

The background of the slide features a large, light gray watermark of the University of Colorado seal. The seal is circular and contains the text "UNIVERSITY OF COLORADO" around the top edge and "1876" at the bottom. In the center, it depicts a figure holding a torch and a book, with the motto "LET YOUR LIGHT SHINE" written above the figure.

Towards a coherent & interactive curriculum in the sciences

Mike Klymkowsky

Molecular, Cellular & Developmental Biology / CU Teach

TRUSE II June

Melanie Cooper (Clemson) & the CLUE team



CU BOULDER Teach
Science & Mathematics

STE&M ER@CU
science, technology, engineering & mathematics education research

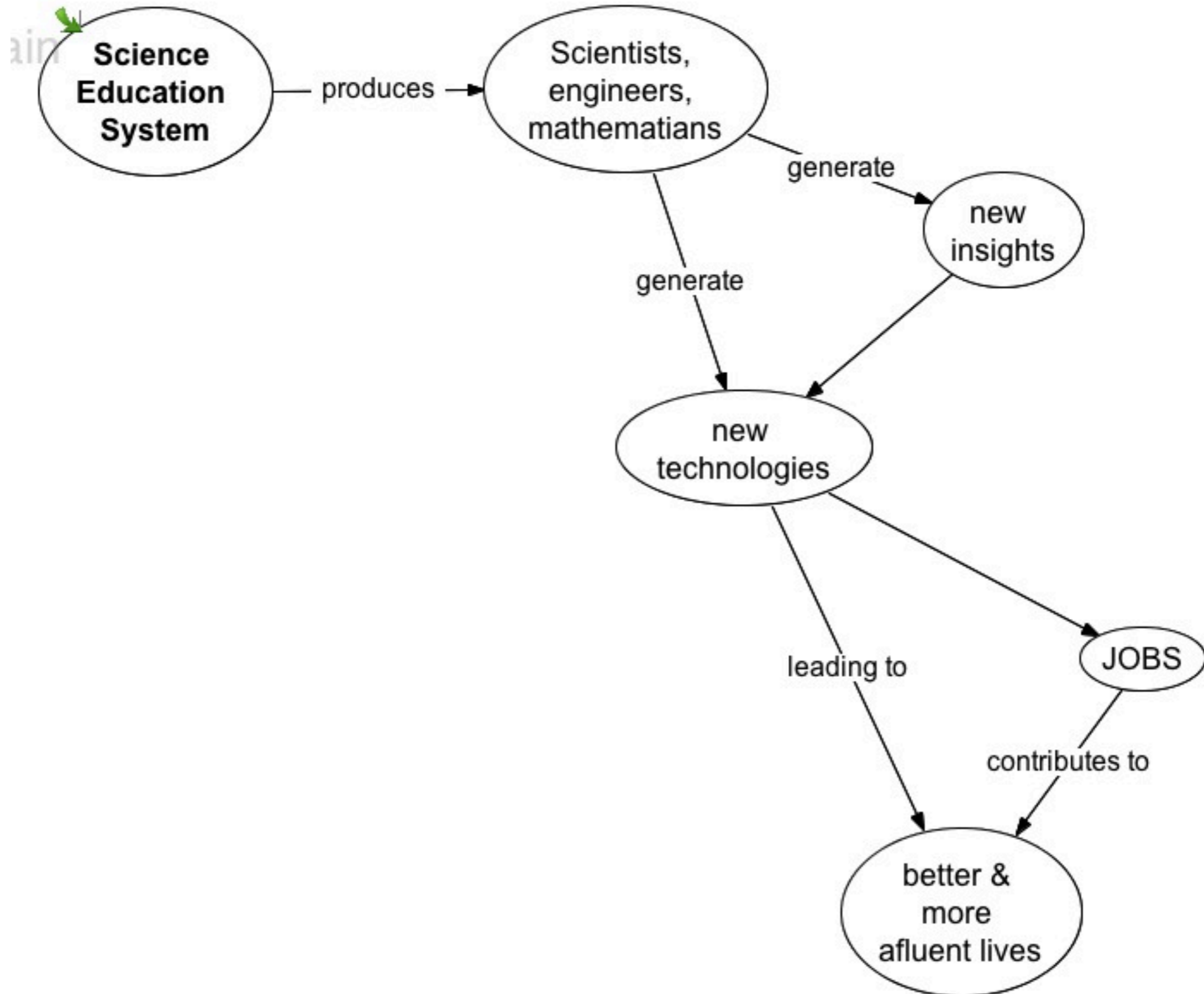
thanks

& our students

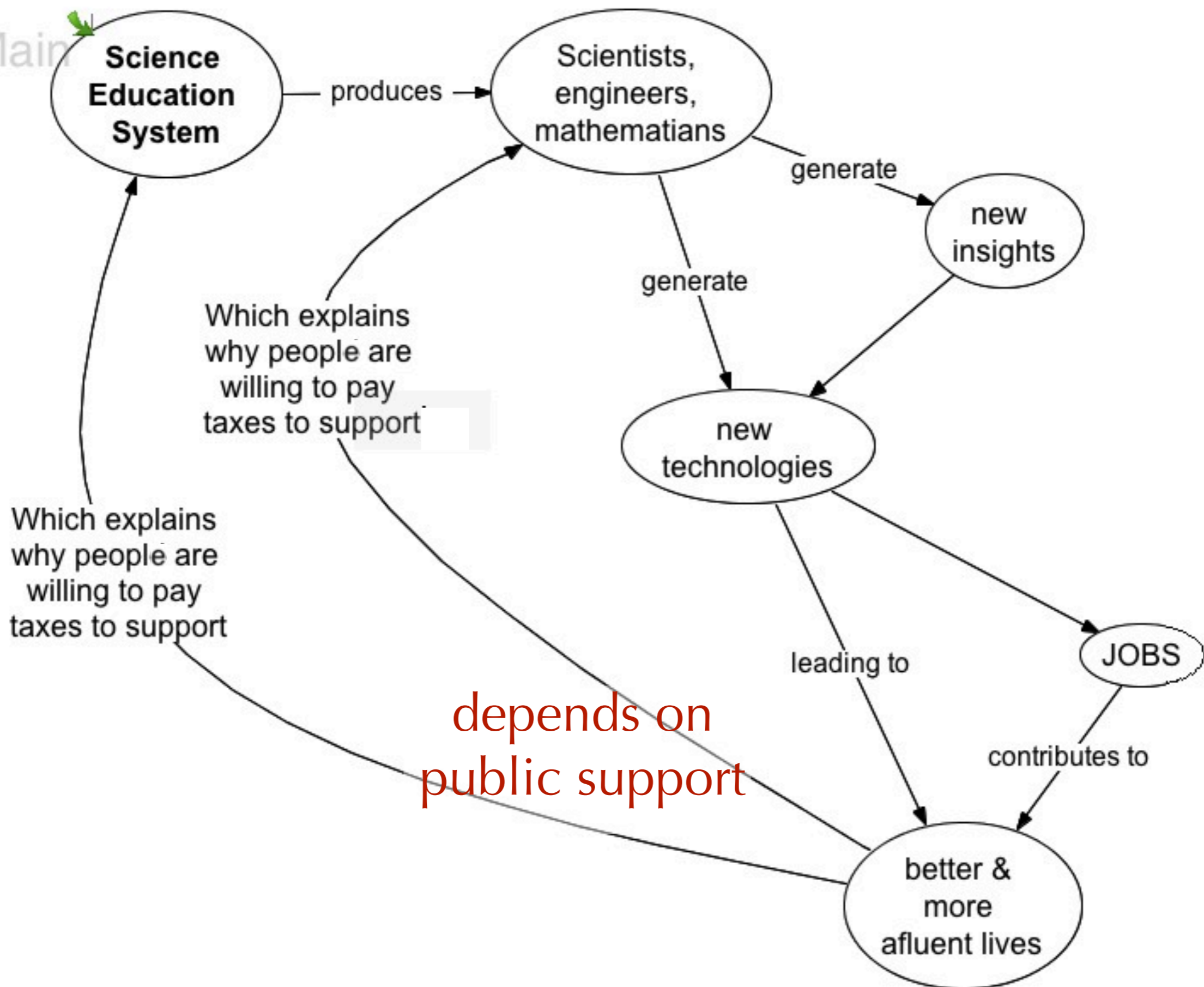


Who & what is science education for?

most obviously

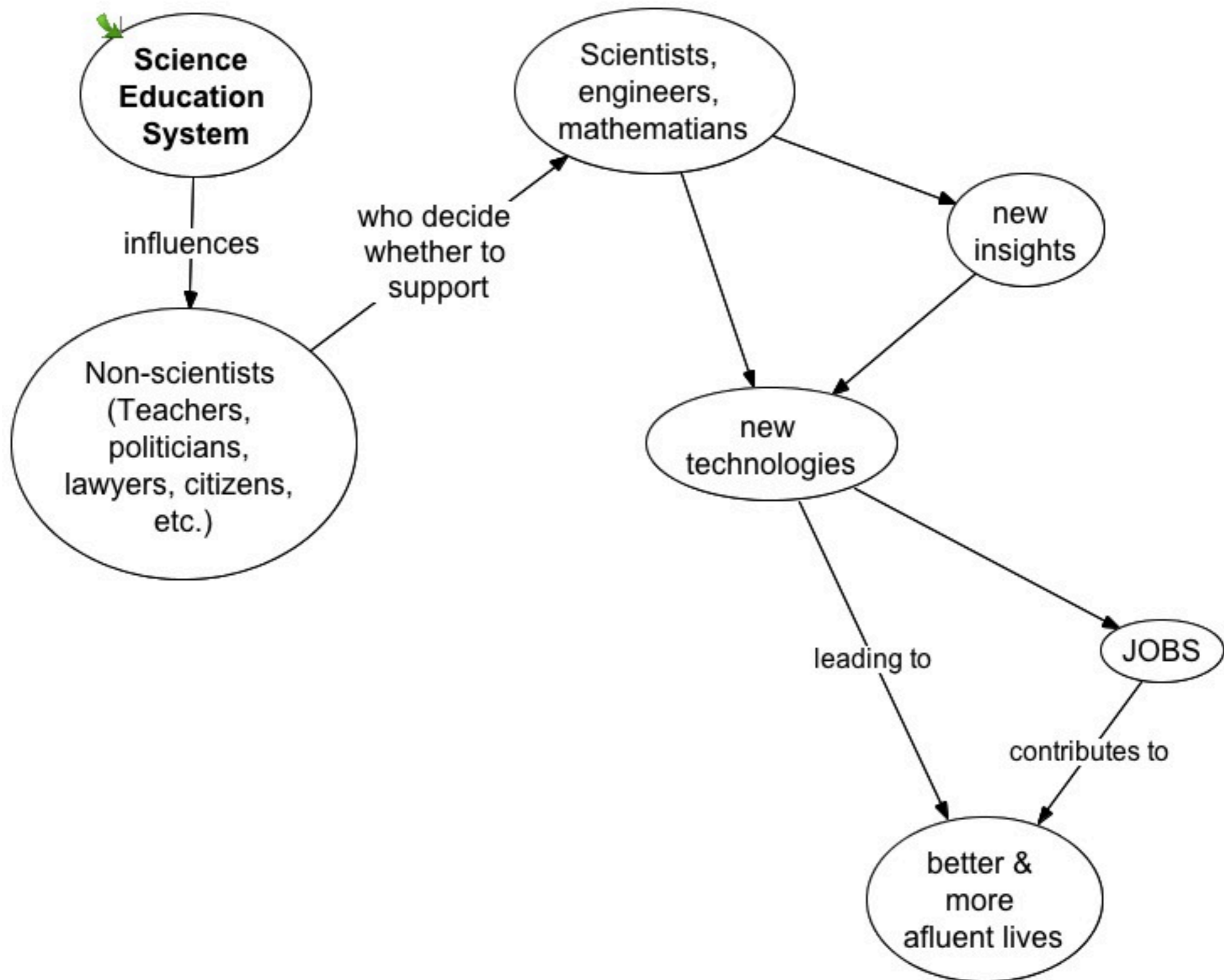


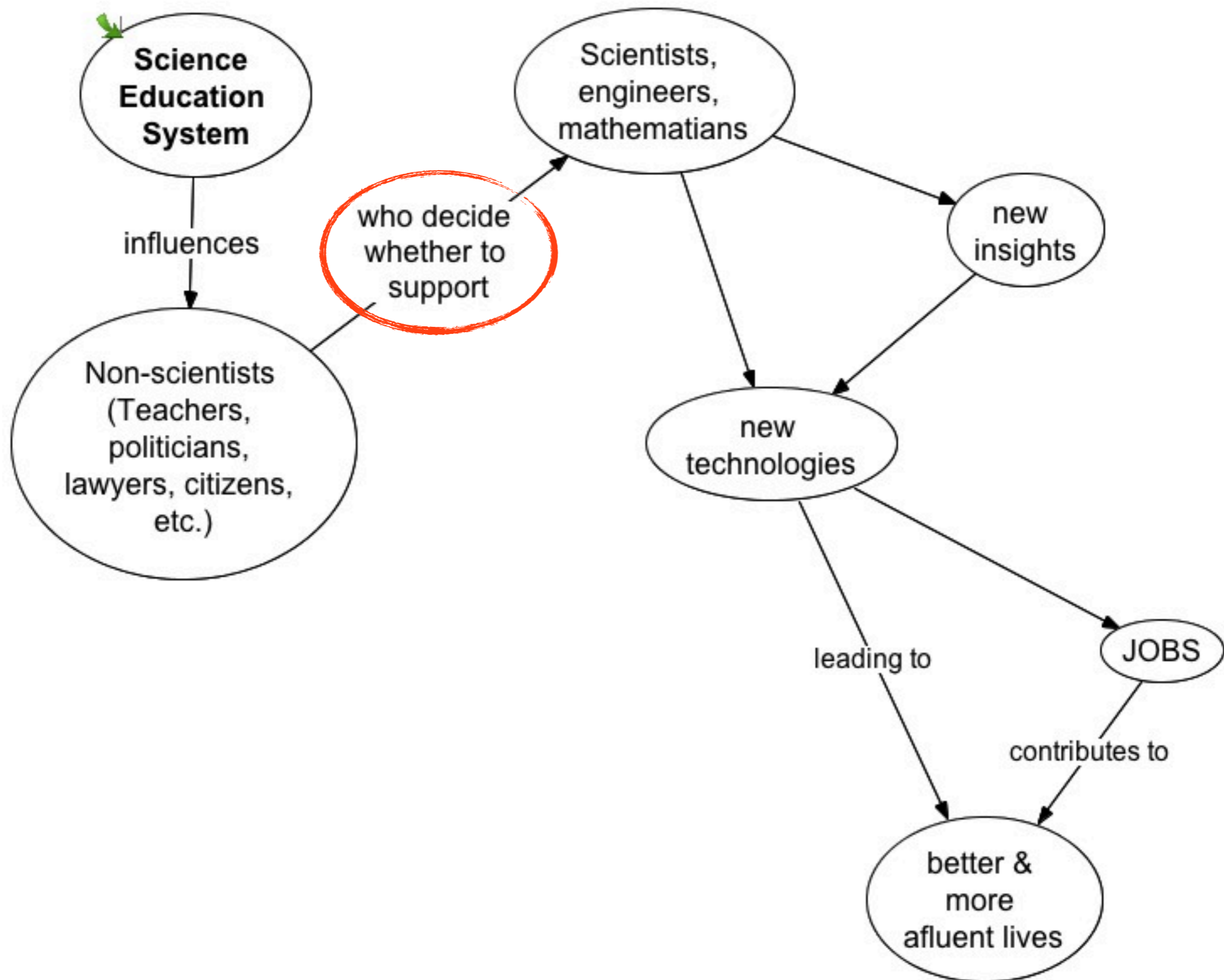
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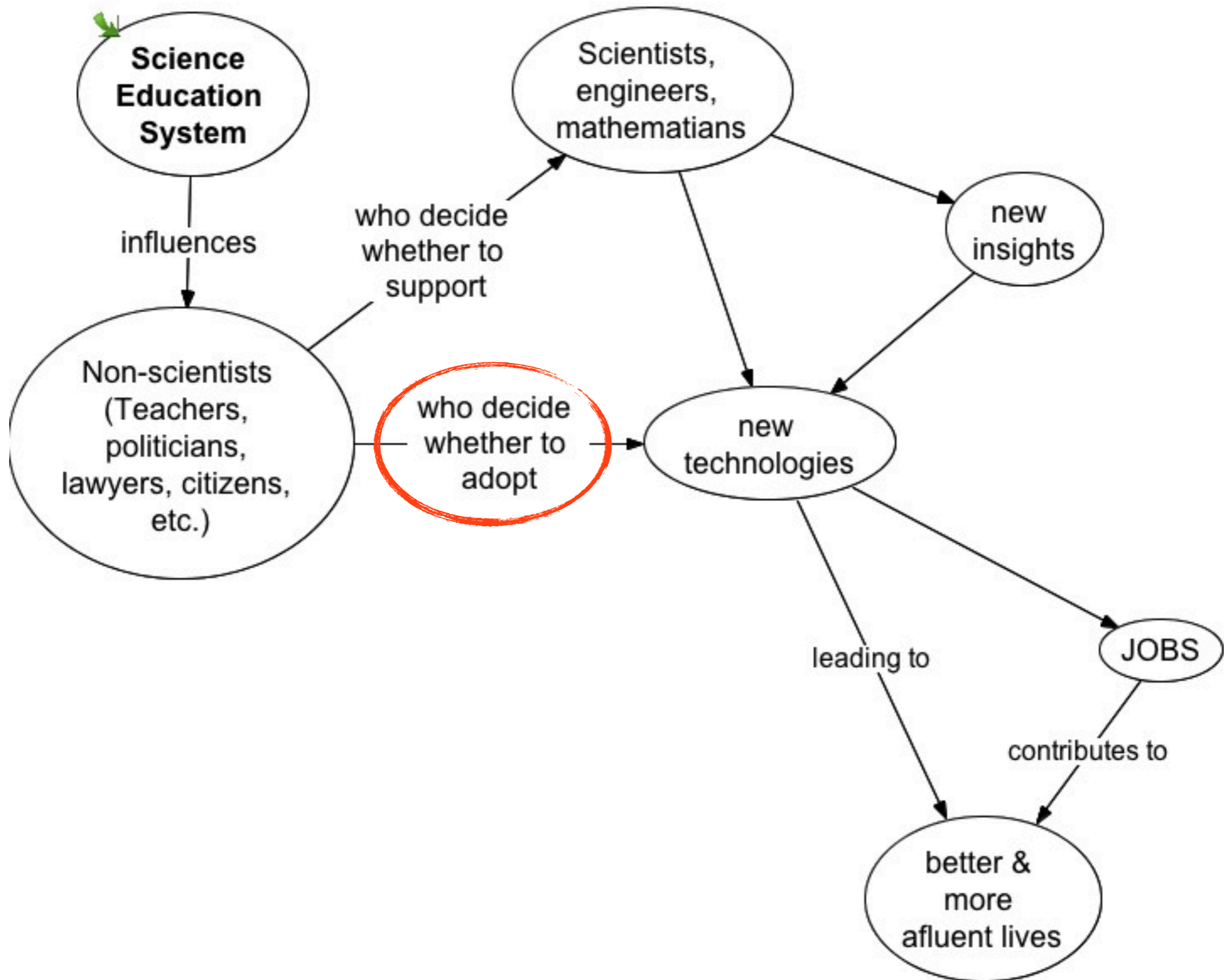


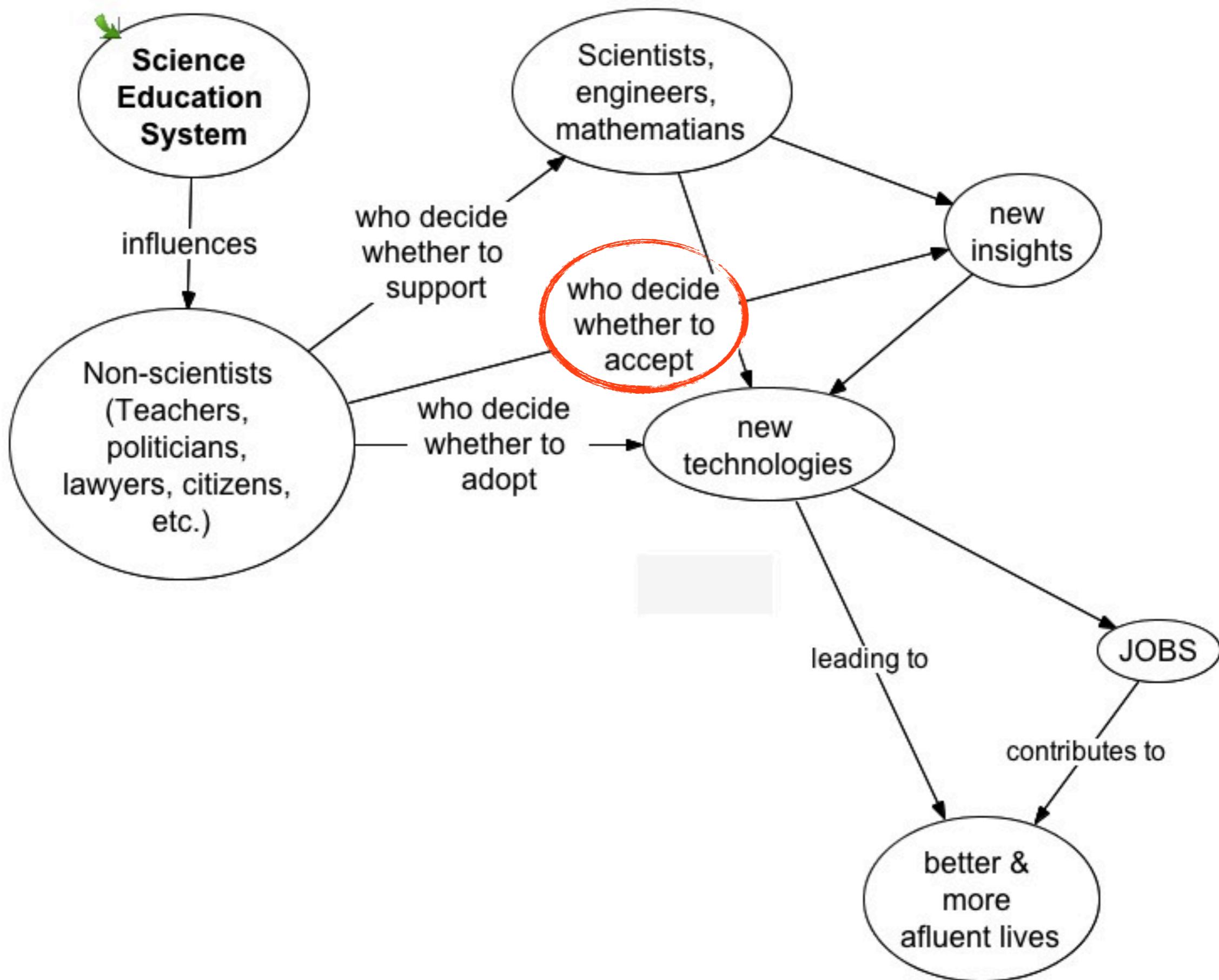


Has effects
beyond the
generation of
scientists











which is not without consequences

DEADLY CHOICES

— HOW THE —
ANTI-VACCINE
— MOVEMENT —
THREATENS
— US ALL —

PAUL A. OFFIT, M.D.

