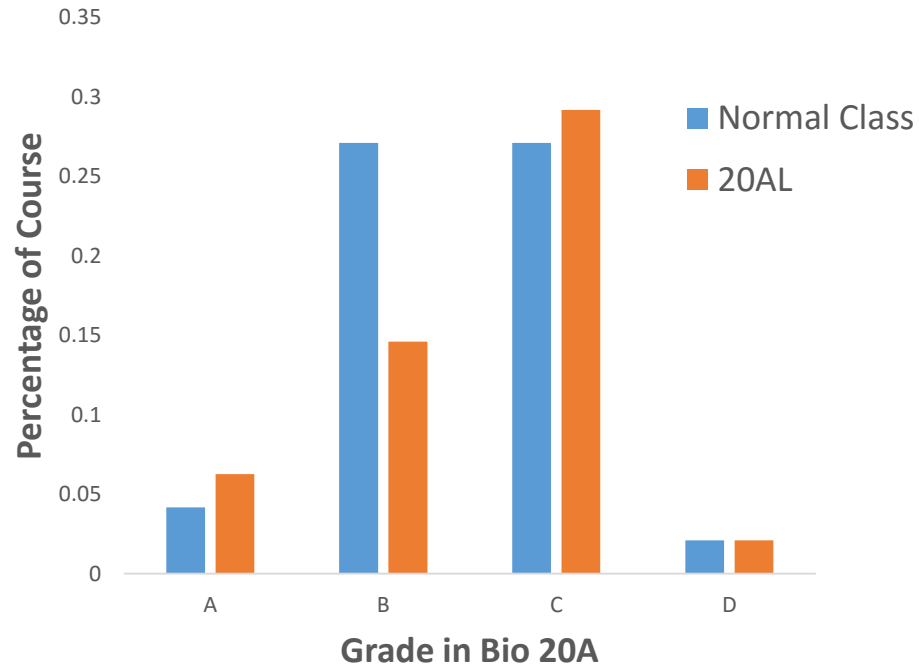
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Taking it to the masses!

# The Students

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Grade coming into bioe Bio 20B



58 students in course  
49 responded to this survey

Full data from institutional  
research is pending

# The first day

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## Equity & Inclusion

### Feedback from students

- *What do I want my instructors to know?*
- *What I am excited to learn about?*



Some real responses:

- I am working 20 hours a week to pay for school
- This is the second time I am taking this class and I am going to work extra hard to pass
- I am scared of talking in class but I am still engaged
- I am scared of math
- I want to be a researcher
- I LOVE biology
- I want to learn about the nervous system
- I have to commute 2 hours to get here
- English is my second language
- You should know that this is not my only class 😊

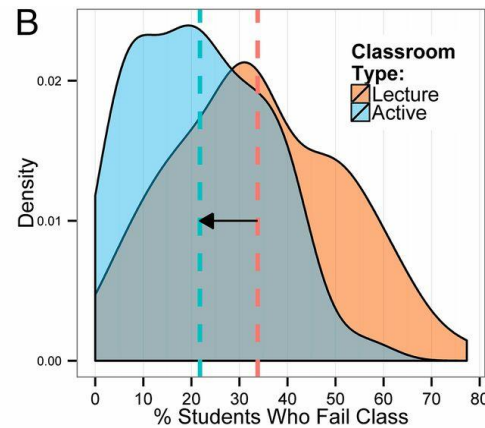
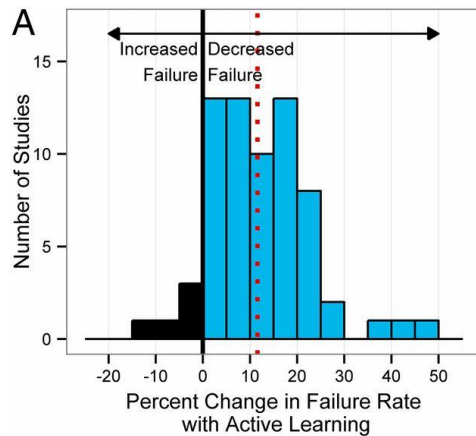
# The first day

Feedback from students

- What do I want my instructors to know?
- What I am excited to learn about?



Make the evidence based case for AL



Scott Freeman et al. PNAS

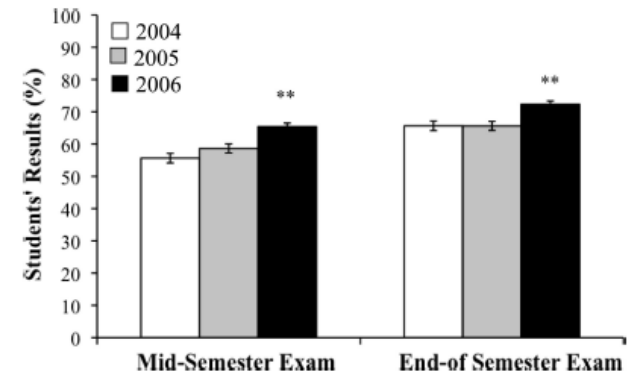


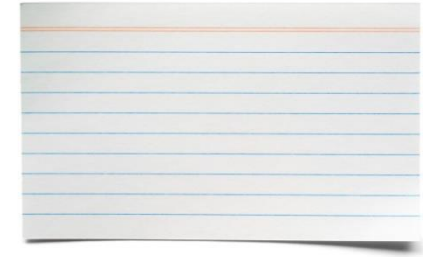
Fig. 5. Comparison among 2004 ( $n = 127$ ), 2005 ( $n = 137$ ), and 2006 ( $n = 169$ ) final examination results for course 536-211 Physiology: Control of Body Function. PRS voting on questions in lectures was not first introduced until 2006. Results are expressed as means  $\pm$  SE. \*\* $P < 0.01$  vs. 2004 and 2005 results (by independent  $t$ -test).

Gauci et al., 2009

# The first day

*Feedback from students*

- What do I want my instructors to know?*
- What I am excited to learn about?*



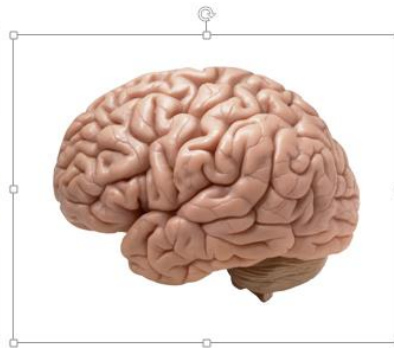
*Make the evidence based case for AL*

## Pair Share on Growth Mindset

With your neighbor...