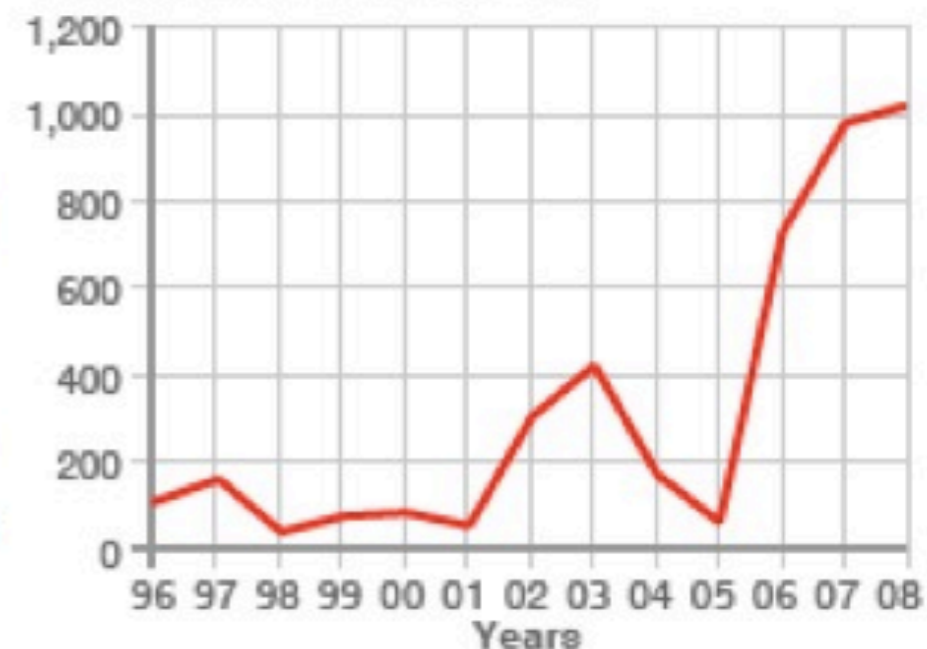
which is not without consequences

NOVEMBER 28, 2008

Reduced MMR Equals More Measles

Reduced uptake of the MMR vaccine, fueled no doubt by anti-vaccine propaganda, has resulted in a recent **significant increase in Measles in the UK** as shown by the graph on the right. And despite what the anti-vaccine twits will tell you, Measles can be a very serious disease. According to the **CDC**:

YEARLY MEASLES CASES



SOURCE: Health Protection Agency

DEADLY CHOICES

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ANTI-VACCINE
MOVEMENT
THREATENS
US ALL

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As many as one out of 20 children with measles gets pneumonia, and about one child in every 1,000 who get measles will develop encephalitis. (This is an inflammation of the brain that can lead to convulsions, and can leave your child deaf or mentally retarded.) For every 1,000 children who get measles, one or two will die from it.

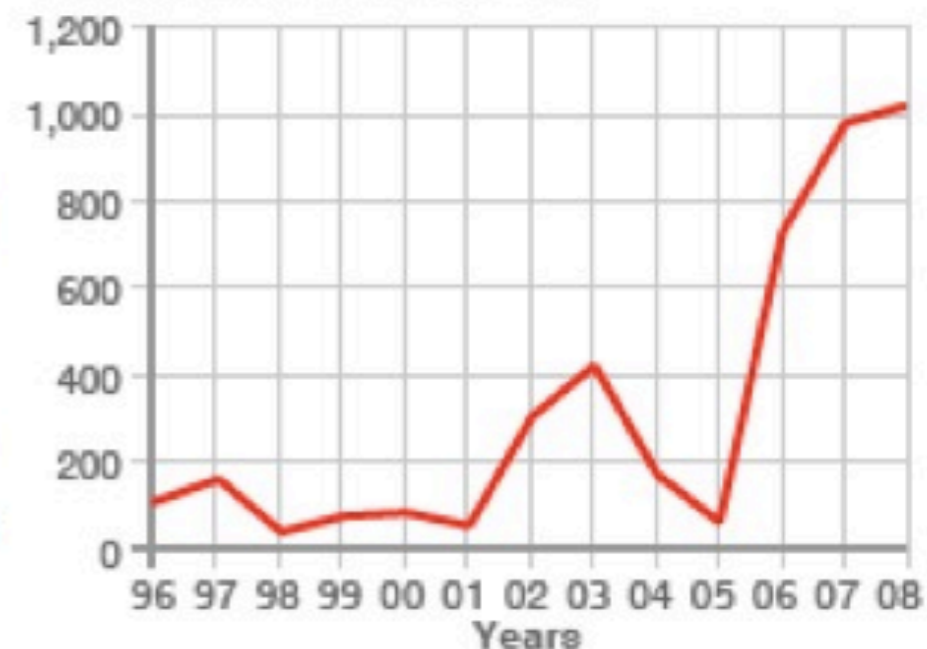
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Louisiana's bold bid to privatize schools

 Recommend  7,300 people recommend this. Be the first of your friends.

By Stephanie Simon

June 1 | Fri Jun 1, 2012 6:04pm EDT

 Tweet  464

The Upperroom Bible Church Academy in New Orleans, a bunker-like building with no windows or playground, also has plenty of slots open. It seeks to bring in 214 voucher students, worth up to \$1.8 million in state funding.

At Eternity Christian Academy in Westlake, pastor-turned-principal Marie Carrier hopes to secure extra space to enroll 135 voucher students, though she now has room for just a few dozen. Her first- through eighth-grade students sit in cubicles for much of the day and move at their own pace through Christian workbooks, such as a beginning science text that explains "what God made" on each of the six days of creation. They are not exposed to the theory of evolution.

"We try to stay away from all those things that might confuse our children," Carrier said.

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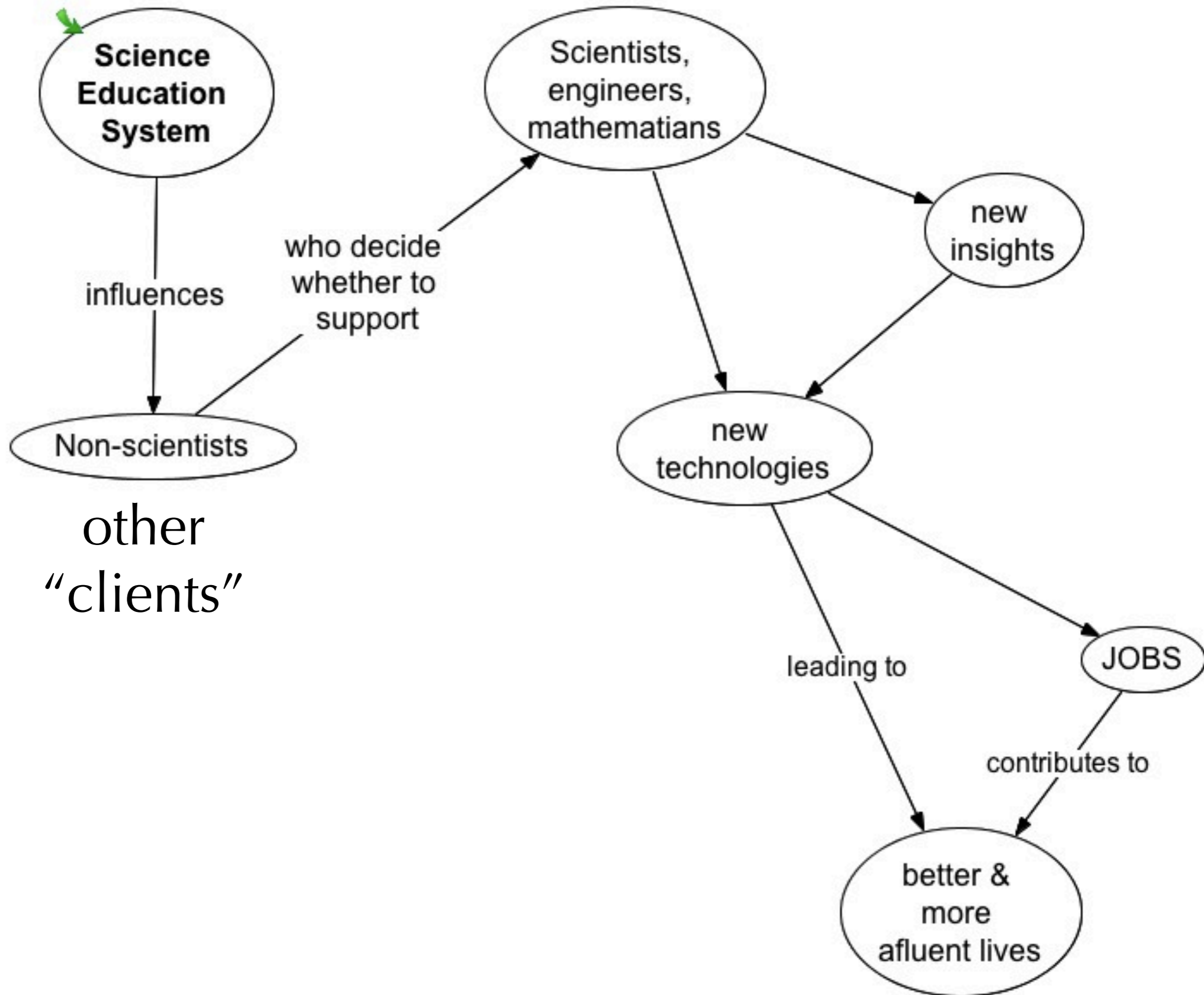
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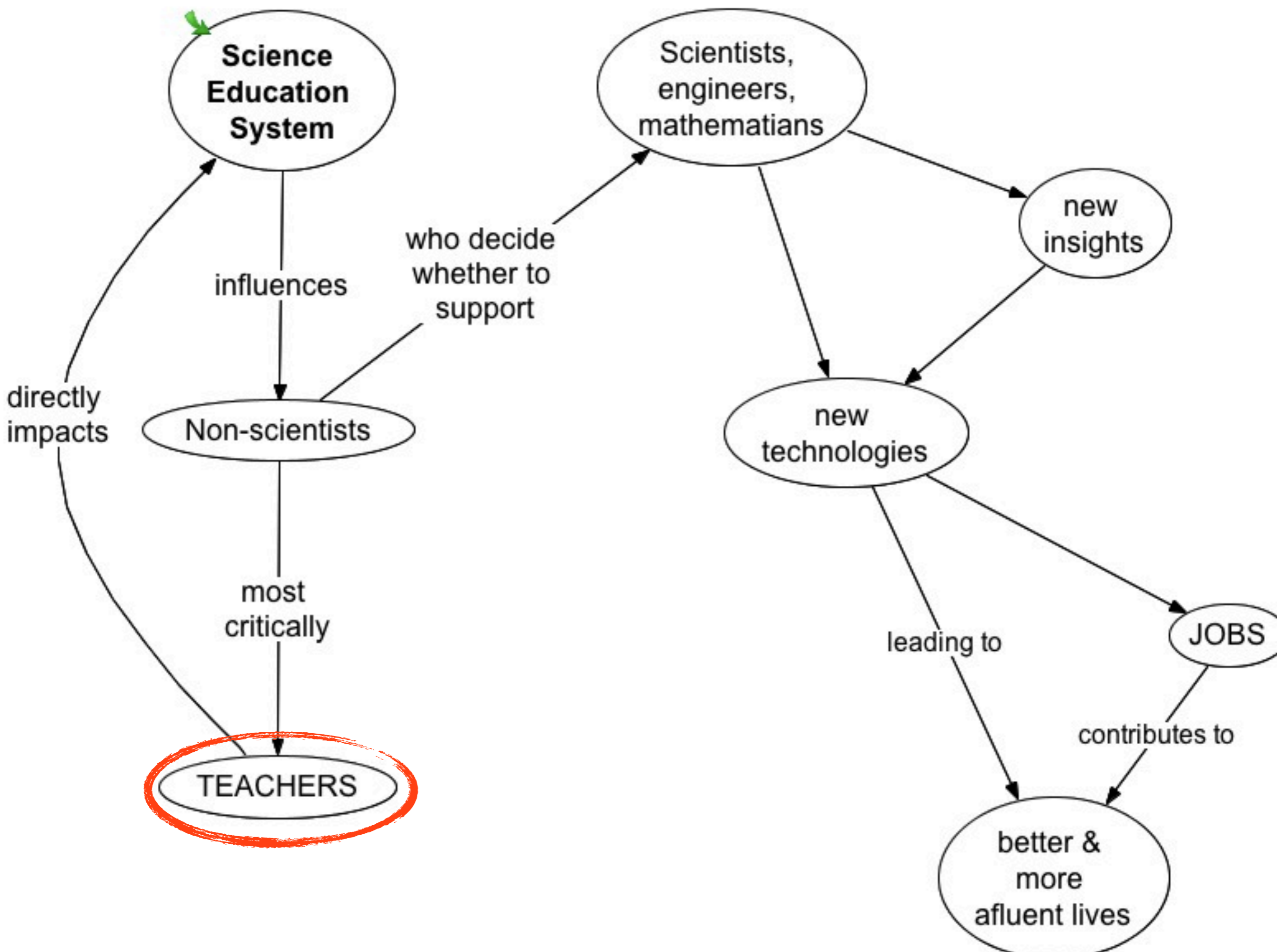
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other
"clients"



Model I: Success is in generating **scientists**

- equivalent of producing best selling authors, accomplished musicians, successful athletes and entrepreneurs)

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- equivalent of producing best selling authors, accomplished musicians, successful athletes and entrepreneurs)

Q: Does this come with unintended consequences for non-scientists?

Model II: Success is teaching people to understand (teach) science: how to read, appreciate music, understand the “game”.

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Q: Should teachers be our “target” audience. Do they represent our most important student population?

Model II: Success is teaching people to understand (teach) science: how to read, appreciate music, understand the “game”.

Q: Should teachers be our “target” audience. Do they represent our most important student population?

Q: Would this improve learning outcomes for all students?

either way, remember that scientific thinking is unnatural (hard)

The New York Times

November 27, 2011



Illustration by David Plankoff

Thinking fast and slow.
Daniel Kahneman (2011)

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November 27, 2011



Illustration by David Plankoff

Thinking fast and slow.
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Now for the Hard Part: The Path to Coherent Curricular Design*

Received for publication, April 12, 2012, and in revised form, April 17, 2012

Michael W. Klymkowsky‡ and Melanie M. Cooper§

From the ‡Department of MCD Biology/CU Teach, University of Colorado, Boulder, Colorado 80309-0347 and §Department of Chemistry, Clemson University, Clemson, South Carolina

Teaching disconcerting scientific ideas

BY MIKE KLYMKOWSKY

Explaining the scientific process will help the public understand why scientists trust their own conclusions.

Strange scientific ideas

- Matter is composed of atoms, which are mostly empty space.
- The universe emerged out of nothing (about 13,700,000,000 years ago).
- There are billions of galaxies, each containing billions of stars.
- Time and space are not distinct.
- All organisms are built from similar building blocks called cells.
- All cells are derived from pre-existing cells in a continual lineage that extends back about 3,500,000,000 years.
- The heavier atoms in our bodies were formed within stars or exploding stars.
- Matter and energy are different versions of the same thing.
- The universe is running down yet expanding at a faster and faster rate.
- Random noise can produce complex structures.
- At the molecular level, everything is reversible.
- A collection of cells can, by itself, produce a self-conscious entity that thinks it is more than a collection of cells.

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Acting as if scientific ideas are obvious, easily mastered, or easy to accepted can leave students feeling stupid.

Strange scientific ideas

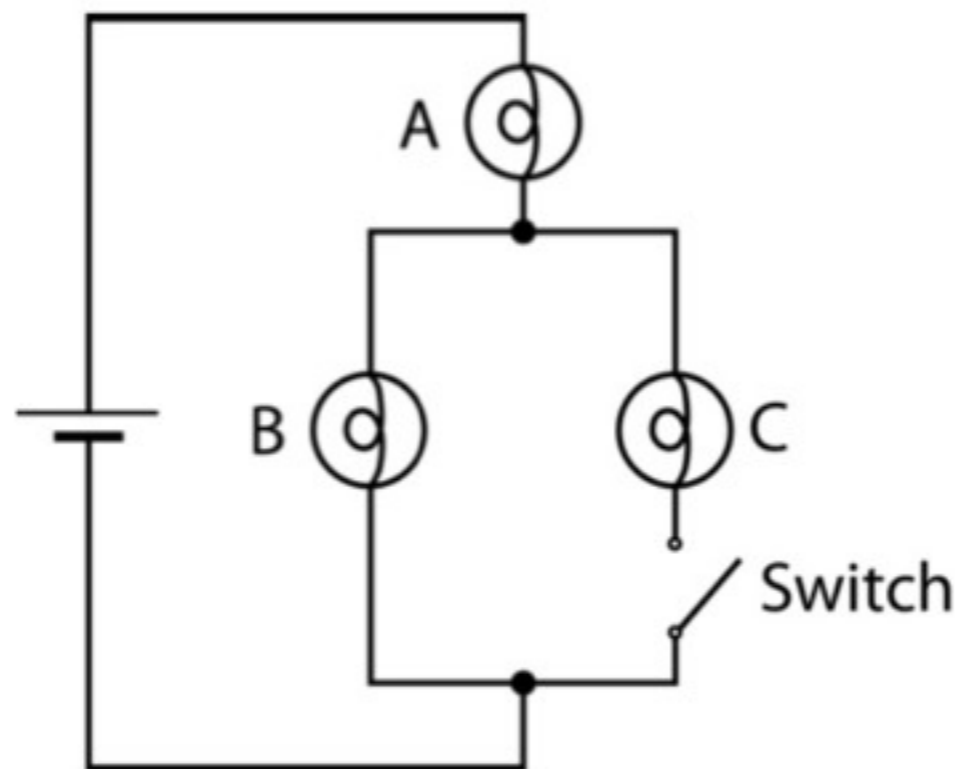
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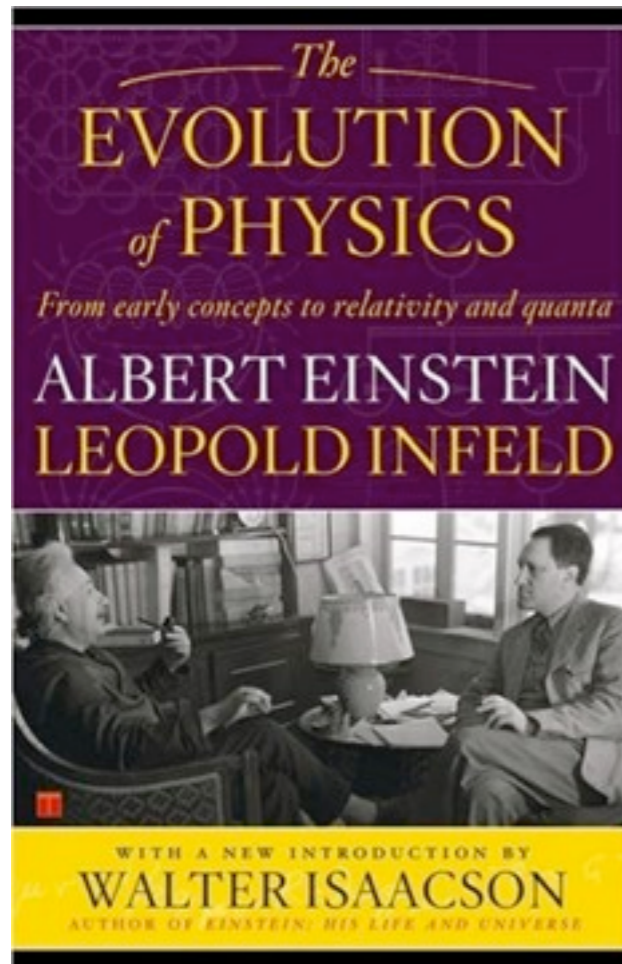
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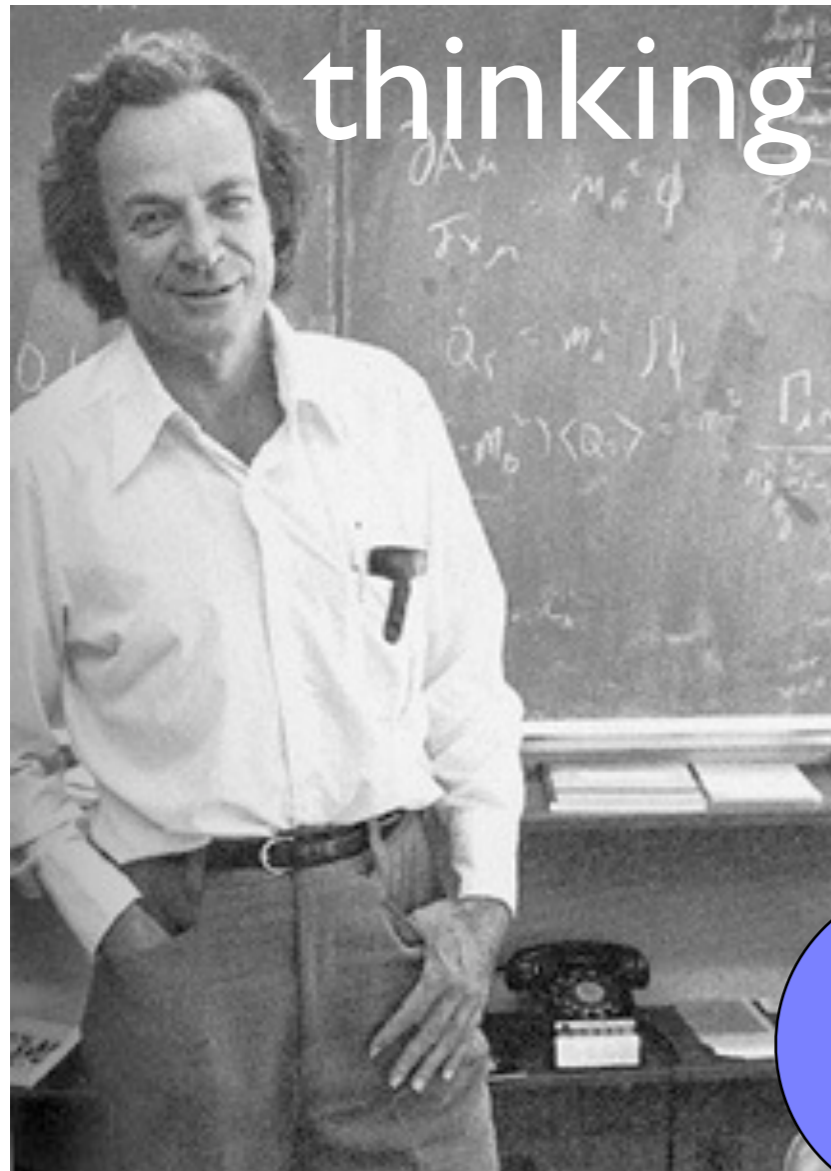
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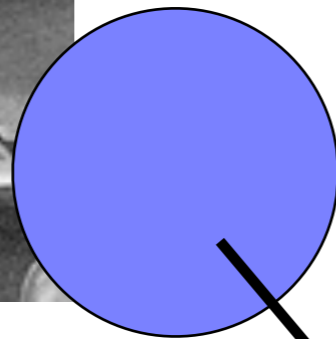
“...new physical concepts are born in the painful struggle with old ideas.”

- Einstein & Infeld, 1967. The Evolution of Physics.

understanding builds on foundations:

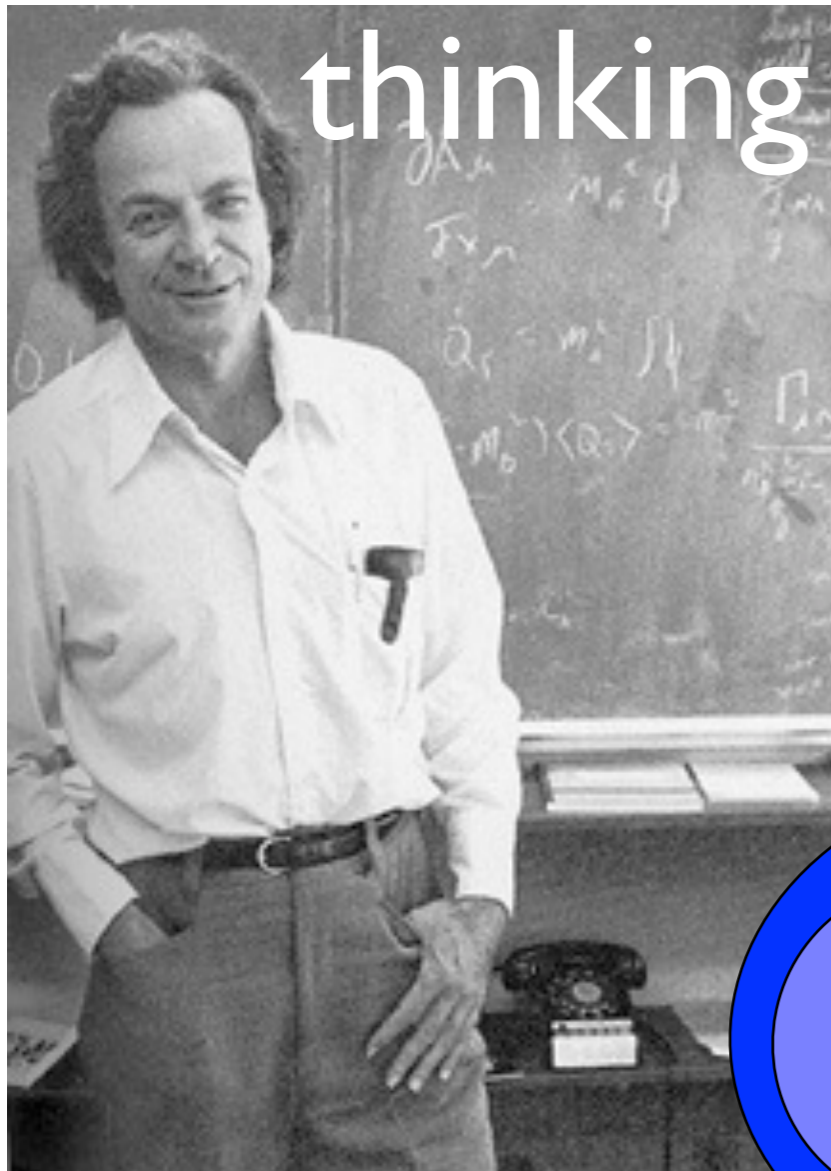


thinking about magnets



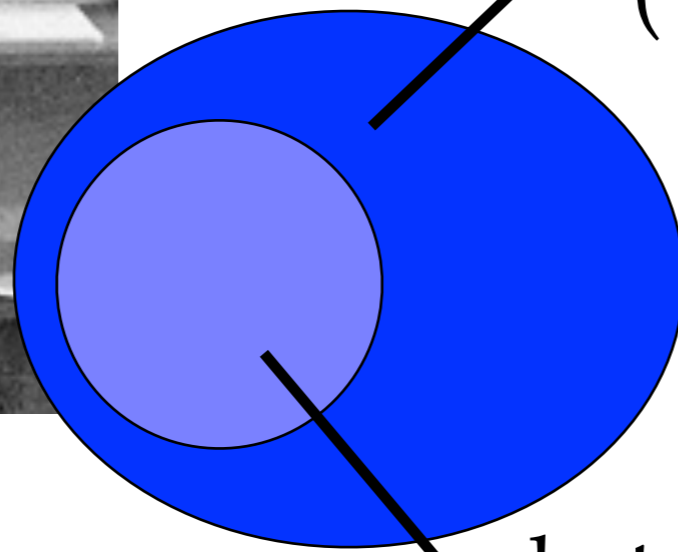
what the student knows
(or thinks they know)

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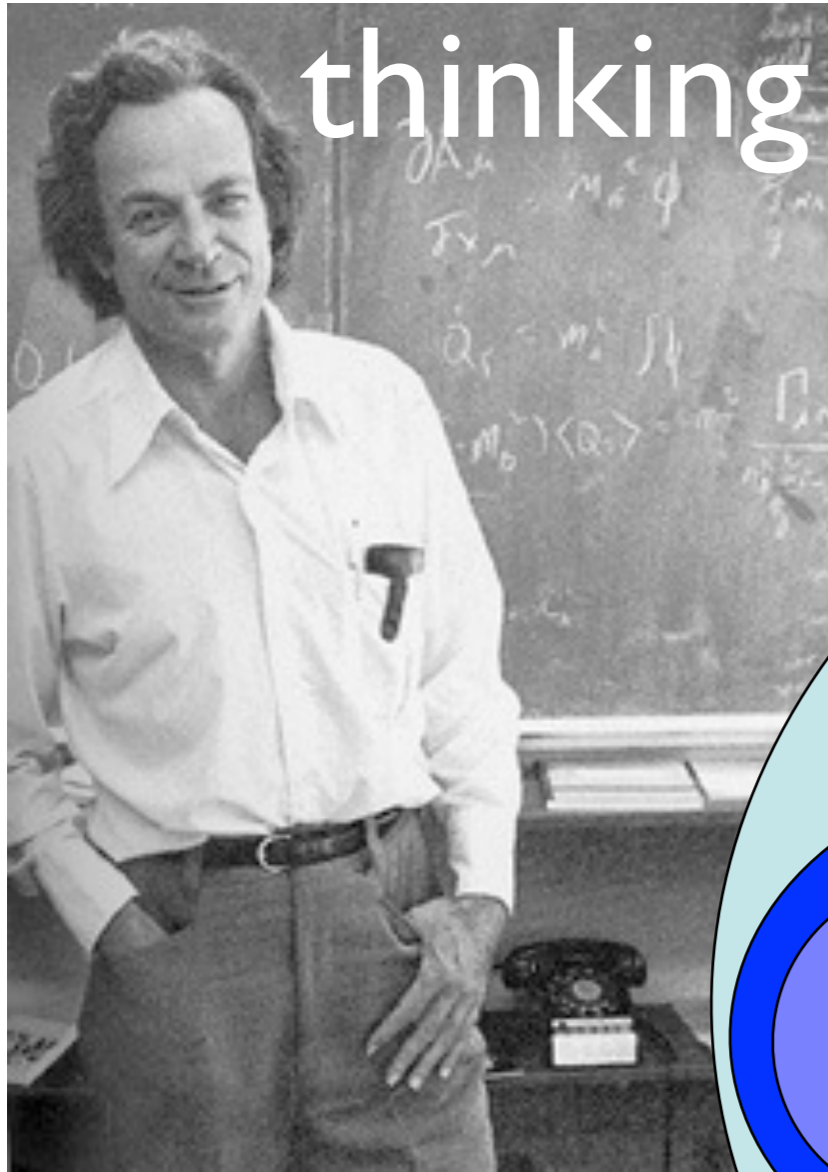


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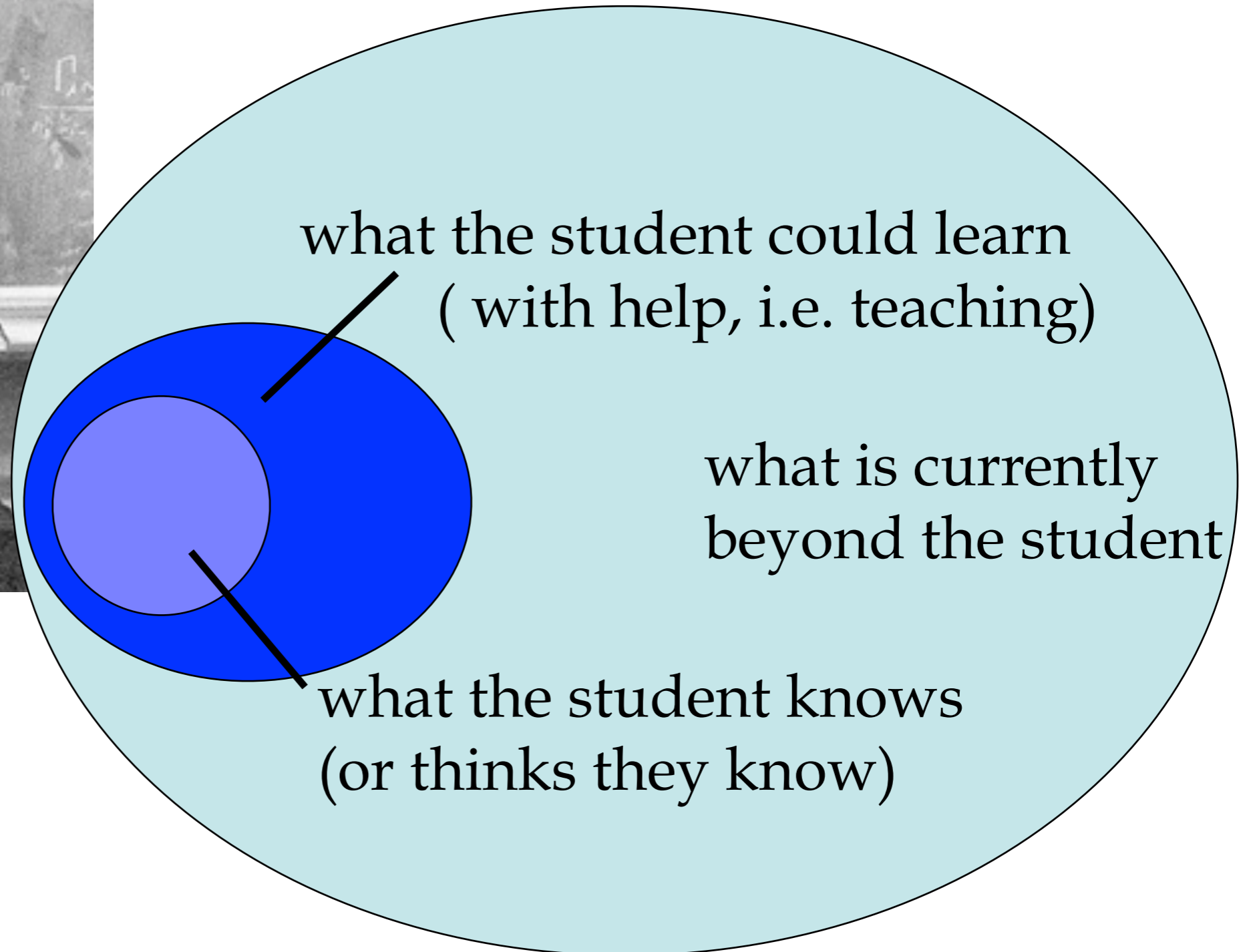
what the student could learn
(with help, i.e. teaching)



what the student knows
(or thinks they know)



thinking about magnets

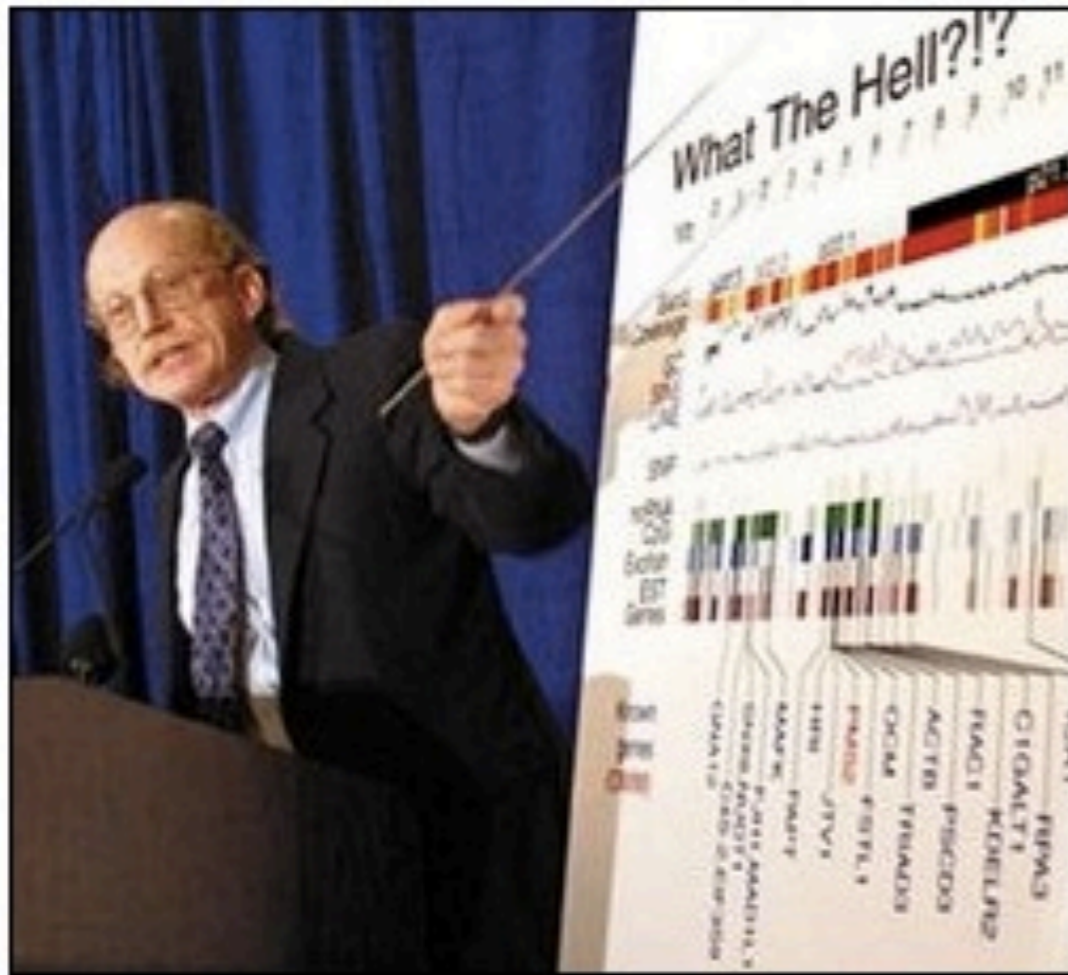


National Science Foundation: Science Hard

JUNE 5, 2002 | ISSUE 45•01 ISSUE 38•21

INDIANAPOLIS—The National Science Foundation's annual symposium concluded Monday, with the 1,500 scientists in attendance reaching the consensus that science is hard.