Is intelligence fixed or flexible?

Deep down...what do you REALLY think?



## What Kind of Mindset Do You Have?



I can learn anything I want to.  
When I'm frustrated, I persevere.  
I want to challenge myself.  
When I fail, I learn.  
Tell me I try hard.  
If you succeed, I'm inspired.  
My effort and attitude determine everything.



I'm either good at it, or I'm not.  
When I'm frustrated, I give up.  
I don't like to be challenged.  
When I fail, I'm no good.  
Tell me I'm smart.  
If you succeed, I feel threatened.  
My abilities determine everything.

# The first day

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*Feedback from students*

- *What do I want my instructors to know?*
- *What I am excited to learn about?*



*Make the evidence based case for AL*

*Pair Share on Growth Mindset*

*Conveyed confidence in their ability to get an A but more importantly deeply learn*

**Equity & Inclusion**

*Course organization, stipulated that things may change, welcomed feedback, told them how we would let them know what they are responsible for knowing.*

# Examples of Activities

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## Staples

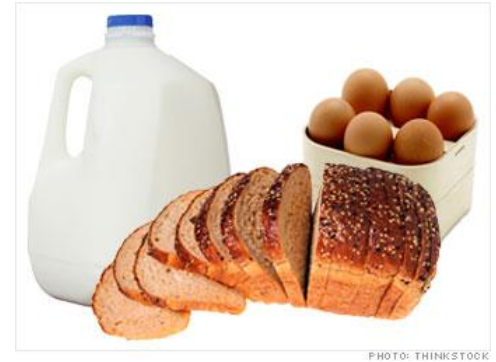
1) *Active Lecture* with clicker questions & “with your neighbor”

*Generally have two types:*

- Checking understanding
- Challenge questions

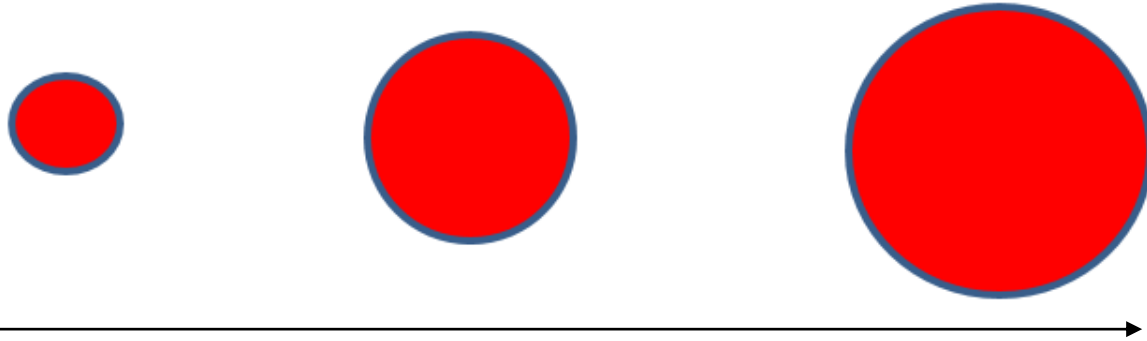
*Skill focused*

- Draw a graph/interpret graph
- Critical thinking
- How would you test X or Y?

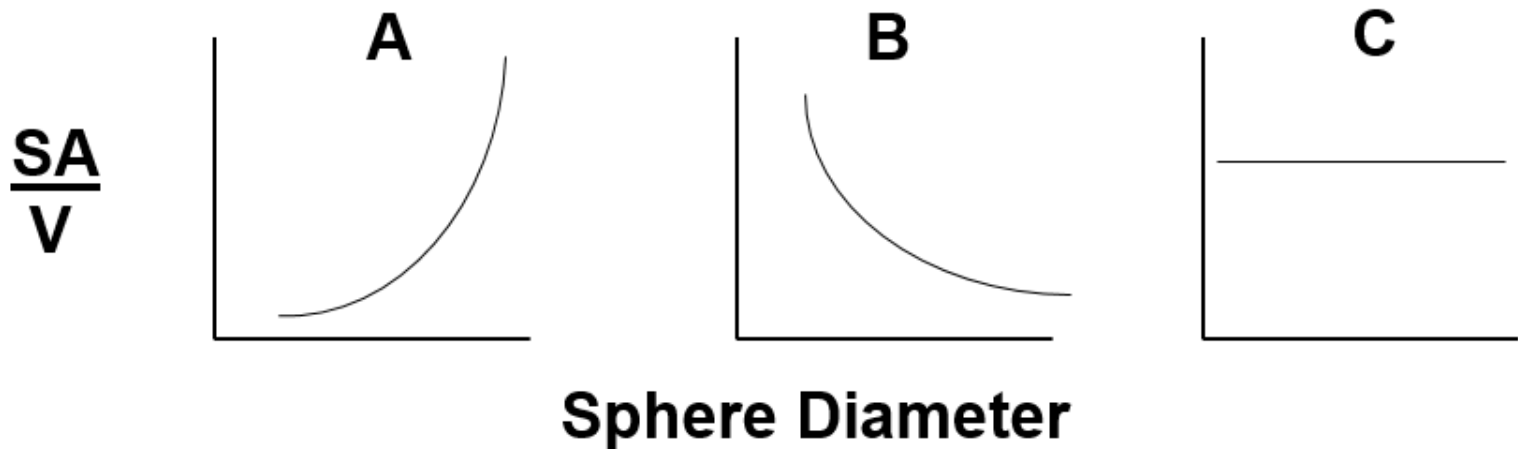




# Examples of Activities



The graph that most accurately describes the surface area to volume ratio of the spheres above is:



# Examples of Activities

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## Staples

1) Active Lecture with clicker questions & “*with your neighbor*”

## 2) *Postering*

- Facilitates peer teaching
- Encourages metacognition
- Encourages revision
- Helps instructor see student’s thinking
- Helpful for:

*Processes*

*Learning a system*

*Working out critical thinking questions*

# Examples of Activities

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## Staples

1) Active Lecture with clicker questions & “*with your neighbor*”