[Enlarge Image](#)



Farian explains the NSF findings.

newly discovered 'Law of Difficulty' holds true for all branches of science, from astronomy to molecular biology and everything in between."

"For centuries, we have embraced the pursuit of scientific knowledge as one of the noblest and worthiest of human endeavors, one leading to the enrichment of mankind both today and for future generations," said keynote speaker and NSF chairman Louis Farian. "However, a breakthrough discovery is challenging our long-held perceptions about our discipline—the discovery that science is really, really hard."

"My area of expertise is the totally impossible science of particle physics," Farian continued, "but, indeed, this

RELATED ARTICLES

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What are we expecting to achieve, exactly?

What are we expecting to achieve, exactly?

What are our “performance expectations” for students?

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What are our “performance expectations” for students?

What do these expectations imply about what students need to know?

What are we expecting to achieve, exactly?

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What do these expectations imply about what students need to know?

Are they attainable?

What are we expecting to achieve, exactly?

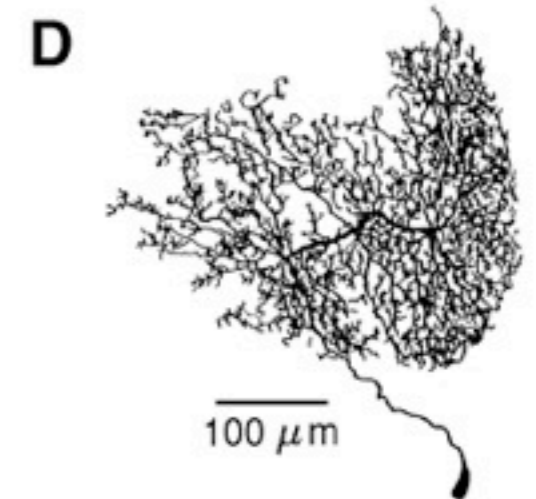
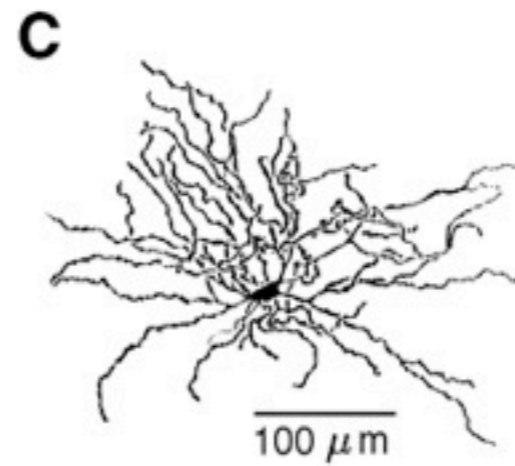
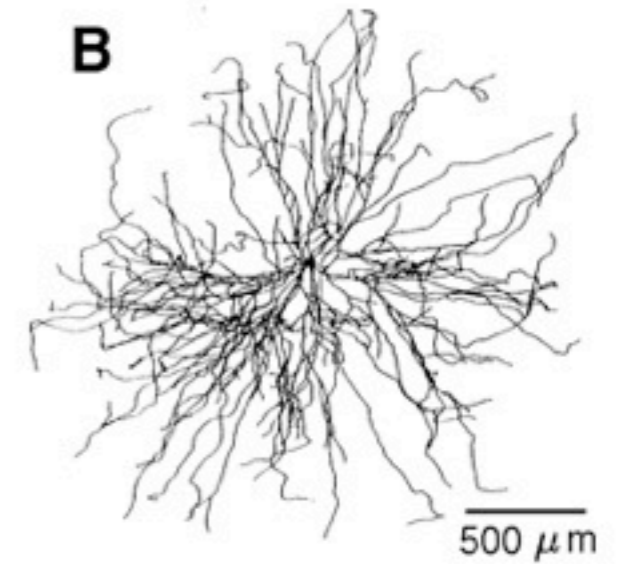
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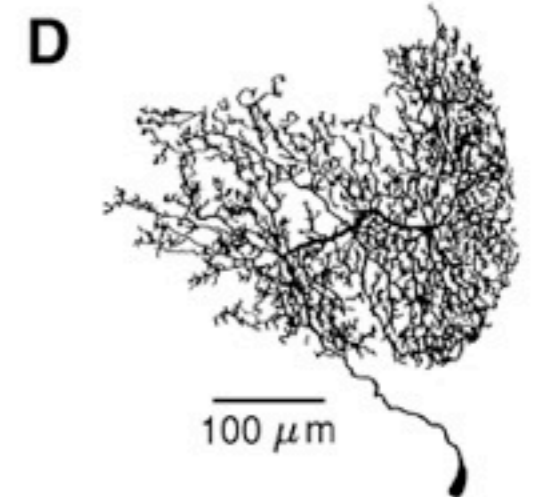
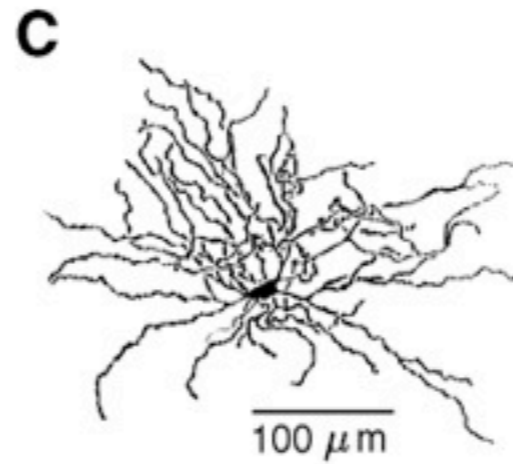
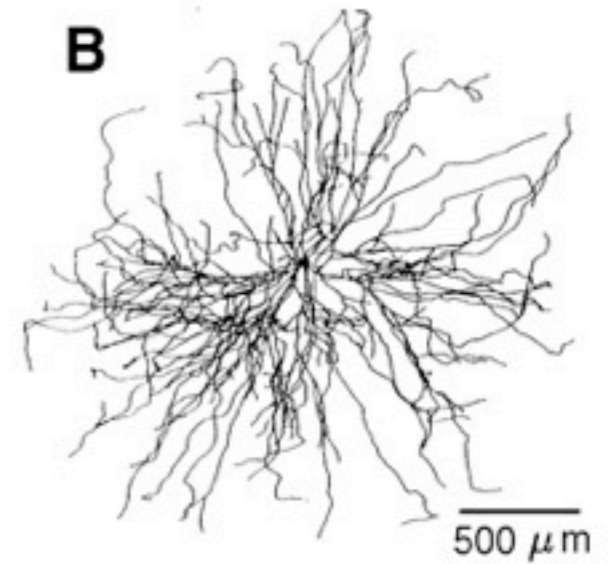
or do they require curriculum redesign?

NGSS draft performance expectation: Model the development of long term memory



NGSS draft performance expectation: Model the development of long term memory

Superficial - cells change



NGSS draft performance expectation:

Model the development of long term memory

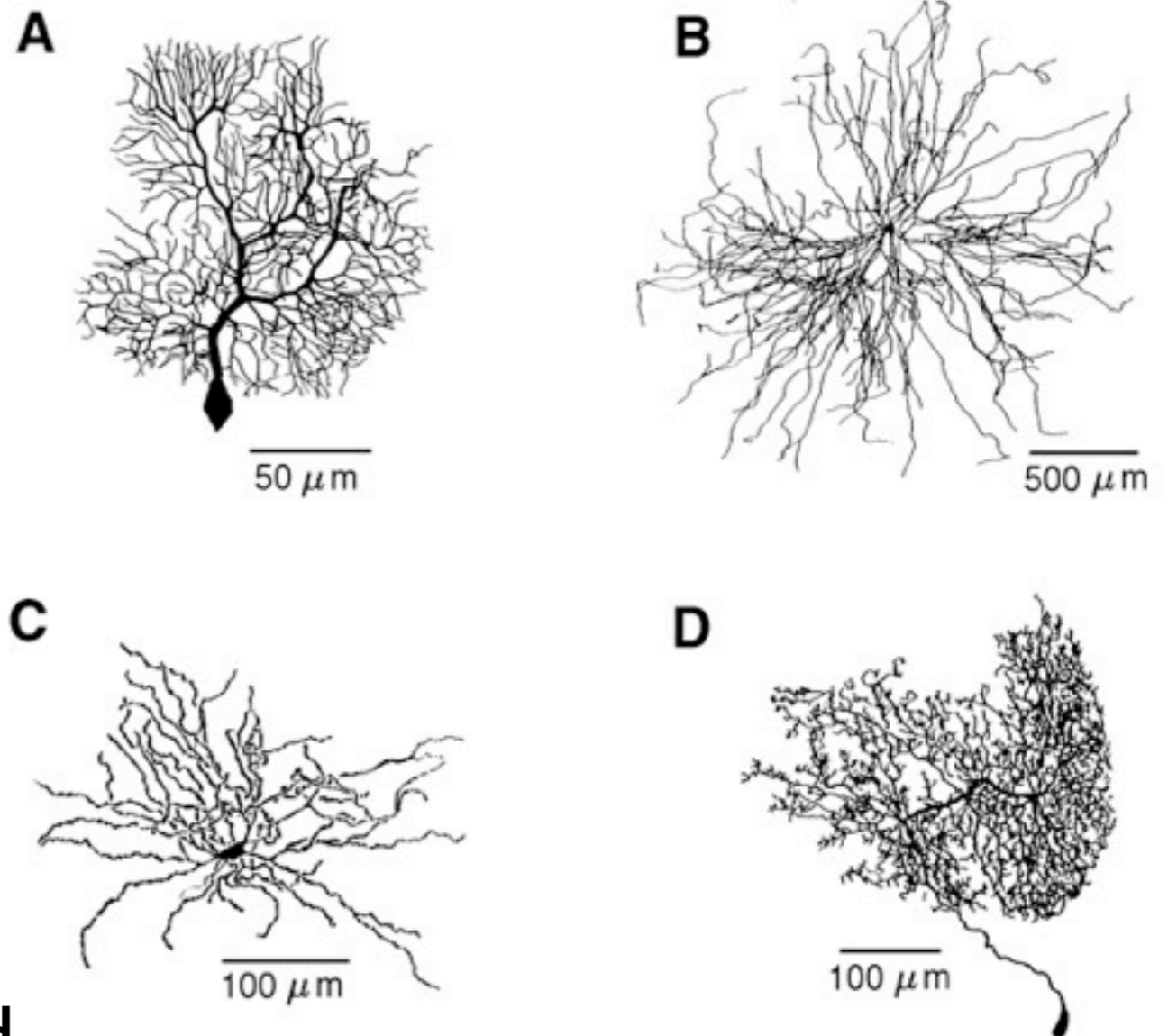
Superficial - cells change

Neurobiological:

sensory inputs combine with pre-existing cellular and network structure (and activity) through coordinated changes cellular morphology,

synaptic position, structure, and

efficiency, the integration of excitatory and inhibitory activity, firing rates, and timing \Rightarrow changes in the activity of the neural networks that is linked (somehow) to detailed memory.



omnis cellula e cellula

knowledge statements & performance expectations

core curriculum design / home

[Evolutionary basics](#)
[Biomolecules](#)
[Molecular machines](#)

[Cell theory](#)
[Cells as systems](#)
[Membrane structure](#)
[Membrane function](#)
[Cytoplasmic systems & intracellular organelles](#)
[Cell polarity: junctions & extracellular matrix](#)
[Genomic organization](#)
[Genes & their expression](#)
[Genetics](#)

[Signaling interactions](#)

[Cell & organismic replication](#)
[Cell death and disease](#)

[The social life of cells](#)

[Highlighter](#)

Rethinking cell & molecular biology curricula

The [design](#) of this project reflects many fruitful interactions, including discussions with faculty from the School of Education (CU Teach) and the general science education community.

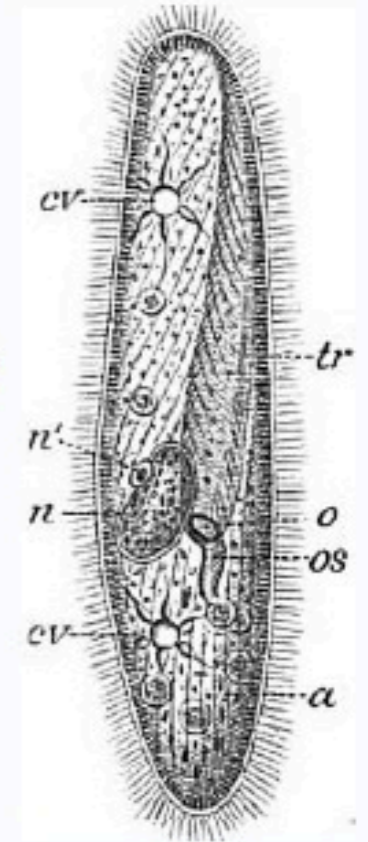
It builds on [lessons learned in the development](#) and evolution of [Biofundamentals](#), an "alternative" introductory course in modern biology (read a student's view of the course [here](#)). Biofundamentals itself was inspired by observations made during the National Science Foundation (NSF)-funded [Biology Concept Inventory \(BCI\)](#) project, a collaboration with Kathy Garvin-Doxas.

Equally critical have been the lessons learned in the course of developing [CLUE: Chemistry, Life, the Universe & Everything](#), an NSF-funded general chemistry curriculum. In particular, CLUE, Biofundamentals, and [OrganicPad](#) inspired the development of [BeSocratic](#), a novel graphics-based formative assessment system, made possible by the brilliant programming efforts of Sam Bryfczynski (now joined by Josiah Hester) and supported in part by funds from the NSF.

Biofundamentals, CLUE and the Cell & Molecular Biology project embrace technologies ([beSocratic](#) and [Highlighter](#)) that make the text socially interactive and evolvable and that provide students with challenging activities to hone their understanding of often [difficult ideas](#).

Finally, we acknowledge the HHMI CourseSource project team, which has emphasized the importance of conceptually coherent and pedagogically effective curricula in modern biology

Comments on the ideas presented here are very welcome
 - Mike Klymkowsky & Melanie Cooper.



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Performance expectations & knowledge statements
<http://virtuallaboratory.colorado.edu/Cell+Molecular/>

a cell biological performance expectation:

Students should be able to:

- i) develop a model for how proteins come to be localized in various parts of the cell; and

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- ii) use that model to predict the effects of mutation on protein localization.

a cell biological performance expectation:

Students should be able to:

- i) develop a model for how various proteins come to be localized in various parts of the cell;
- and ii) use that model to predict the effects of mutation on protein localization.

Should include:

- polypeptide synthesis occurs in the cytoplasm
 - it is vectorial (from N- to C-terminus).
- often continues until polypeptide's synthesis is complete
 - can be paused allowing movement to the endoplasmic reticulum (ER)
 - based on signal sequence/SRP and its interaction with ER-associated receptor/translocation complex
 - allows translation to resume, polypeptide enters ER membrane
- cytoplasmic polypeptides that are components of multisubunit proteins interact with molecular chaperones to assemble protein.
- presumes the pre-existence of cellular machinery (The Cell Theory).
- some cytoplasmic proteins remain cytoplasmic because they are too large to move (via diffusion) through the nuclear pore complex (NPC)
- some, small enough to pass through the NPC, accumulate within the nucleus because they bind to intranuclear components, such as chromatin.

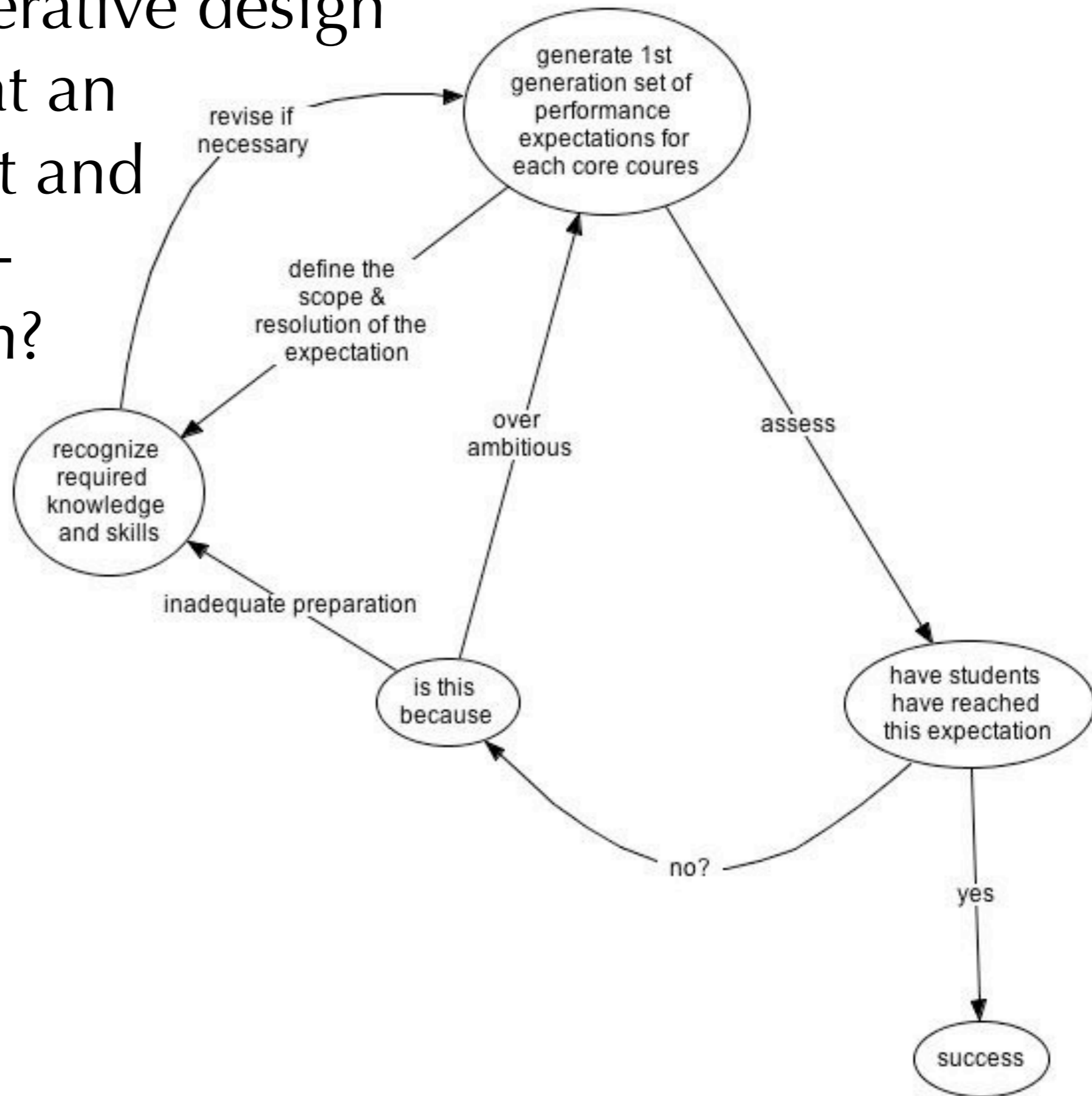
a cell biological performance expectation:

students should be able to develop a model for how various proteins come to be located in various parts of the cell, and based on that model predict the effects of various types of mutations on protein localization

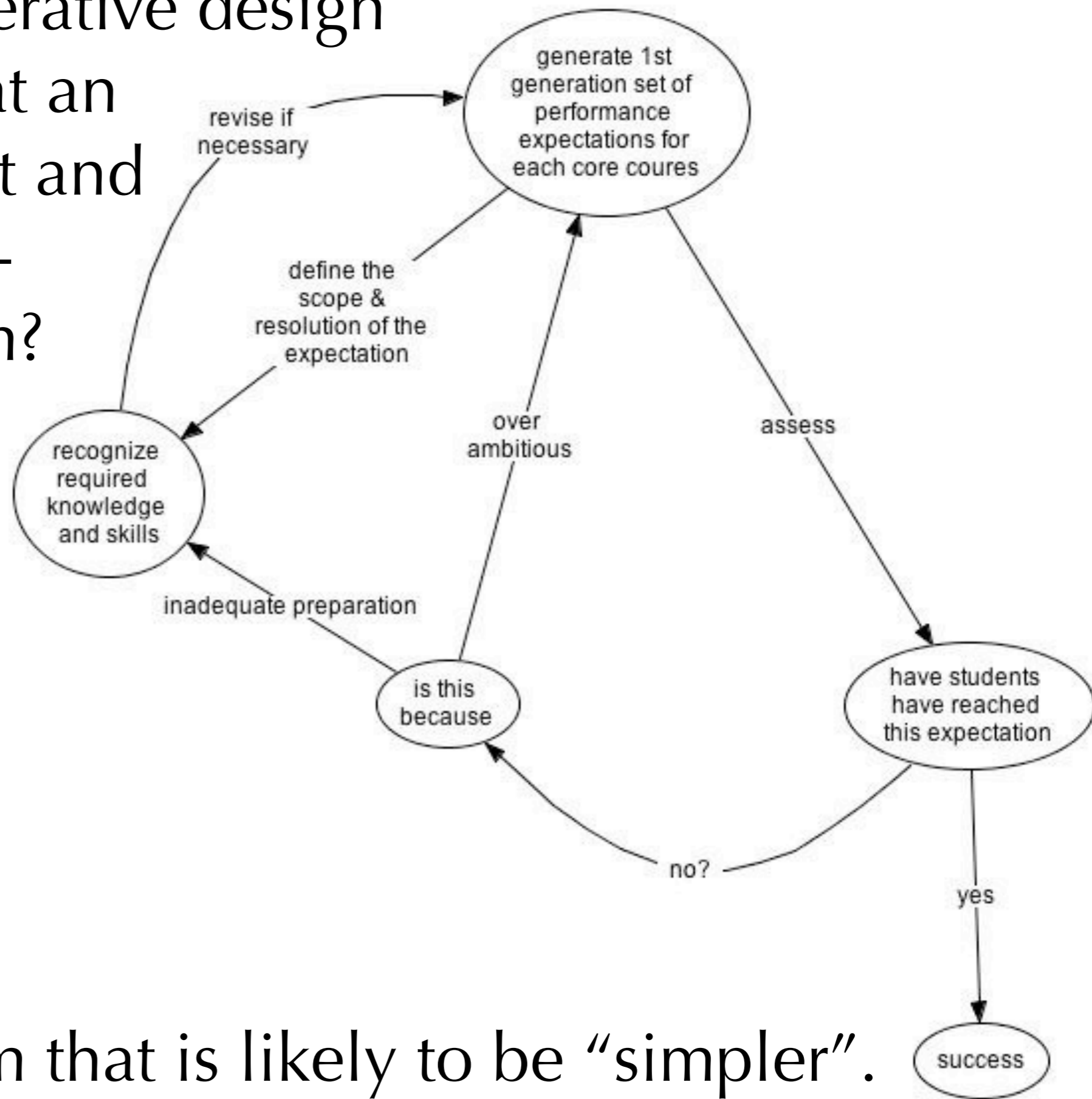
Should include (continue):

- Larger, nuclearly localized molecules rely on various active, that is, generally ATP-hydrolysis coupled, reactions, involving the NPC transport machinery and nuclear localization sequences (NLS).
- nuclear exclusion sequence (NES) lead to active transport out of the nucleus
- Both NLSs and NESs activity can be regulated
- macromolecular assembly can involve transport of components from the cytoplasm to the nucleus and then back to the cytoplasm.
- cytoplasmic proteins can be transported into intracellular organelles.
- ER protein can be localized to various various organelles or delivered to the plasma membrane
- proteins secreted from one cell can be taken up by other cells
- There are various “quality control” mechanisms that facilitate the return of mislocalized proteins to their correct locations or refolding, and if unsuccessful, the degradation of misfolded or mislocalized proteins

Can we use a reiterative design process to arrive at an effective, coherent and realistic “learning-based” curriculum?



Can we use a reiterative design process to arrive at an effective, coherent and realistic “learning-based” curriculum?



A curriculum that is likely to be “simpler”.

to be a little provocative....

should the needs of (say) a molecular biology curriculum ...

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“dictate” the goals of the physics, chemistry,
and mathematics courses we require our
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alternatively, can one be (scientifically) literate within out a
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alternatively, can one be (scientifically) literate within out a
foundational understanding of modern biology?

How can we cooperate effectively?
Can we all get along?

We are involved in three efforts at building coherent courses.

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BIOFUNDAMENTALS

(to replace intro. molecular biology)

We are involved in three efforts at building coherent courses.

BIOFUNDAMENTALS

(to replace intro. molecular biology)

CLUE-CHEMISTRY (w. *Melanie Cooper*)

(to replace intro. chemistry)

We are involved in three efforts at building coherent courses.

BIOFUNDAMENTALS

(to replace intro. molecular biology)

CLUE-CHEMISTRY (w. Melanie Cooper)

(to replace intro. chemistry)

CALCULUS, STOCHASTICS & MODELING (CSM)

(w. Eric Stade)(to replace calculus 1)

Biofundamentals™

[syllabus](#) | [home](#) | [blog](#)
course information - question

Being Biofundamental
Science & its Methods

Life's Origins

Evolution's logic

Speciation & Extinction

Adaptation & Selection

Predators, Prey & Mates

Non-Adaptive Processes

Homology & Analogy

Water & Life's Structure

Lipids & Membranes

Getting through Membranes

Carriers, Pore & Pumps

A (very) little thermodynamics

Capturing Energy

Storing Energy

Eukaryotic Symbiosis

Chemical basis of heredity

Nucleic Acid Structure

DNA replication

Mutations & Repair

Peptide Bonds & Polypeptides

Making Polypeptides

Assembling Proteins

Regulating Protein Activities

Regulating gene expression

Regulatory networks

Cell Divison

Life cycles & Sex

Stem Cells & Differentiation

Cellular communities

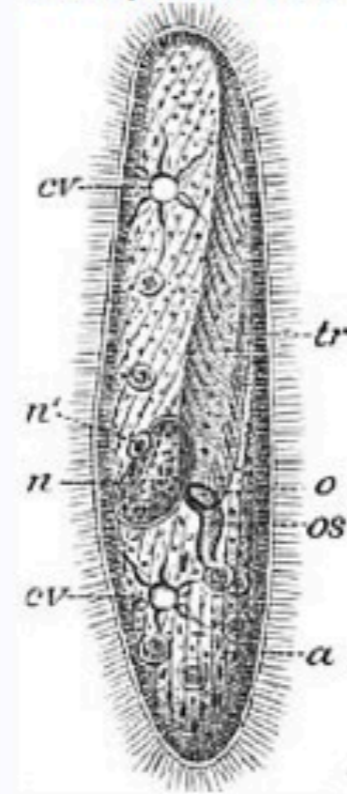
Eusocial and Antisocial

behavior

Biofundamentals™ (MCDB 1150) is an attempt to build a more conceptually coherent and rigorous introductory course in modern biology ([testimonial](#))

We use an interactive teaching style and web tools to "get Socratic". You need to read and engage with the text and embedded assignments **before** class.

To insure that you do, higher functions are incorporated throughout.



As you read you can leave comments and respond to the comments left by other students. Class time is spent considering the most difficult ideas.

Our goal is to help students master and apply difficult ideas, not sort them.

We use a novel testing strategy that includes "I know it now!©" tests designed to allow students to demonstrate their understanding of key ideas. →[IKiN LiNK](#)←

Biofundamentals™ is part of a larger course and curricular redesign effort described in more detail at the beSocratic.colorado.edu web site.

 **MERLOT**
Multimedia Educational Resource
for Learning and Online Teaching

Highlighter



 +1 3 people +1'd this

link to: [rethinking the cell & molecular biology curriculum](#)

LISTEN TO STUDENTS (talking to students)

because the population has no genetic memory of its size. Each generation is an independent event. The final result is that the population eventually drifts to fixation; change is possible; the population has become homozygous.

A different population, isolated from the first, also undergoes genetic drift. It may become homozygous for allele "A", whereas the first population became homozygous for allele "a".

As time goes on, isolated populations diverge from each other. The genetic variation originally present within populations now appears as differences between them (Suzuki et al., 1989. An Introduction to Genetic Analysis).

These non-selective, sampling-based effects are one reason why we see variation in whether a particular trait is adaptive or not. It really depends on the environment.

The end result of founder effects, bottlenecks, and genetic drift is that a trait is represented in a population by chance.

Questions to answer

1. Consider the various ways that the effects of a bottleneck might differ from those that can you identify?
2. Based on the Java Genetic Drift applet, is it possible for an allele to go from 5% to 100% of the population simply by genetic drift. The graph showed this.
3. How does selection act to limit the effects of genetic drift?
4. Is it possible for a genetic bottleneck to increase the frequency of a deleterious trait than before the bottleneck? How does this occur?
5. Assume that all members of a population that pass through a bottleneck have a deleterious trait; can the population survive and, if so, how would selection act on the population after the bottleneck?

Questions to ponder

- What limits the "size" of the founder effect or a bottleneck effect?
- Does passing through a bottleneck improve or hamper a population's ability to adapt to a new environment?

Based on the Java Ge[...]

Sort by Oldest

Share This Highlight Via Twitter Facebook Email

student 1 Posted 2 months ago

Based on the Java Genetic Drift applet, it IS possible for an allele to go from 5% to 100% of the population simply by genetic drift. The graph showed this.

Respond Share

student 2 Posted 2 months ago

I agree, especially in smaller populations

Respond Share

student 3 Posted 2 months ago

I also agree, no matter how unlikely it may be. However, within smaller populations, if the trait is truly advantageous it is more likely to be seen.

Respond Share

student 4 Posted 2 months ago

Even though this is possible, would it really happen in a population? If so, can anyone think of an example to help me understand?

Respond Share

student 5 Posted 2 months ago

Yes. but that happens under low possibility.

Respond Share

http://virtuallaboratory.colorado.edu/Biofundamentals/lectureNotes/Topic I E_Evo.htm

**USE Multiple FORMS OF
“CONCEPTUAL ASSESSMENT”**



Biological Concepts Instrument (BCI): A diagnostic tool for revealing student thinking

Michael W. Klymkowsky, Sonia M. Underwood, R. Kathleen Garvin-Doxas

(Submitted on 20 Dec 2010)

A key to effective teaching is an awareness and accurate understanding of the thinking and implicit assumptions that students bring to the subject to be learned. In the absence of extensive Socratic interactions with students, one strategy to assess student thinking involves the use of concept inventories (CIs). CIs are typically multiple-choice assessments, constructed based on research into student thinking and language, and designed to reveal the presence of common misconceptions and implicit assumptions pertaining to a particular facet of a subject. Here we describe the open-source Biological Concepts Instrument (BCI), a diagnostic, multiple-choice instrument designed to provide instructors with a preliminary map of a number of basic ideas in molecular level biology. We describe the strategy behind its design, the research upon which it is based, item construction, and its possible uses as a means to reveal and address persistent and often unrecognized conceptual obstacles.

Text

Subjects: **Other Quantitative Biology (q-bio.OT)**

Cite as: [arXiv:1012.4501v1](https://arxiv.org/abs/1012.4501v1) [q-bio.OT]



BCI word cloud

Molecular level thinking

Q25: Imagine an ADP molecule inside a bacterial cell. Which best describes how it would manage to "find" an ATP synthase so that it could become an ATP molecule?

- a. It would follow the hydrogen ion flow.
- b. The ATP synthase would grab it.
- c. Its electronegativity would attract it to the ATP synthase.
- d. It would actively be pumped to the right area.
- e. Random movements would bring it to the ATP synthase.

Molecular level thinking

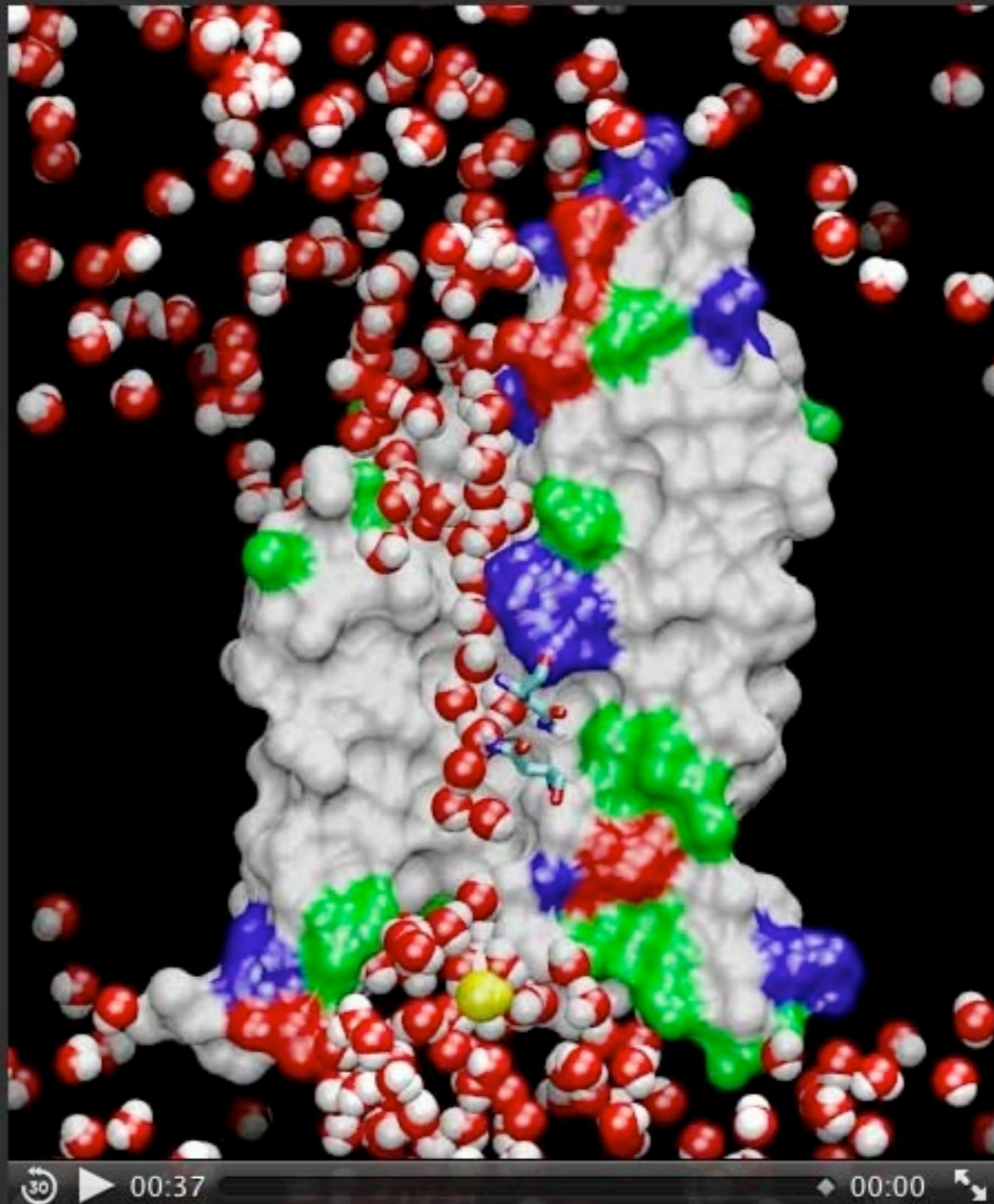
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- e. Random movements would bring it to the ATP synthase.

Diffusion and drift group (Questions 1, 5, 25, 29, 30):



Random processes seem impossible



http://www.nobelprize.org/nobel_prizes/chemistry/laureates/2003/chemanim1.mpg

Article

Understanding Randomness and its Impact on Student Learning: Lessons Learned from Building the Biology Concept Inventory (BCI)

Kathy Garvin-Doxas* and Michael W. Klymkowsky[†]

*Center for Integrated Plasma Studies and [†]Molecular, Cellular, and Developmental Biology Department, University of Colorado, Boulder, CO 80309

Submitted August 23, 2007; Revised January 14, 2008; Accepted February 7, 2008
Monitoring Editor: Bruce Alberts

CBE—Life Sciences Education
Vol. 7, 227–233, Summer 2008

Because mutations are random, the cannot lead to useful effects; either evolution cannot occur or must require (supernatural) guidance!

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The difficult of randomness

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Our suggestion:

Address the issue head on, illustrate/emphasize that stochastic processes (however silly they look) “work” at the molecular level.

What do responses tell us?

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BCI Q: 23 An individual, "A", displays two distinct traits. A single, but different gene controls each trait. You examine A's offspring, of which there are hundreds, and find that most display either the same two traits displayed by A, or neither trait. There are, however, rare offspring that display one or the other trait, but not both.

- A. The genes controlling the two traits are located on different chromosomes.
- B. The genes controlling the two traits are located close together on a single chromosome.
- C. The genes controlling the two traits are located at opposite ends of the same chromosome.

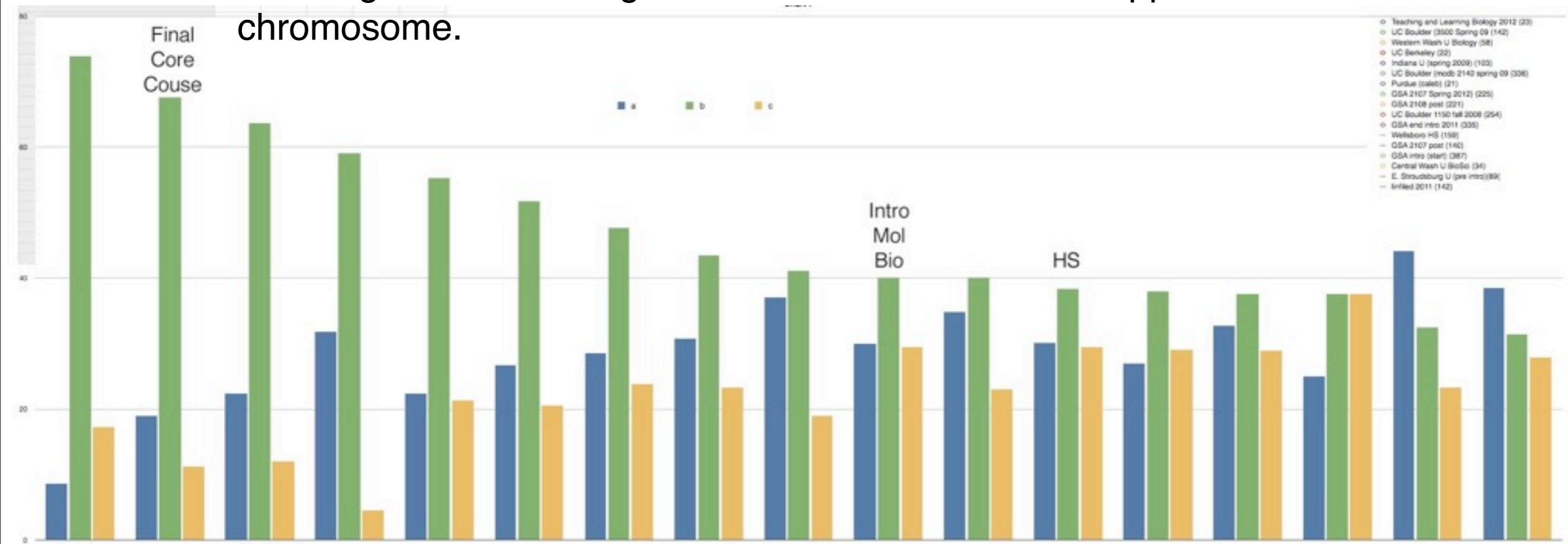
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A. The genes controlling the two traits are located on different chromosomes.

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C. The genes controlling the two traits are located at opposite ends of the same chromosome.



Leaves unanswered, whether students understand why (biologically) gene linkage is important?

Look at other student thinking other ways....

Look at other student thinking other ways....

Using more informative instruments:

Look at other student thinking other ways....

Using more informative instruments:

graphic analysis (Trujillo et al 2012. *BAMBED* **40**:100)

Look at other student thinking other ways....

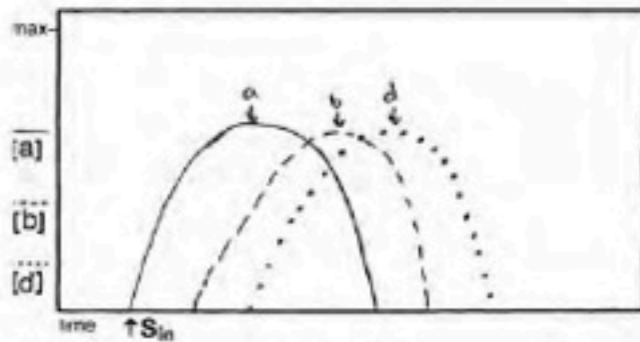
Using more informative instruments:

graphic analysis (Trujillo et al 2012. *BAMBED* **40**:100)

Look at other student thinking other ways....