2) Postering

### 3) *Worksheets*

- Targeted typically difficult topics
- Aimed at giving them practice
- Basic to challenge questions
- Always went over in class

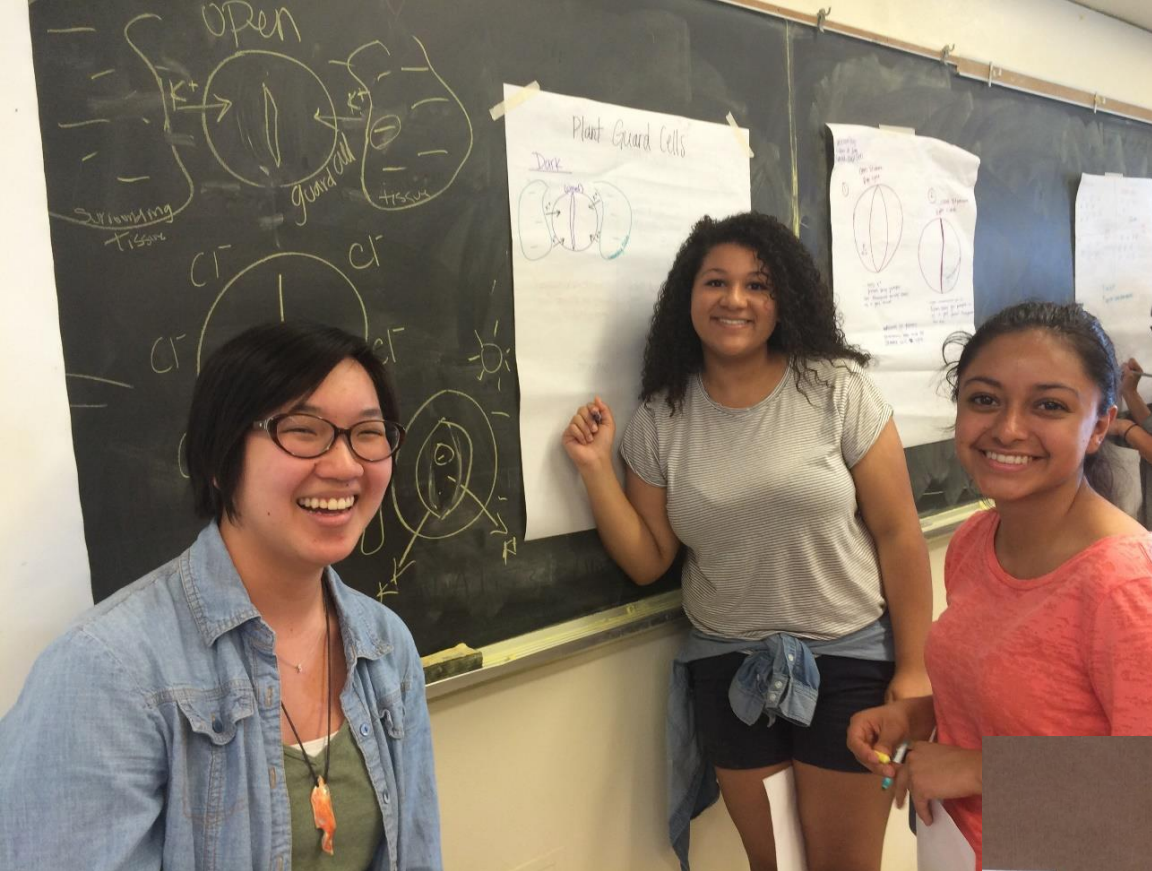
# *Examples of Activities*

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## Additional activities

- Evidence and Modeling x 3
- Graph Speed Data-ing
- Act out an action potential
- Human demonstration of tree growth
- Human demonstration of water transport
  
- Rap Genius
  - *Quarter long project focused on reading and interpreting scientific literature using rap genius annotation website*
  - *Culminated in a mock “conference”*

**Equity & Inclusion**



*Given evidence statements*

*Asked to build a model that fits the data*

- *Guard cell function*
- *Transport of nutrients across intestine*
- *Effects of neurotoxins*

*Evidence & Modeling of how guard cells open & close*



FILTER BY All Annotations

Add a comment

amyloidogenic processing pathways [5]. This pathway is initiated when APP undergoes proteolytic cleavage by  $\beta$ -secretase, encoded by the *BACE* gene. This cleavage produces a soluble extracellular/luminal fragment of APP (sAPP $\beta$ ) and a membrane spanning C-terminal fragment ( $\beta$ CTF/C99). The  $\gamma$ -secretase complex then cleaves  $\beta$ CTF to produce A $\beta$  peptides and the APP intracellular domain (AICD) [5]. A $\beta$  peptides of a variety of lengths are produced but A $\beta_{40}$  and A $\beta_{42}$  are the major isoforms produced in the central nervous system (CNS). Compared to A $\beta_{40}$ , A $\beta_{42}$  is more prone to oligomerization and has been shown to be more neurotoxic [6].

APP also undergoes an alternative proteolytic processing pathway termed the non-amyloidogenic pathway. In this pathway,  $\alpha$ -secretase initially cleaves APP, rather than  $\beta$ -secretase, to produce a soluble extracellular/luminal fragment of APP (sAPP $\alpha$ ) and a membrane spanning C-terminal fragment ( $\alpha$ CTF/C83). Again, the  $\gamma$ -secretase complex then cleaves  $\alpha$ CTF to produce the P3 peptide and AICD [5].

APP proteolysis is an important step towards development of AD. Therefore, it is important to identify genes and pharmaceuticals that modulate APP metabolism and A $\beta$  production and clearance. Developing *in vivo* disease models has proven crucial to illuminating disease mechanisms, since *in vitro* studies do not always represent the natural physiological complexity of the tissue and/or organism. In particular, the fruit fly, *Drosophila melanogaster*, has been tremendously important and influential in furthering our understanding of the mechanisms of many forms of neurodegenerative diseases, including AD [7], [8], [9], [10], [11].