Using more informative instruments: graphic analysis (Trujillo et al 2012. BAMBED 40:100)

- a, b, and d will decrease over time because of the negative feedback loop
- a will be made before b, which will be made before d
- a will be at its maximum level when d starts to be produced
- d will decrease after b decreases, which will decrease after a decrease
- when d is high, a is low

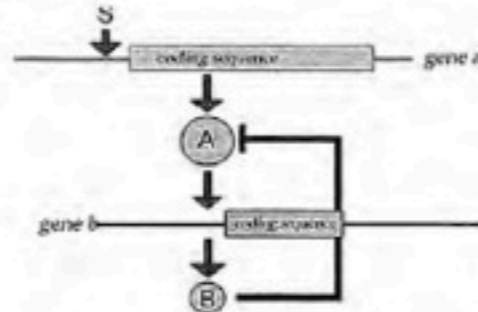


a

Now for one final complexity. Let us assume that polypeptide B acts negatively on polypeptide A.

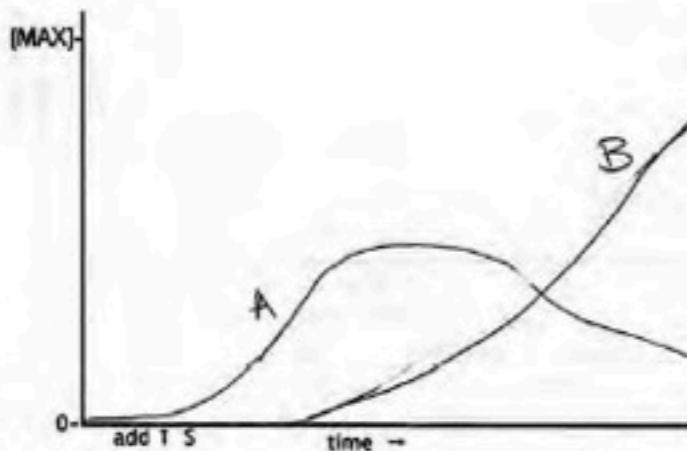
This negative interaction could involve inhibition of A's activity or it could increase the rate at which A is degraded.

b

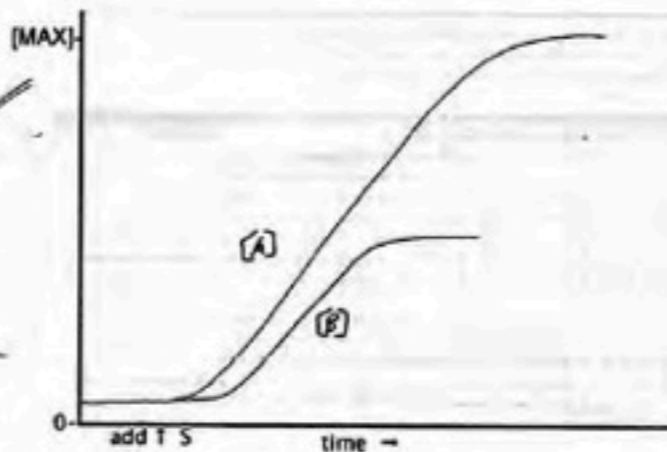


Draw a graph of [A] and [B] over time. As before expression of gene a begins with the addition of S (and S remains present throughout the experiment).

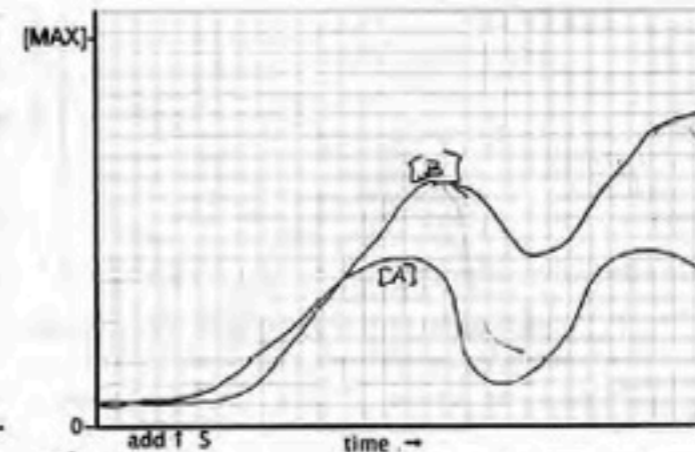
Indicate which assumption you are making:
 B enhances the degradation of A or B inhibits the activity of A.
 Describe how your choice influences your graph.



b



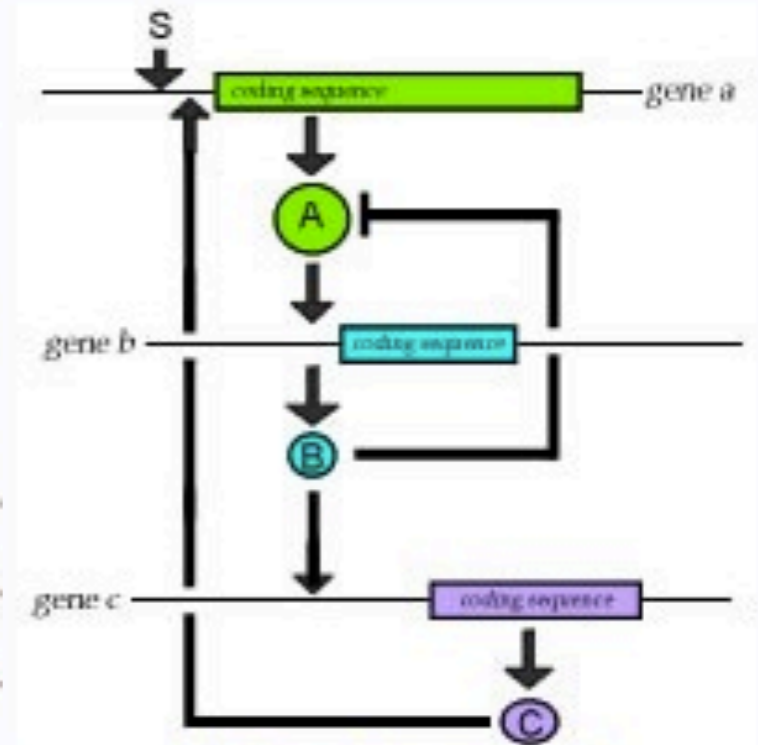
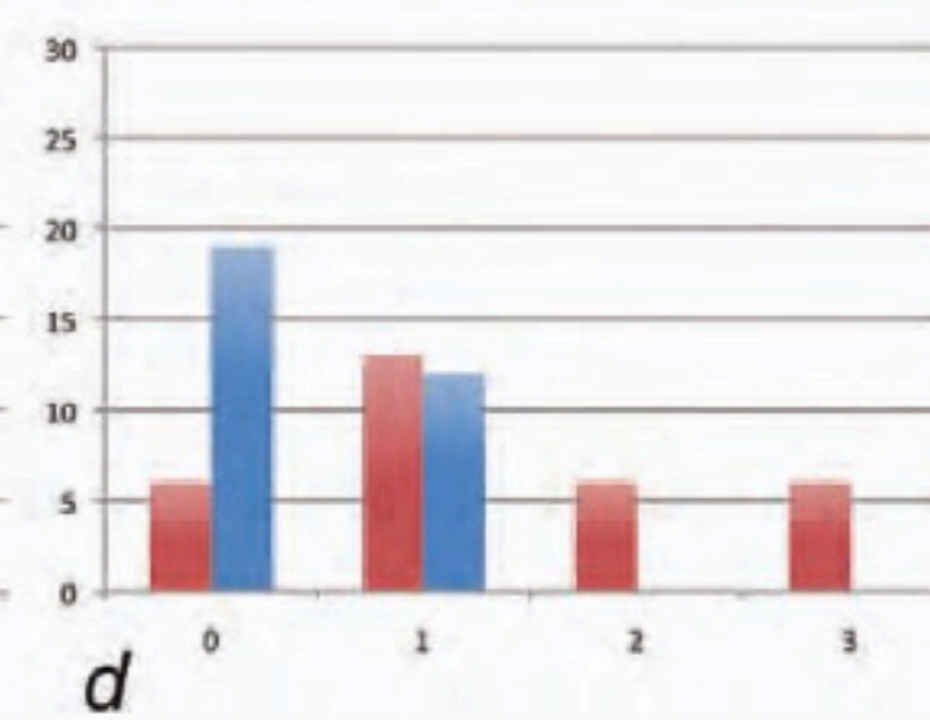
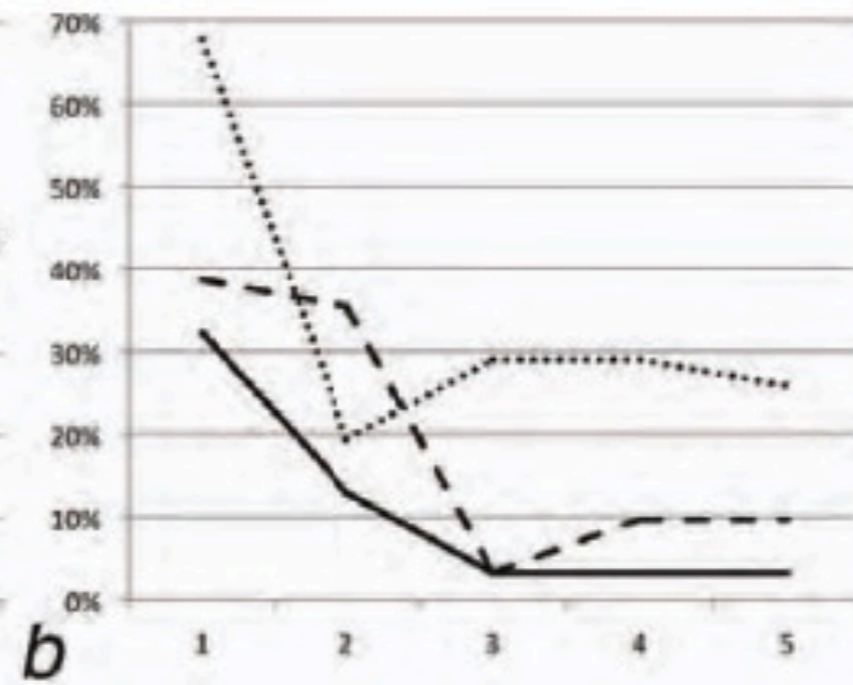
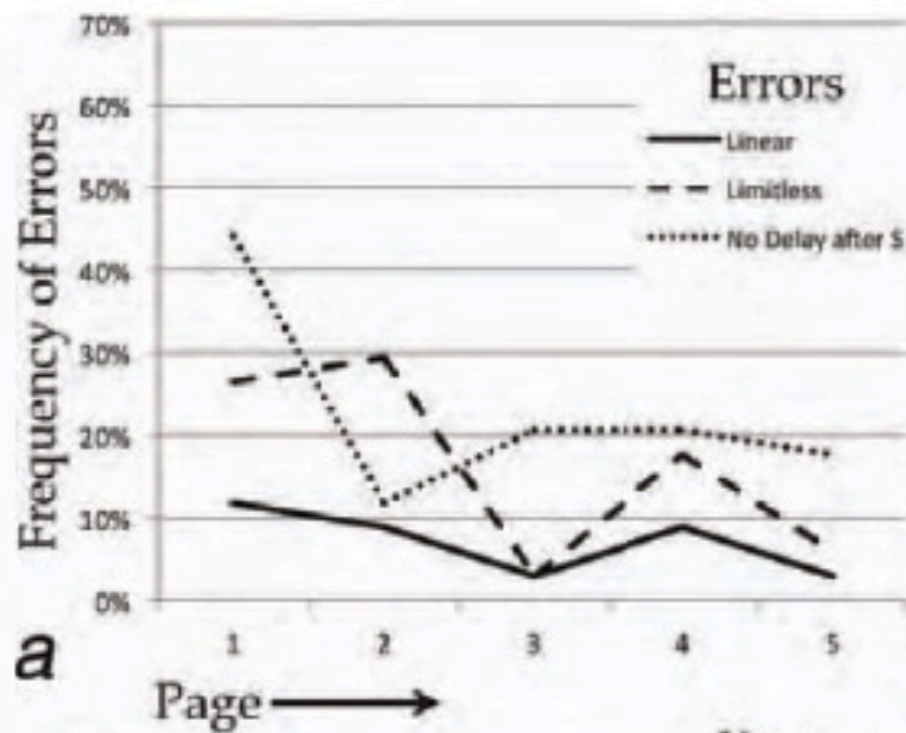
c



d

Discovering student thinking ...

Thinking about networks:




Go! to the tutorial

BeSocratic

http://besocratic.clemson.edu/ Reader Google

News Lab PubMed CLUE arts Dish BioFundamentals

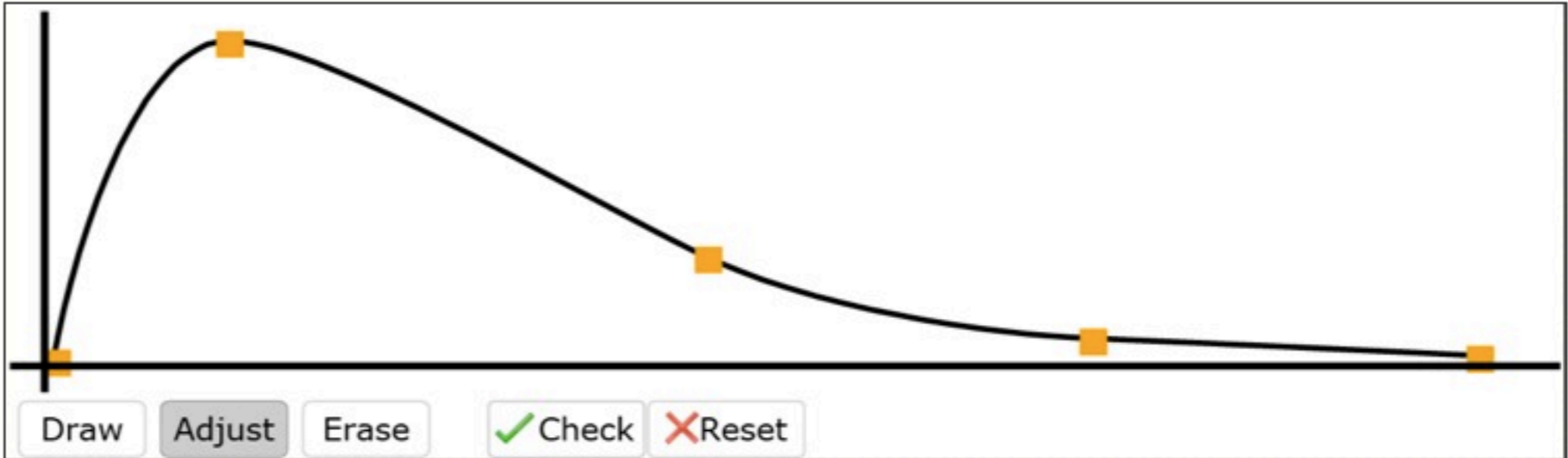
 **BeSocratic™**
Socratic Activities and Feedback

[Sign In](#) | [Sign Up for Free](#)

[Home](#) [Features](#) [About](#) [Demos](#) [Videos](#) [Help](#)

Meaningful Assessment of Free-form Drawings

BeSocratic is a flexible, web-based system that recognizes and responds to free-form student input. Using a pen or touch interface, students may respond naturally to questions posed by the system. At the same time, the structures that students draw are rigid enough that they may be automatically evaluated. This allows the applications to prompt the student with multi-tiered feedback when the students are having difficulties. Furthermore, BeSocratic allows for teachers to analyze the student work in a variety of meaningful ways and give insights into a student's thought process.



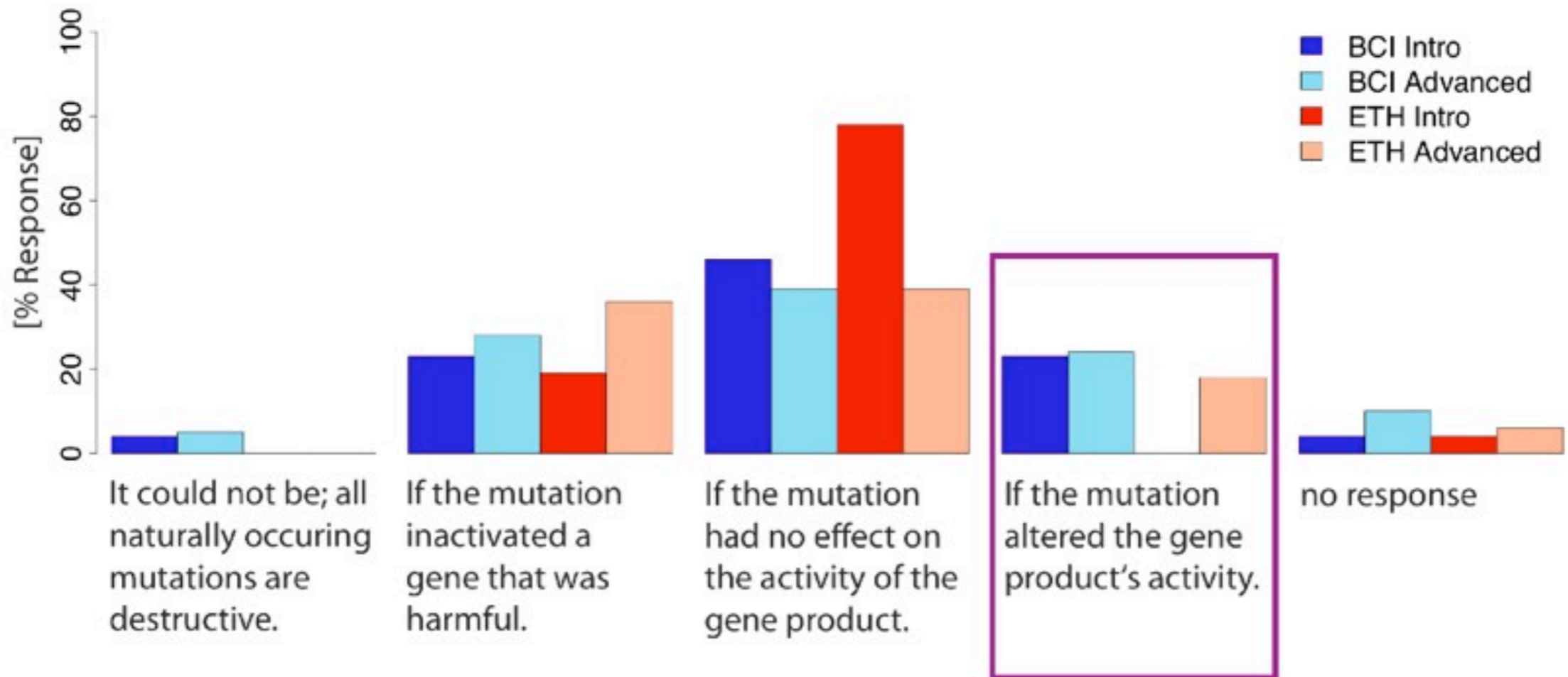
We are developing tutorials and formative assessment activities based on the available evidence and

Discussed further by Melanie Cooper (tomorrow)

Look at other student thinking other ways....



Look at other student thinking other ways....