*Drosophila* endogenously express orthologues to the human APP [12],  $\alpha$ -secretase [13], [14], and  $\gamma$ -secretase [15], [16], [17], [18]. Recently, a functional *Drosophila* homolog of the BACE ( $\beta$ -secretase) family of proteins has also been identified [19]. Though the *Drosophila* homolog to human APP, *Appl*, does not contain significant sequence similarity within the A $\beta$  region of human APP [12], there is recent evidence suggesting

In this pathway,  $\alpha$ -secretase initially cleaves APP, rather than  $\beta$ -secretase, to produce a soluble extracellular/luminal fragment of APP (sAPP $\alpha$ ) and a membrane spanning C-terminal fragment ( $\alpha$ CTF/C83). Again, the  $\gamma$ -secretase complex then cleaves  $\alpha$ CTF to produce the P3 peptide and AICD



sorlium

4 months

Instead of the beta-secretase cleaving the APP a different protein does it, which is alpha-secretase. This produces similar protein fragments as the beta-secretase that also undergo cleavage again by  $\gamma$ -secretase, but it is implied that they are less harmful than AB40 and AB42 which are produced by the initial cleavage by beta-secretase.

report abuse

Upvote



Add a comment

A $\beta$  production and clearance

bio20b\_wed\_11\_group8

5 mo

# “Mock Conference” at end of Course

- *Students presented posters*
- *Authors of papers discussed careers & role of failure in science*

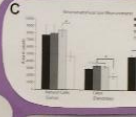




# Characterization of a *Drosophila* Alzheimer's Disease Model: Pharmacological Rescue of Cognitive Defects

## Evidence

The evidence shows that ELAV<sup>1</sup> RNAi rescues the memory deficit in the *Drosophila* Alzheimer's disease model. ELAV<sup>1</sup> RNAi also rescues the cognitive deficit in the *Drosophila* Alzheimer's disease model. ELAV<sup>1</sup> RNAi also rescues the cognitive deficit in the *Drosophila* Alzheimer's disease model.



## Reasoning



## Future Directions

The next step in research for Alzheimer's is likely to be to identify the specific pathways that are involved in the cognitive deficits seen in the *Drosophila* Alzheimer's disease model. This will allow us to identify the specific pathways that are involved in the cognitive deficits seen in the *Drosophila* Alzheimer's disease model.

## Questions

1. How does ELAV<sup>1</sup> RNAi rescue the cognitive deficit in the *Drosophila* Alzheimer's disease model?
2. What are the specific pathways that are involved in the cognitive deficits seen in the *Drosophila* Alzheimer's disease model?
3. How does ELAV<sup>1</sup> RNAi rescue the cognitive deficit in the *Drosophila* Alzheimer's disease model?

# Running, Swimming and Diving Modifies Neuroprotecting Globins in the Mammalian Brain



Authors: Williams TM, Zavanelli M, Miller MA, Goldberg RA, Morledge M, Casper D, Pabst DA, McLellan W, Cantin LP, Kiger DS

MAMMALIAN HAVE UNIQUE ADAPTATIONS COMPARED TO TERRESTRIAL MAMMALS BUT WHAT MAKES THESE ANIMALS INTERESTING FROM A LACK OF OXYGEN. MAMMALIAN ARE ABLE TO PROTECT THEIR BRAIN FROM ISCHEMIA UNDER VARIOUS CIRCUMSTANCES. MAMMALIAN ARE ABLE TO PROTECT THEIR BRAIN FROM ISCHEMIA UNDER VARIOUS CIRCUMSTANCES. MAMMALIAN ARE ABLE TO PROTECT THEIR BRAIN FROM ISCHEMIA UNDER VARIOUS CIRCUMSTANCES.



Figure 1: Concentration of Hb & HMO in the brain over time.



High concentration of Hb in Marine Mammal - High amount of Hb allows oxygen to move more easily under the blood oxygen limited pressure.

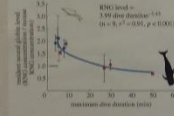


Figure 2: Marine mammals have higher concentration of Hb & HMO in the brain compared to terrestrial animals.

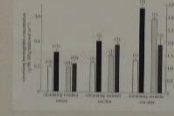


Figure 3: Marine mammals have higher concentration of Hb & HMO in the brain compared to terrestrial animals.

Future Directions: Other areas could be done on other deep sleep animals. Prospects for marine mammals are very exciting. Other possible subjects are animals that typically live at very high altitudes. Can't remember exactly. Measure membrane oxygen saturation before they start a dive and after they finish, and before they get back. Using FMO Enzymes. Can't remember exactly for these mammals. We think the parts of the body that have lower affinity for oxygen than normal for mammals. The main candidates for research in marine mammals are: Applications for hypoxemia (low blood oxygen). Which could be derived from fetal blood vessel oxygen. Applications to diseases that reduce the oxygen levels, such as sickle cell.

- ### Questions for the authors
1. In humans we don't use 100% of the oxygen extracted in our blood. Do these mammals breathe more than 100% of the oxygen they O previously stored upon breathing?
  2. Are the parts of the mammalian brain where hemoglobin is less concentrated? If so what do these parts contain?
  3. What would an oxygen-hemoglobin curve for these mammals look like? Can we measure that the affinity for O2 is lower than terrestrial mammals?
  4. Are there any differences in the structure of hemoglobin, myoglobin, or myoglobin in deep sleep animals or during a quantitative difference?





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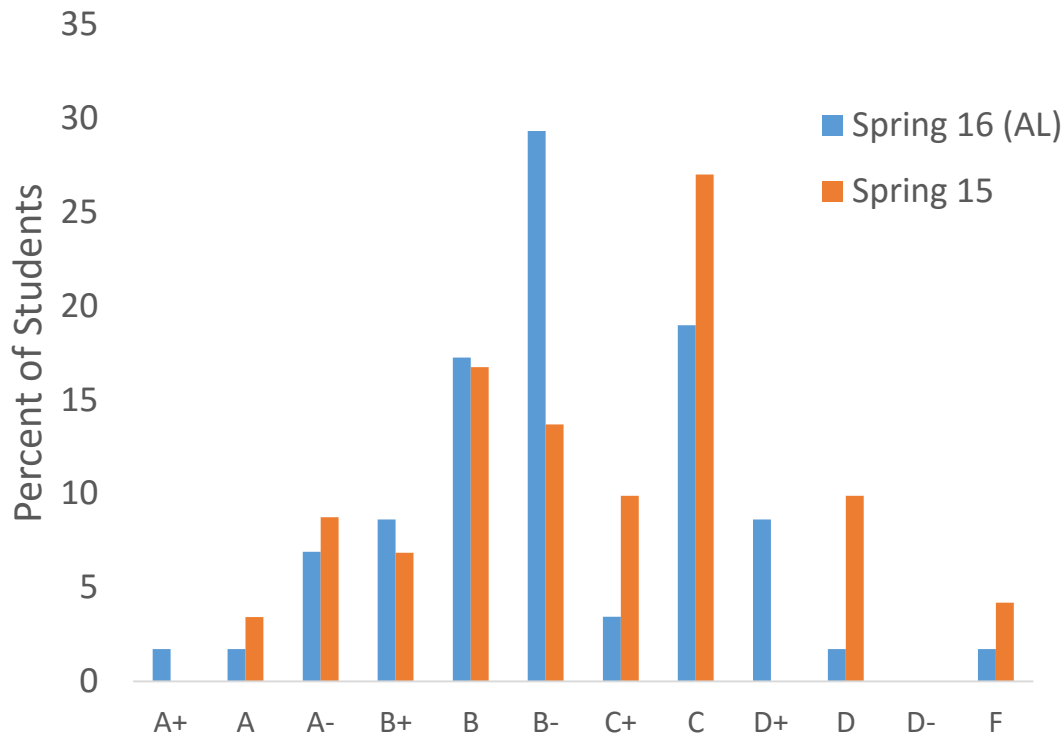
Taking it to the masses!

# Were we successful?

## Grades

Data sources:

- 1) grades
- 2) end of quarter survey



- Failure rate – nearly identical
- Moved C students up to B- ??

*Caveat: Very different sample sizes....*

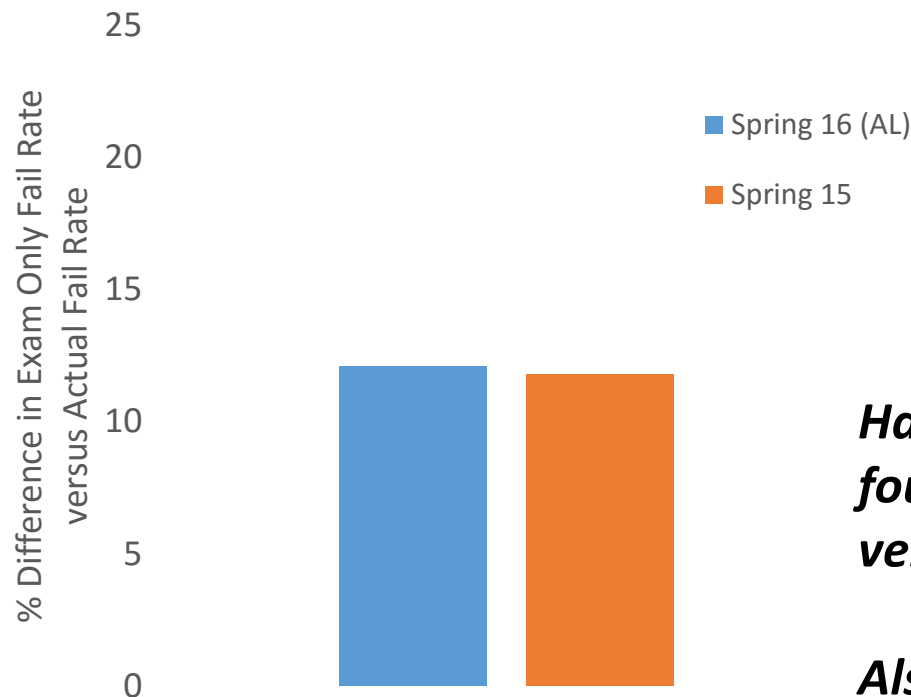
***Do active learning exercise just result in grade inflation because of more non-exam points?***

# Were we successful?

## Grades

Data sources:

- 1) grades
- 2) end of quarter survey



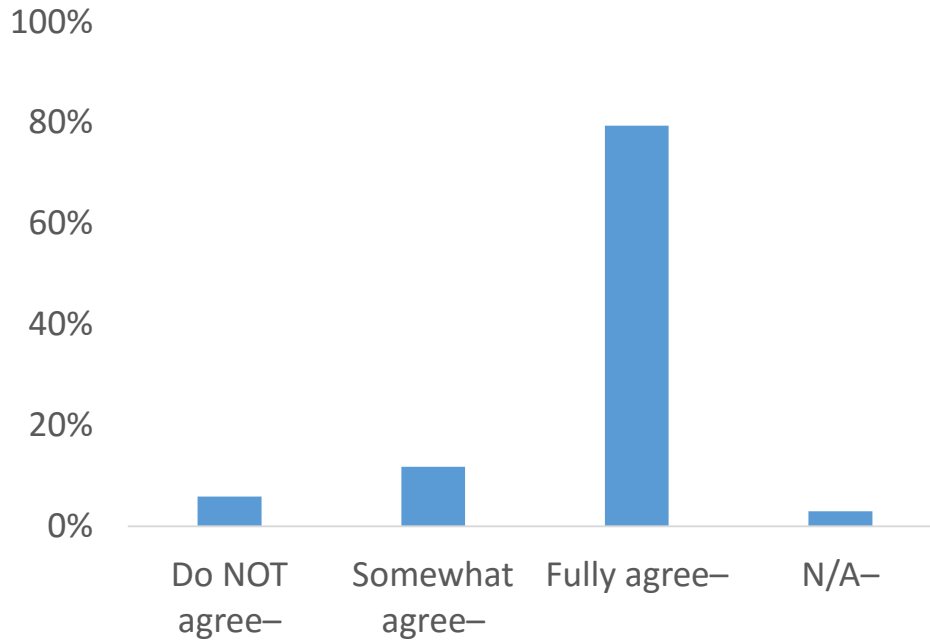
- Virtually identical rate of students would have failed based on exam scores only