Using more informative instruments

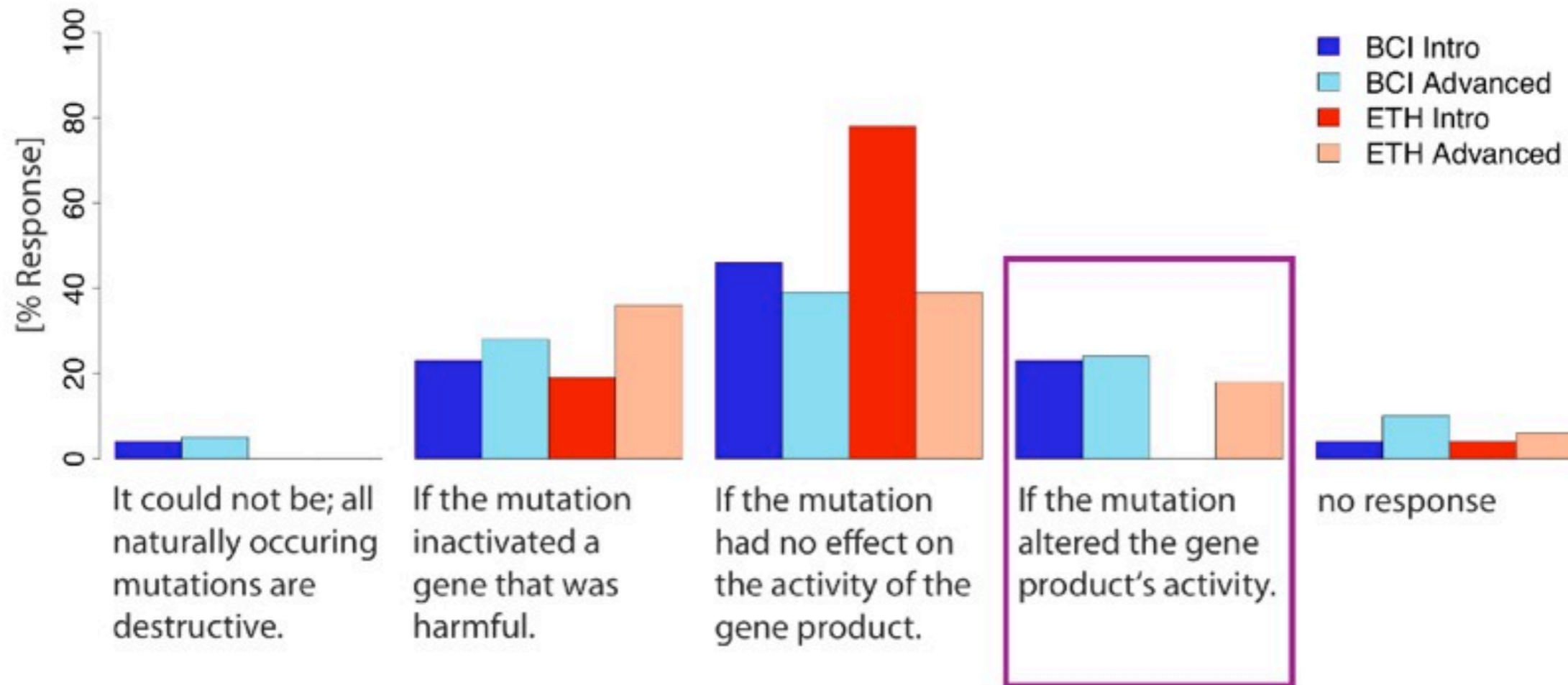
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Using more informative instruments

- textual analysis (Hensen et al 2012. Biology Open, online)

Look at other student thinking other ways....



Using more informative instruments

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Provide context through Muller's morphs

(1932). Further studies on the nature and causes of gene mutations. Sixth Int. Cong. Genet. 1, 213–255.

Provide context through Muller's morphs

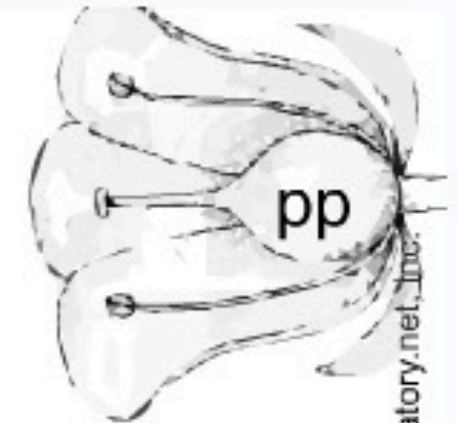
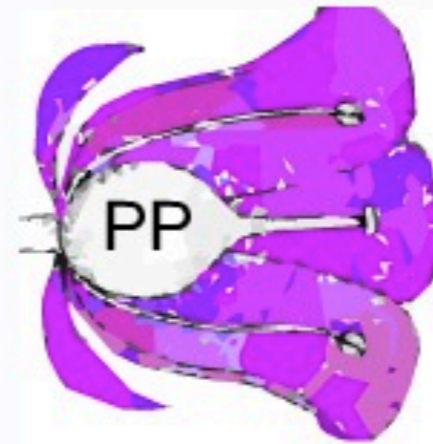
(1932). Further studies on the nature and causes of gene mutations. Sixth Int. Cong. Genet. 1, 213–255.

Mendel's Factors & Muller's mutations – page 1 of 9

As you almost certainly already know (and can [review here](#)), in the 1800's Mendel deduced the existence of genetic factors (which we now know as genes).

These factors are passed unaltered from parent to offspring - but importantly, not all factors held by a parent are transmitted ([note on simplifications](#)).

Each parent has two copies of each gene, but one and only one copy is transmitted to any particular offspring. Which copy is transmitted is random (stochastic).



Mendel's hypothesis:
Pure-bred lines carry two copies of the factor that determines flower color.



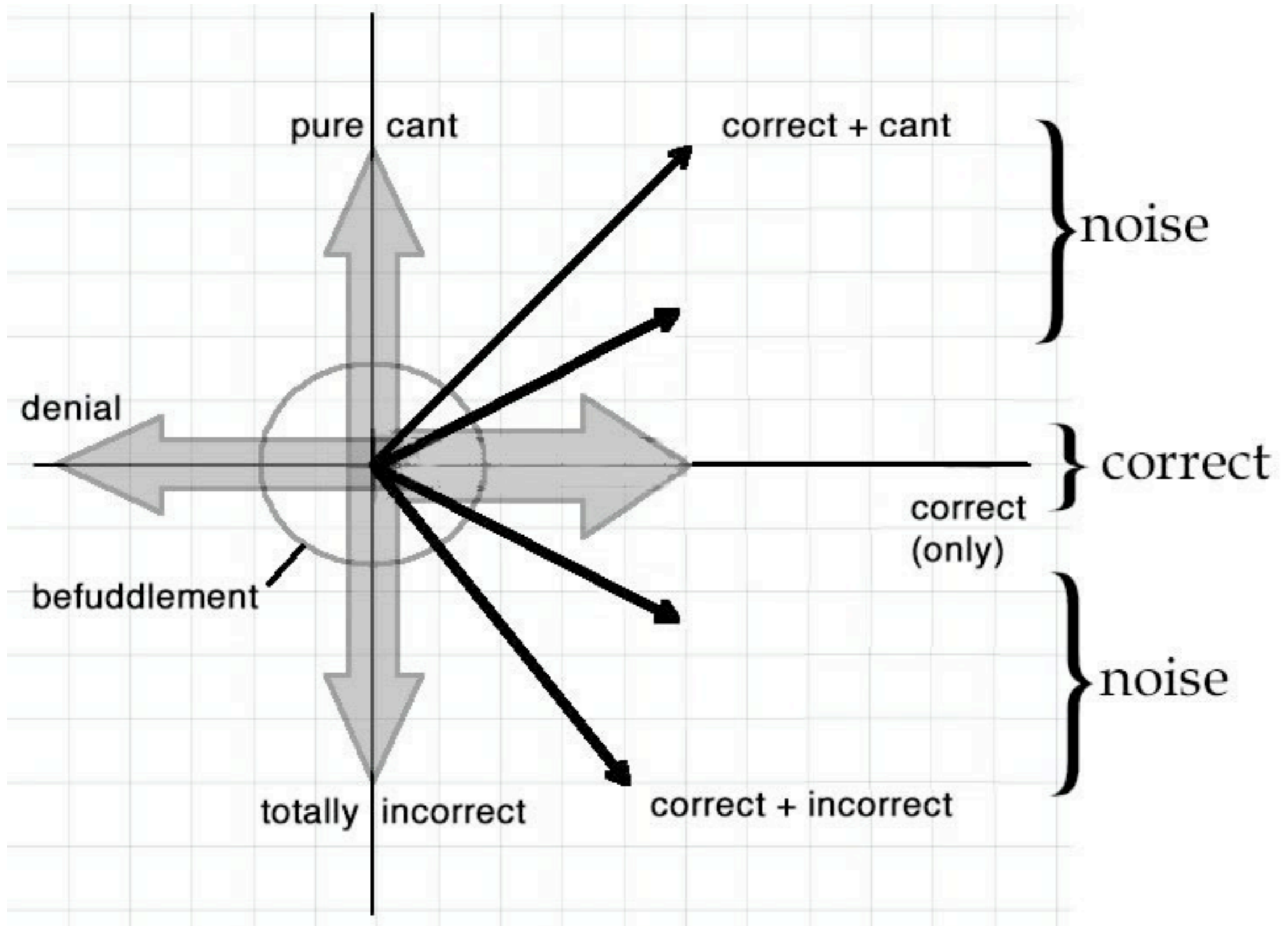
©2001 - virtuallaboratory.net, Inc.

As you answer these questions, remember, there is rarely a single correct answer.

What did Mendel know about the physical nature of his factors?

Submit/Next

Coding student responses



Henson, Cooper & Klymkowsky. 2012. Turning randomness into meaning at the molecular level using Muller's morphs. *Biology Open*, in press.

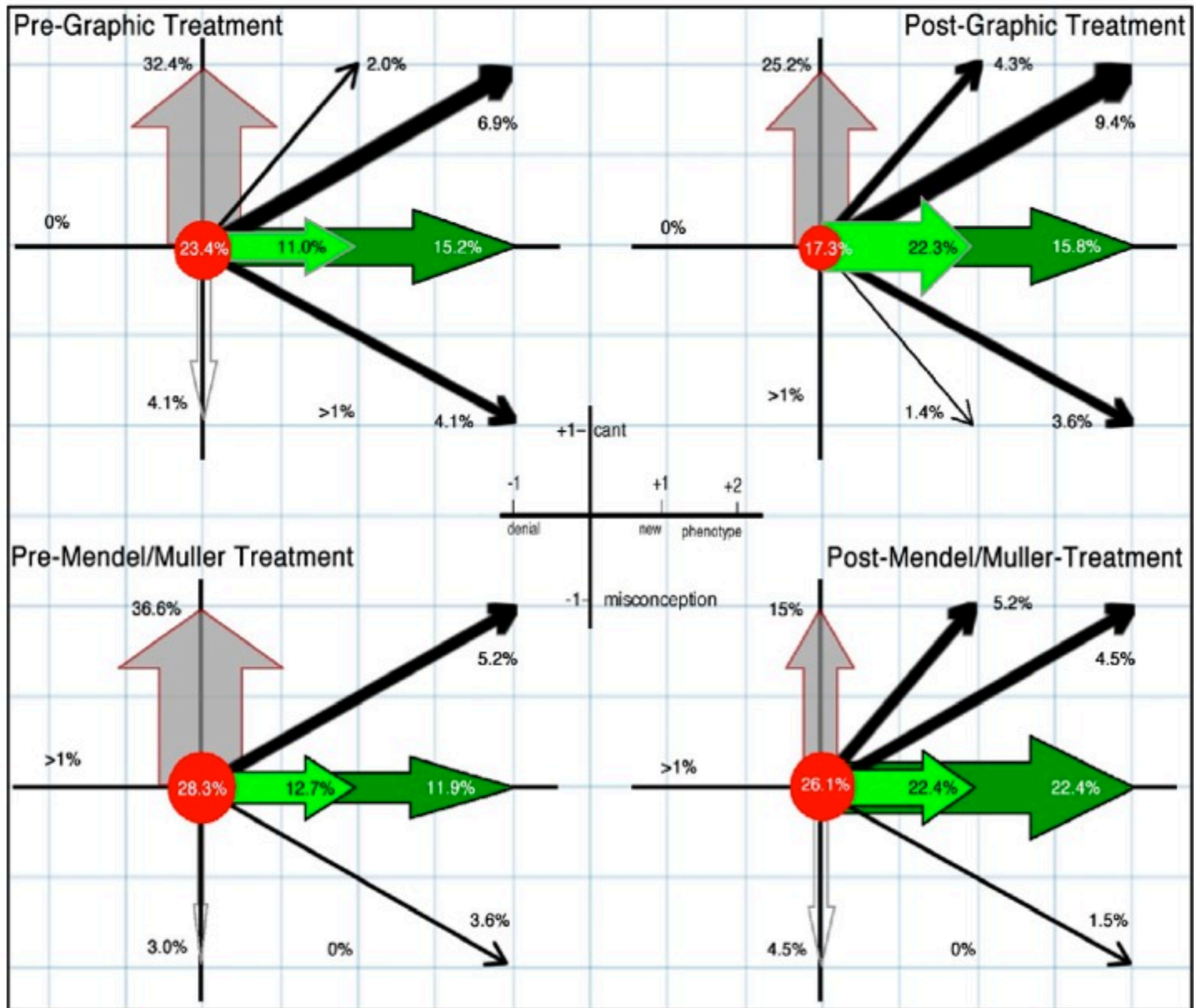


Fig. 3. Changes in student thinking. Students were asked to work through either the Graphical Thinking (top panels) or the Mendel/Muller (bottom panels) activities in groups. Student responses to the “How might a mutation be creative?” question pre- (left panels) and post- (right panels) treatment were analyzed.

“The puzzling conclusion is that although individual FCI responses are not reliable, the FCI total score is highly reliable.”

“The puzzling conclusion is that although individual FCI responses are not reliable, the FCI total score is highly reliable.”

“This study confirms that the total FCI score reliably measures a single concept, although our analysis is silent as to the nature of this concept.”

The under appreciated pitfalls of various instruments (channeling Michael Whitman)

The under appreciated pitfalls of various instruments (channeling Michael Whitman)

The story was always the same: our ability to predict performance at the school was negligible. Our forecasts were better than blind guesses, but not by much. -
Daniel Kahneman

WYSIATI: “What you see is all there is.”

WYSIATI: “What you see is all there is.”

Simple check:

Ask students to explain why “incorrect” choices are wrong.

Control Chemistry

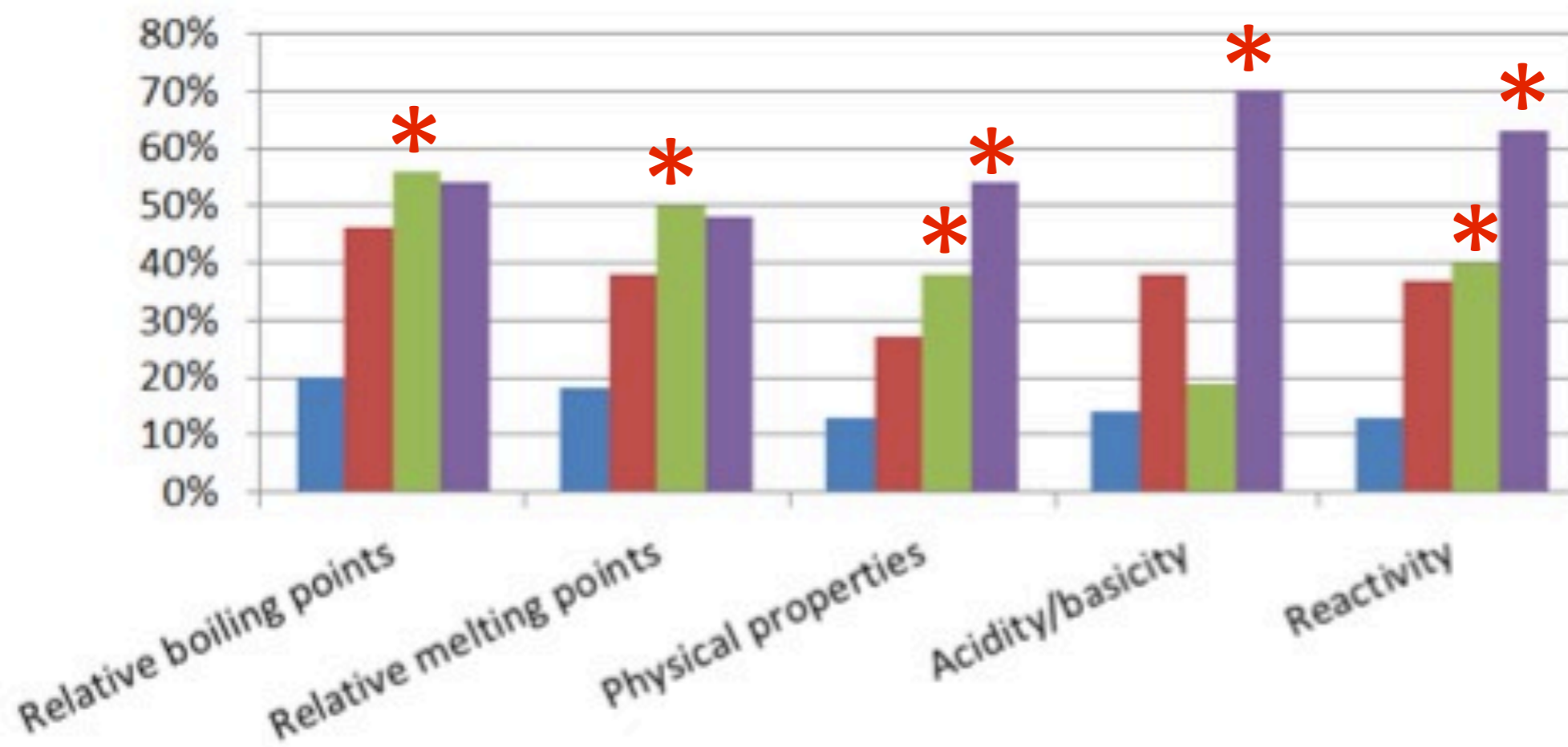
Post Fall Control

End of Spring Control

Post Fall Treatment

End of Spring Treatment

CLUE Chemistry

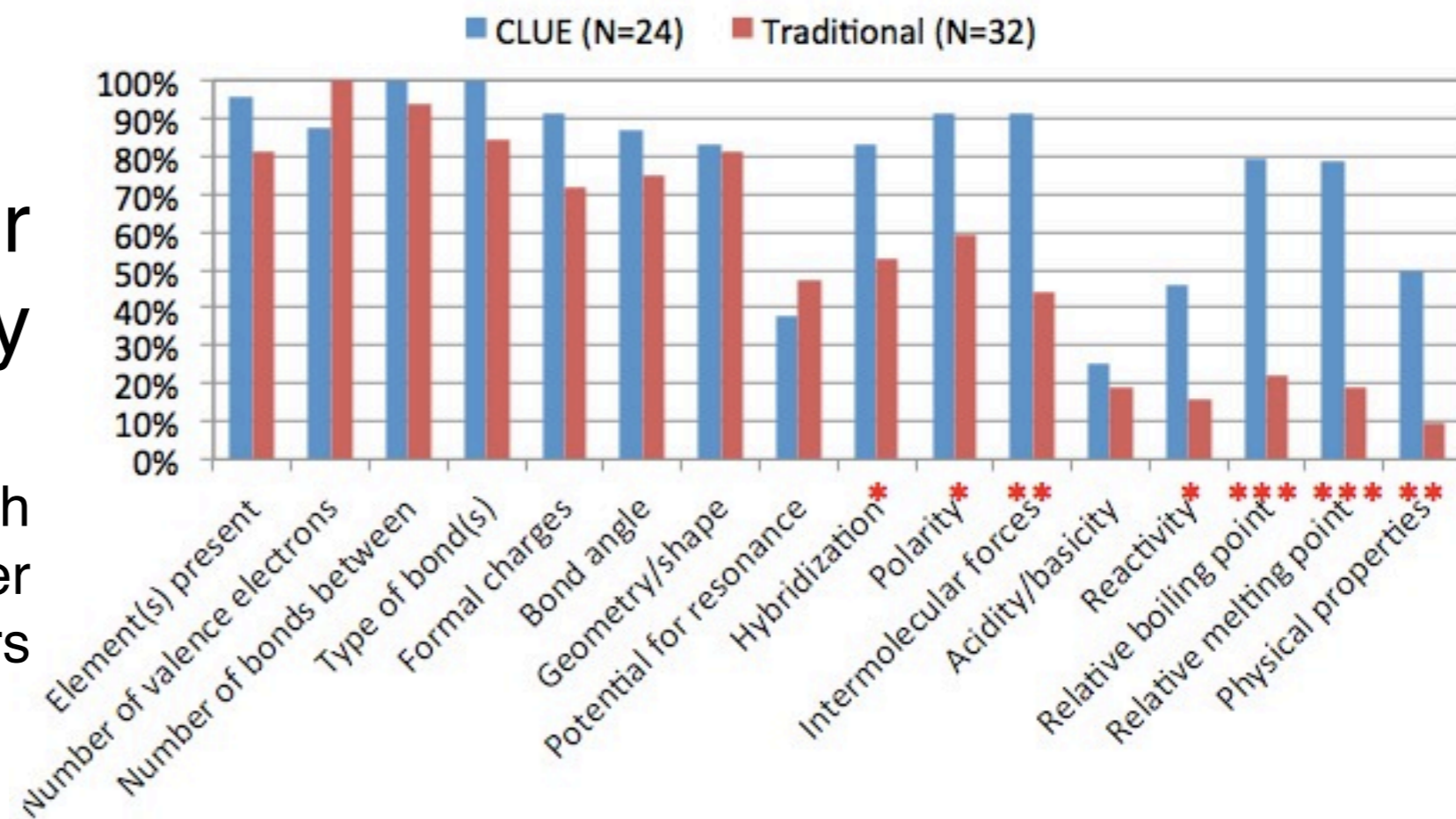


Chemical Formula	Post Fall Control	Post Fall Treatment	p-value	End of Spring Control	End of Spring Treatment	p-value
Relative boiling points	20%	56%	< .001	46%	54%	.512
Relative melting points	18%	50%	< .001	38%	48%	.311
Physical properties	13%	38%	< .001	27%	54%	.003
Acidity/basicity	14%	19%	.408	38%	70%	< .001
Reactivity	13%	40%	< .001	37%	63%	.005