***Haak and Freeman, 2011 (Science) found the same for controlled study with very large sample size.***

***Also found exams tended to get harder in AL courses.***

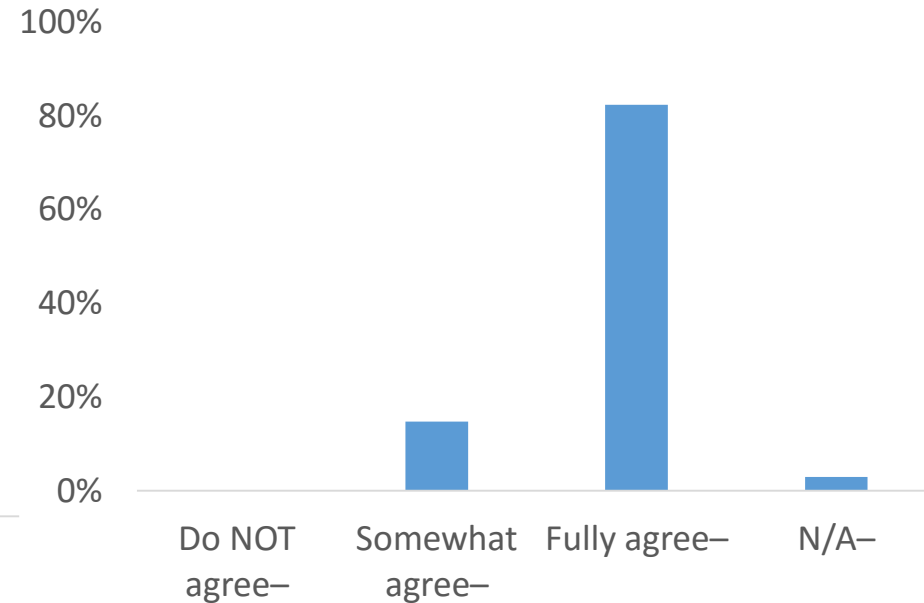
# Were we successful?

## STEM Identity & Equity and Inclusion



**I felt that the course helped me feel more like a scientist compared to when I started the course.**

## End of quarter survey

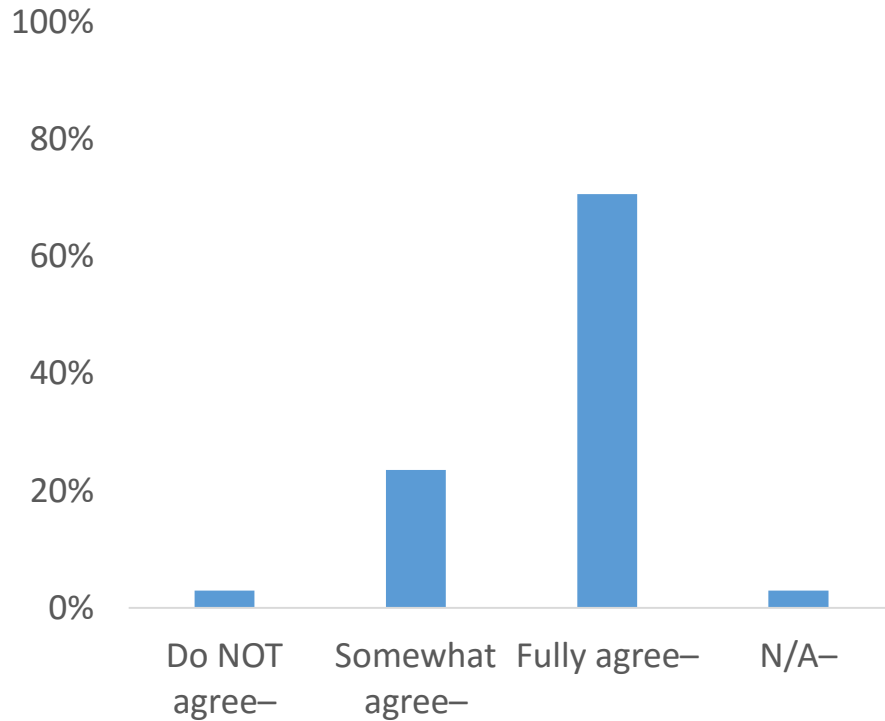


**I felt the course was equitable & Inclusive**

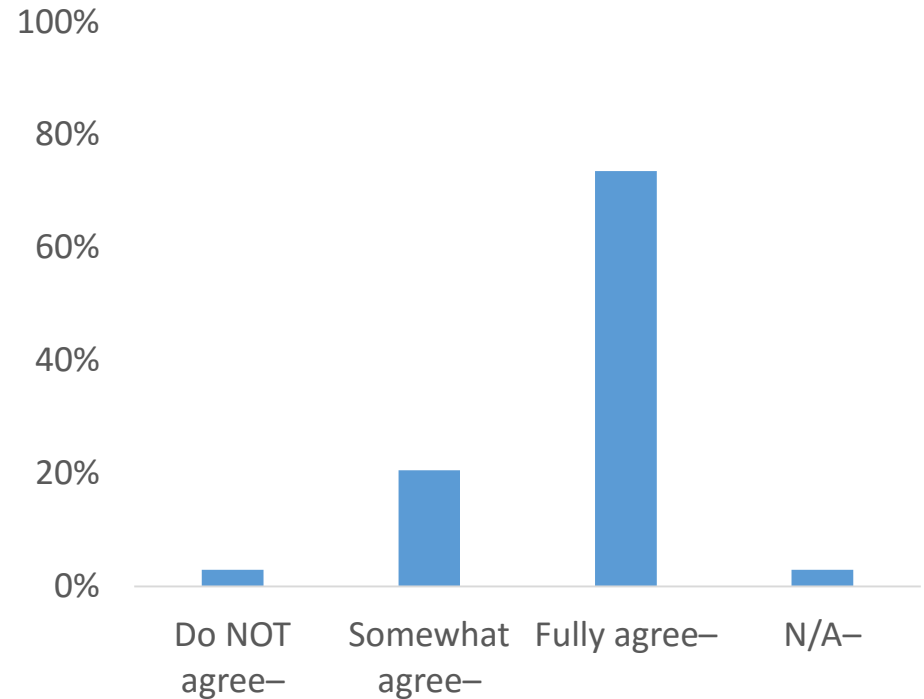
# Were we successful?

End of quarter  
survey

## Skills or STEM Practices



**I feel more comfortable interpreting graphs compared to when I started the course.**



**I feel more comfortable translating jargon into plain language compared to when I started the course.**

# Were we successful?

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## Typical Positive Statements

*“The poster activities were extremely helpful. Especially when we had to explain our diagrams/answers to another group.”*

*“Physically modeling biological phenomena makes you remember things better. There is not a single activity or worksheet we did that I do not feel was useful.”*

*“I would study for two weeks for exams in my other bio classes and would not retain as much information and would not do nearly as well. I walk away from this class with a strong memory of everything we learned throughout this entire course.”*

## End of quarter survey

## Typical Areas of Improvement

*“This class was very demanding. I liked the style of learning a lot but I didn't like the worksheets we had to do. Often I felt that they were unfairly graded (too many points taken off) and that we didn't go over the answers enough for a full understanding of the concept. Often lecture seemed rushed.”*

*“Better time management so we have time to talk about an activity after we do it”*

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Taking it to the masses!

# *What will we change?*

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- *Pare down the number of activities*
- *Reduce active lecture more by using more original videos*
- *Use section to finish worksheets if needed*
- *Learning outcomes posted at the beginning of the course*
- *Potentially use sections to do synthesis questions*
- *Have learning assistants & teaching assistants always sit among the class rather than clustered together*
- *Incorporate more exam questions that try to get at skills*



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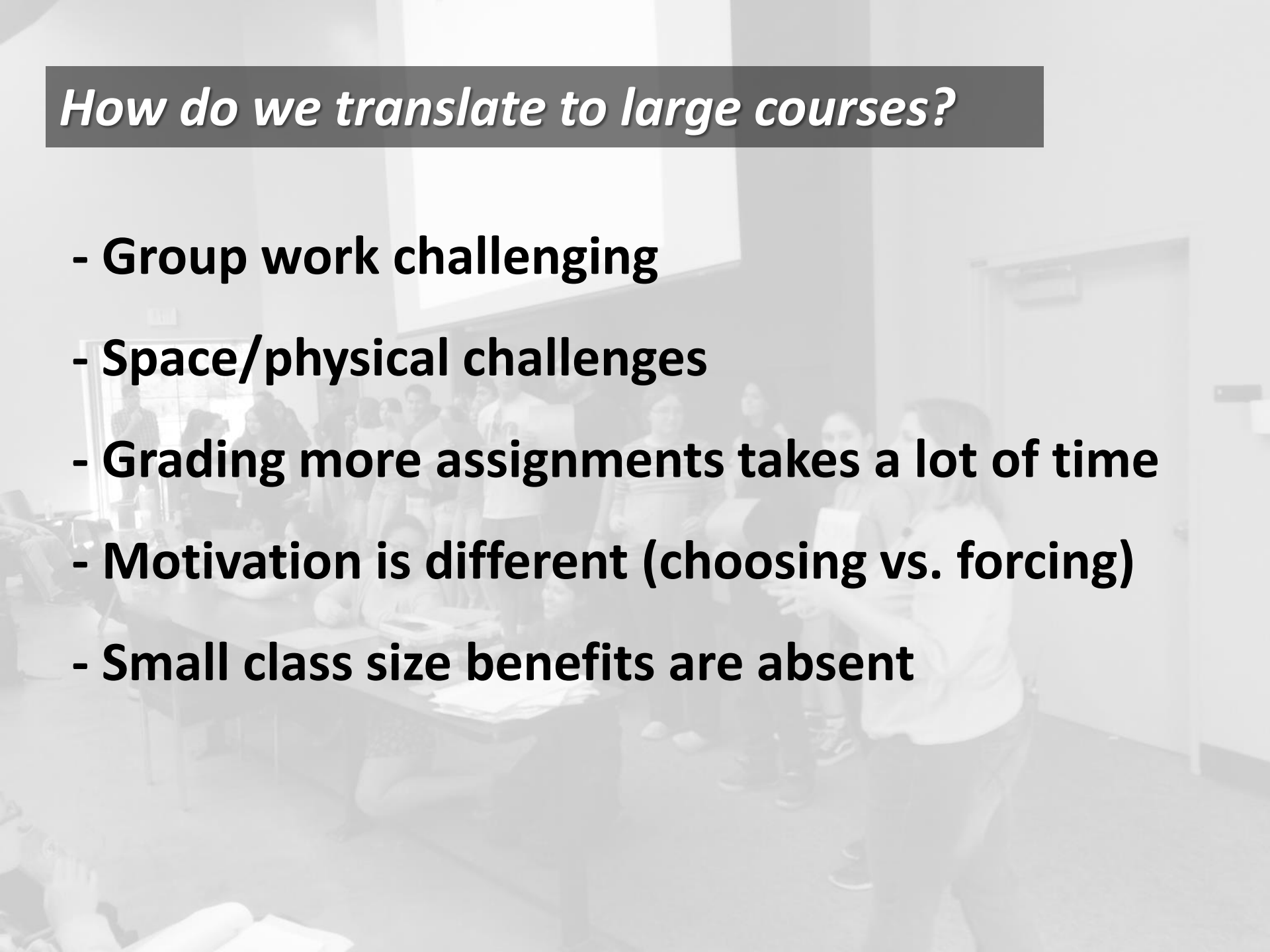
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What would we change?

5

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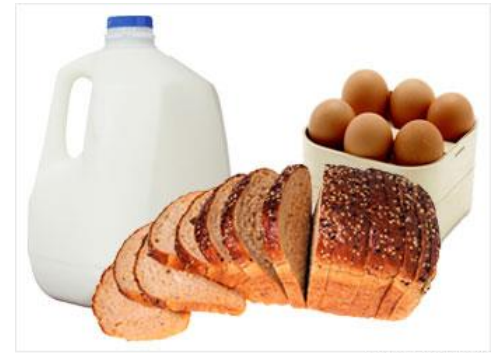
## *How do we translate to large courses?*

- **Group work challenging**
  - **Space/physical challenges**
  - **Grading more assignments takes a lot of time**
  - **Motivation is different (choosing vs. forcing)**
  - **Small class size benefits are absent**
- 