Development and Assessment of a Molecular Structure and Properties Learning Progression, Cooper, Underwood, Hilley & Klymkowsky, "in press"

After first semester of intro. chemistry

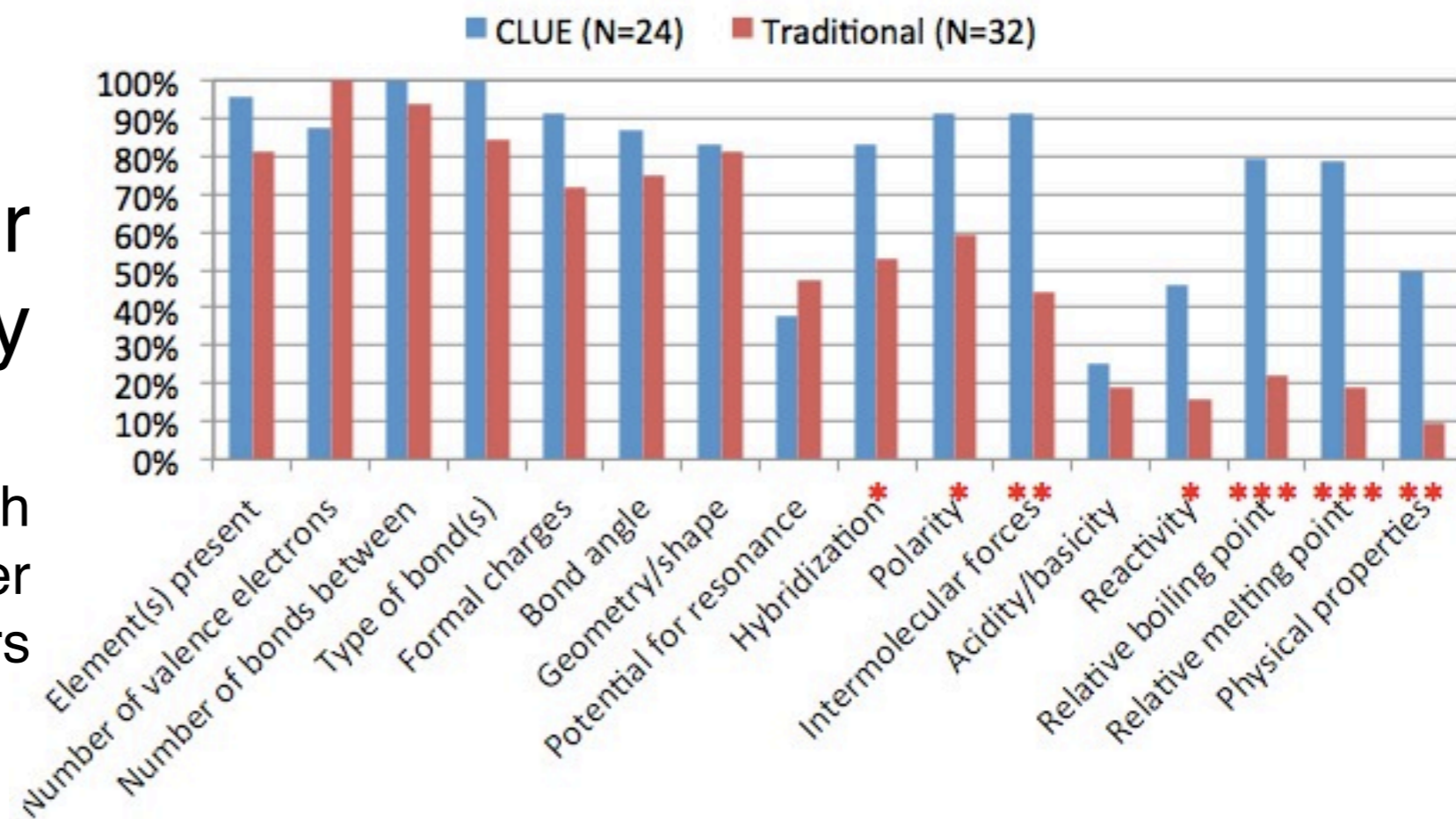
n.b. started with 90 in each group and followed them over two years



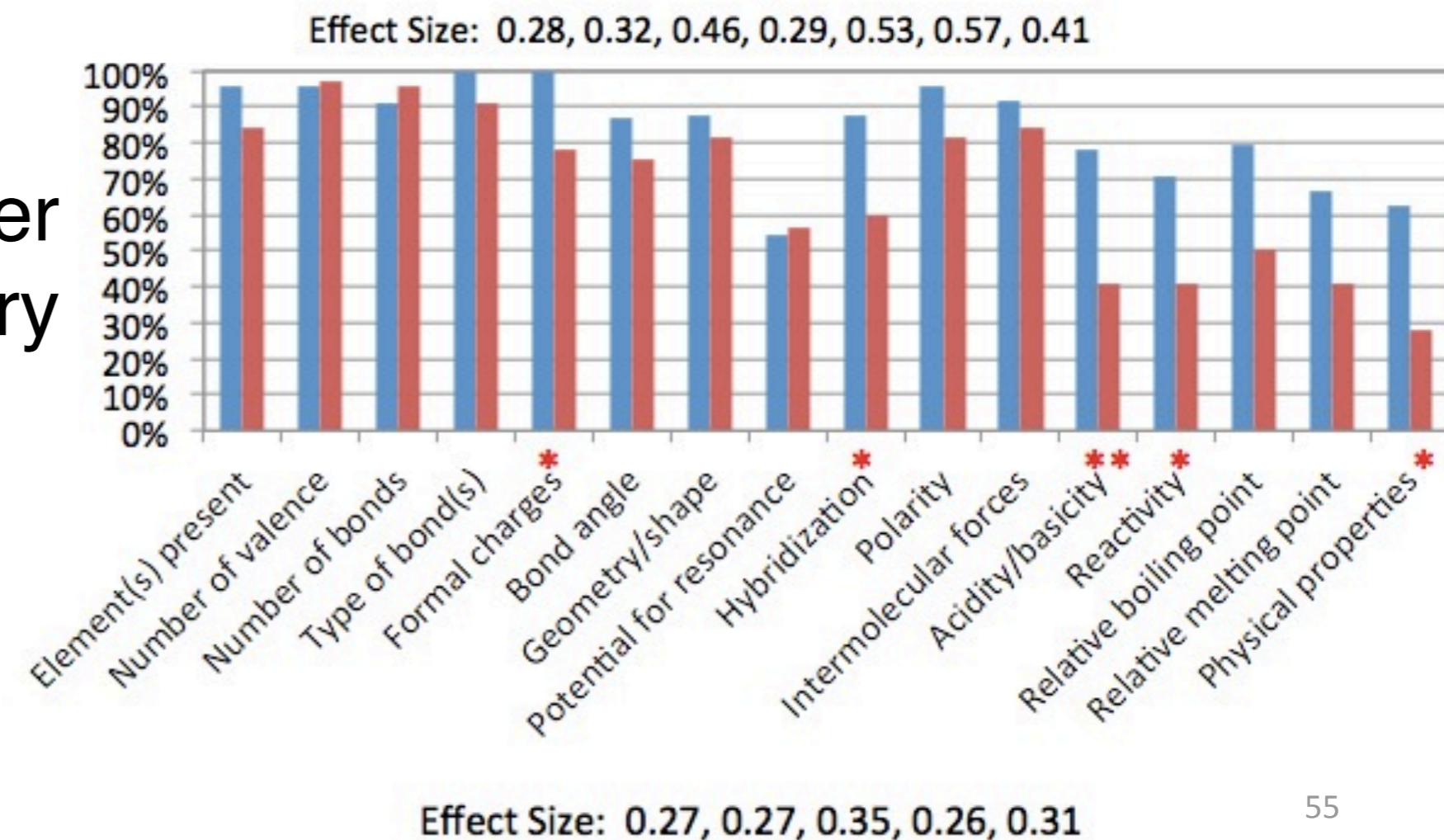
Effect Size: 0.28, 0.32, 0.46, 0.29, 0.53, 0.57, 0.41

After first semester of intro. chemistry

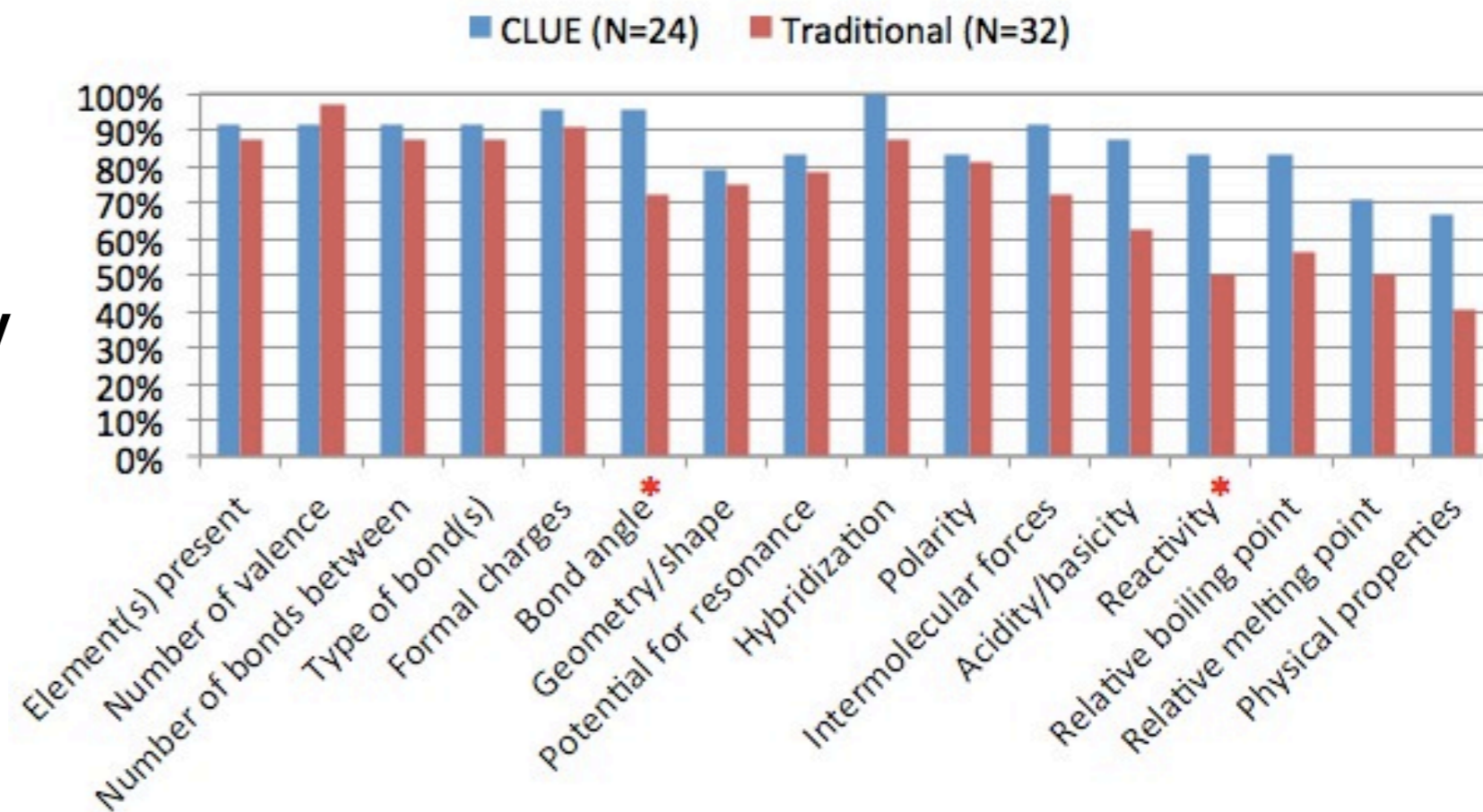
n.b. started with 90 in each group and followed them over two years



After second semester of intro. chemistry

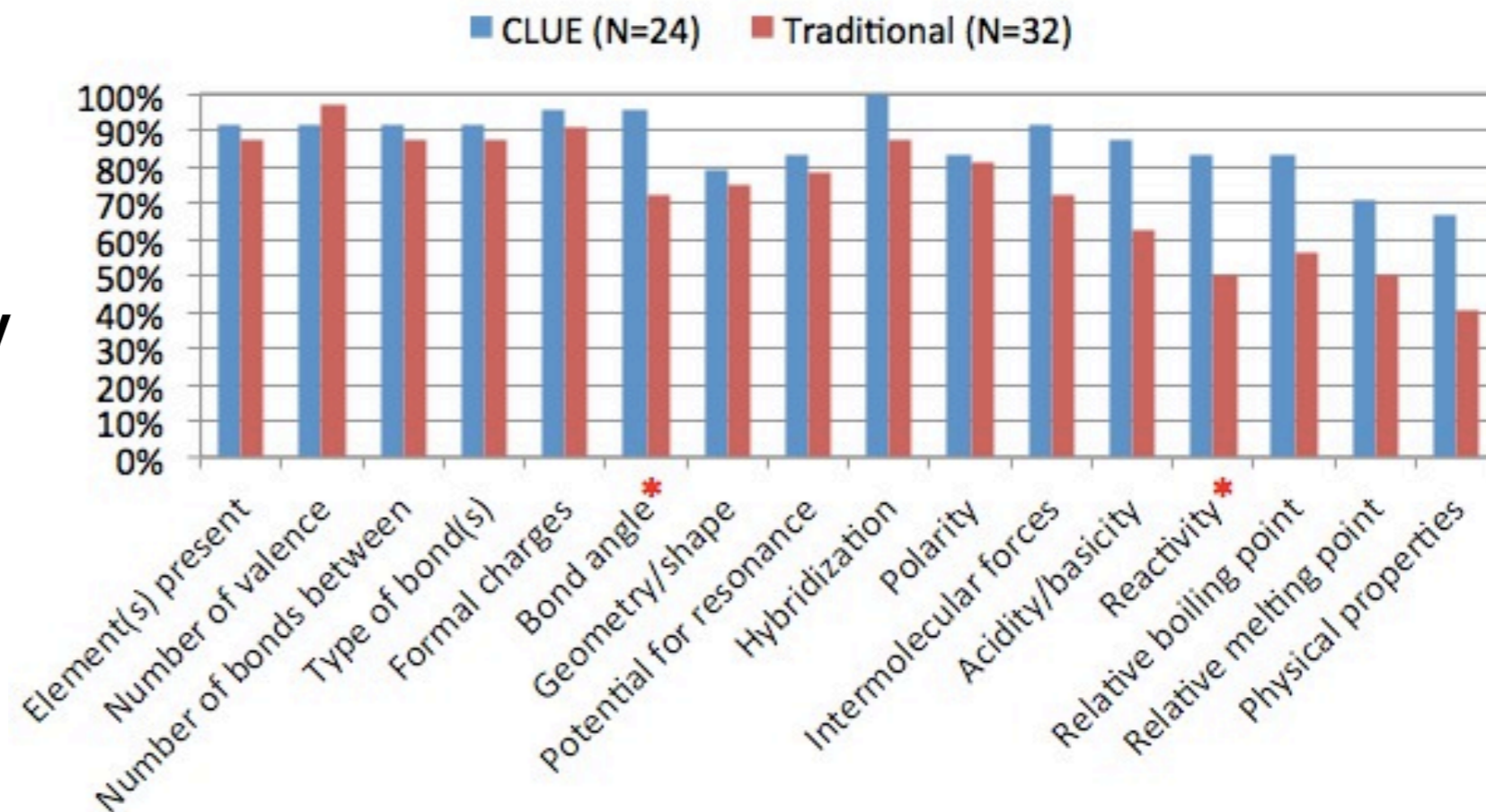


After one semester of organic chemistry



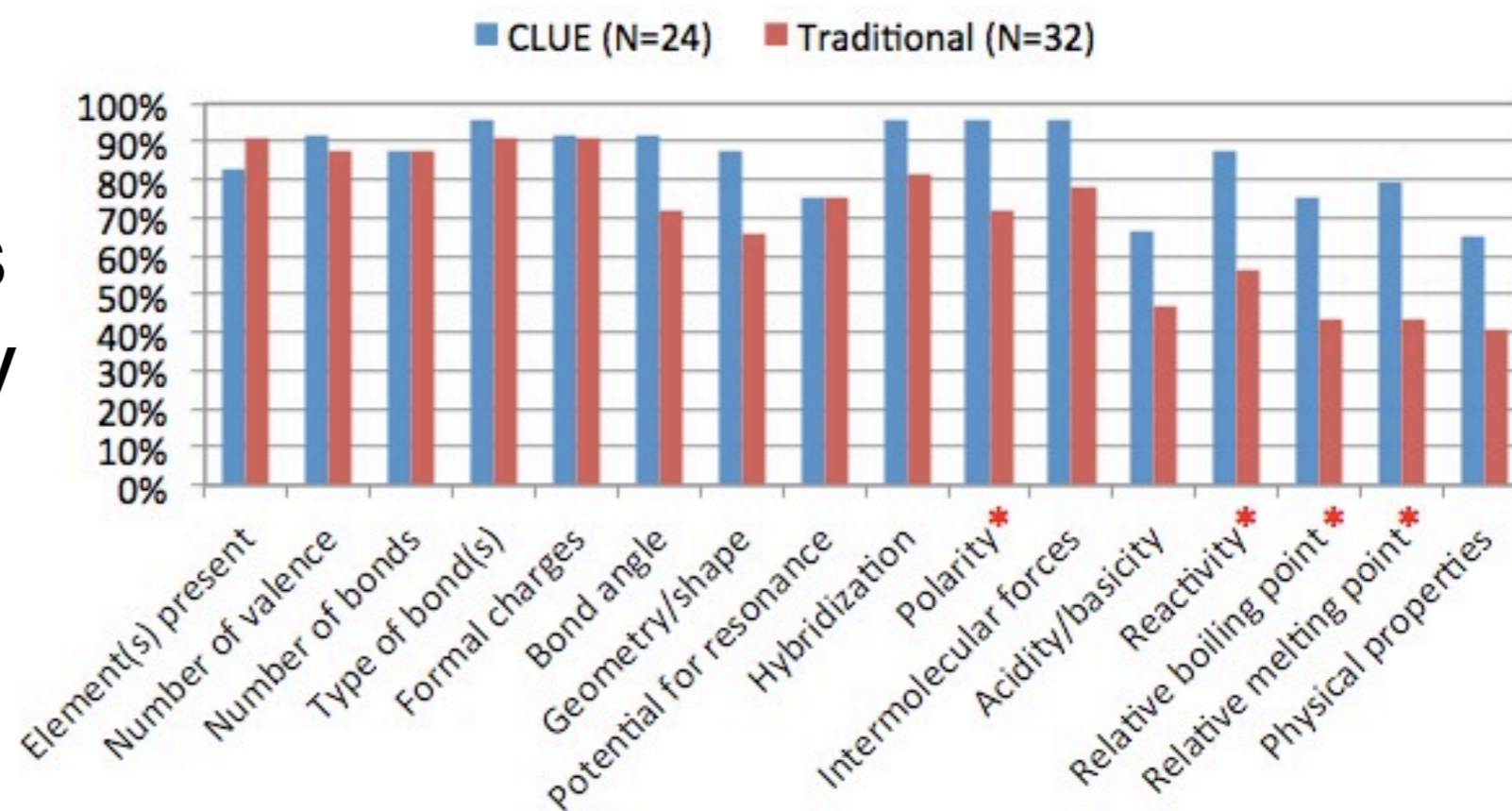
Effect Size: 0.26, 0.30, 0.28, 0.32

After one semester
of organic chemistry



Effect Size: 0.26, 0.30, 0.28, 0.32

After two semesters
of organic chemistry



Effect Size: 0.26, 0.30, 0.28, 0.32

Summary:

We have evidence that coherently designed curriculum are more effective in promoting student learning.

Course/curriculum need to be:

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Course/curriculum need to be: **relevant**
rigorous
coherent

Summary:

We have evidence that coherently designed curriculum are more effective in promoting student learning.

Course/curriculum need to be: **relevant**
rigorous
coherent
realistic