# *Taking it to the masses!*

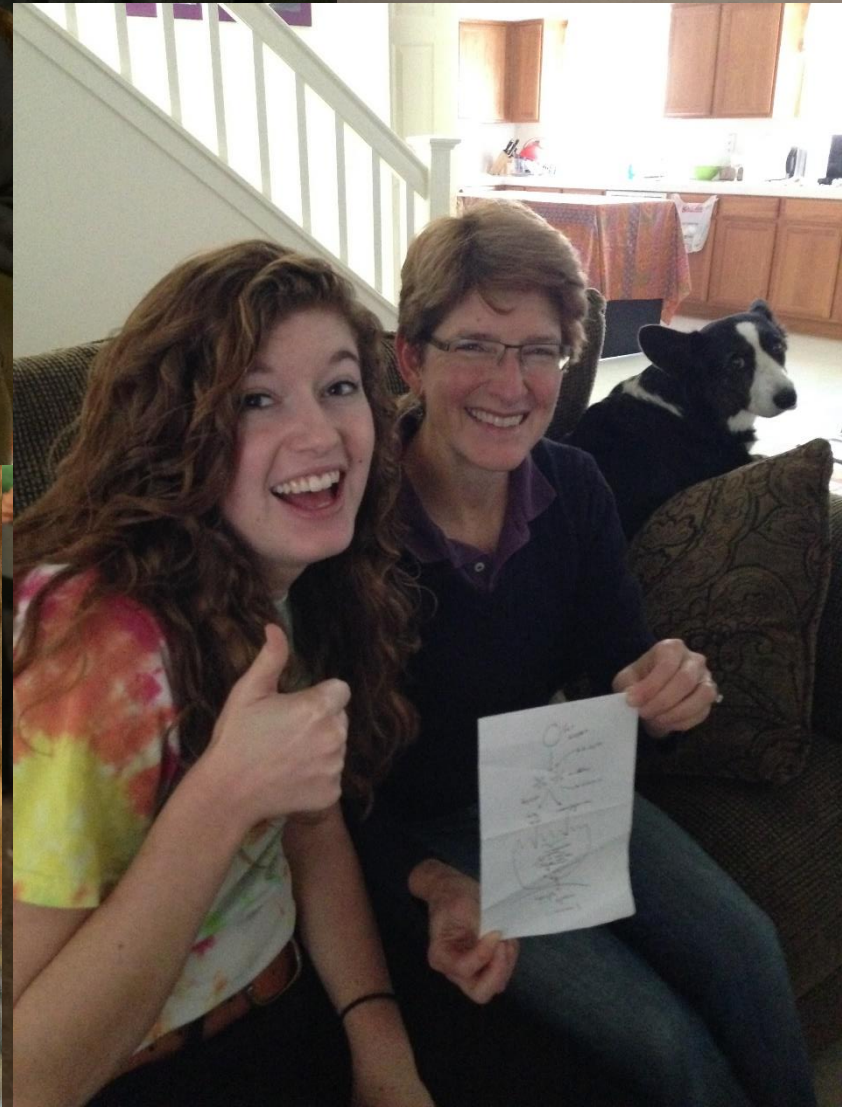
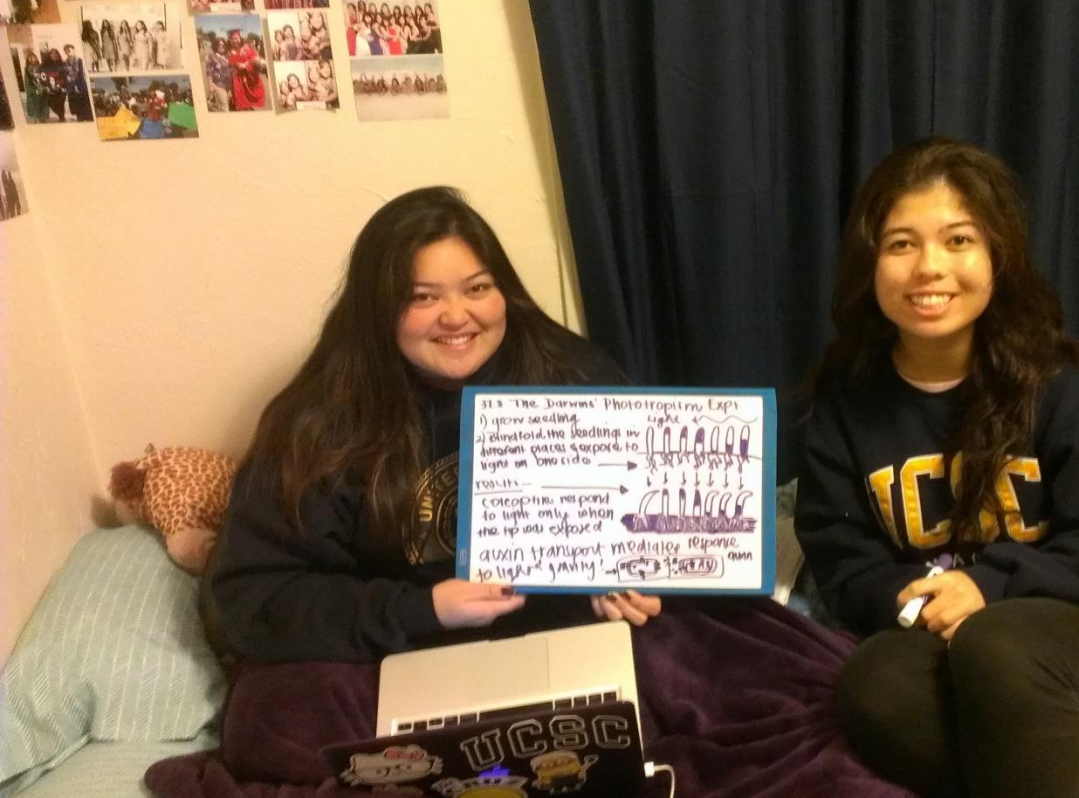
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The staples....tried, true...and cheap!  
*(potentially considered a phase 1)*



Reintegrating activities into large lecture with modifications

- *Evidence and modeling/Rap Genius with modification*
- *Giving graphs/critical thinking questions in lecture and giving time to struggle*
- *Worksheets as homework, go over in class after giving students a chance to work with neighbors*
- *Humans-as-props! Fish bowl demos*
- *A note about TAs & LAs*



# *What I have Learned?*

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- Forced me to focus even more on what very specific ideas/concepts I want students to master
- Refocus on skills – with each concept that we cover, how can I work in time to practice a skill
- Less is more
- Organization is really important
- Transparency with students is really important
- Some things are going to flop – contexting at the beginning is important

# *Did the concerns we started with come to pass?*

- Will students do the work needed outside of class?
- How much content are we going to have to cut?
- Will students engage or will it be like a whole quarter of flopping at open mic night?
- Will our evaluations suffer?
- How much extra time/work is this going to take?
- Will they really do better?



Susie and I....

# Questions?

Mid quarter!



End of Quarter!

