

# Recitation 01

Principles, Ethics, & Practices



MAS.885  
Spring 2026

HOW TO GROW  
(ALMOST) ANYTHING

**Instructors**  
David S. Kong (MIT Media Lab)  
George Church (Harvard)  
Joseph Jacobson (MIT Media Lab)



# Agenda

**Semester Preview & Homework 01 Requirements** - 1 min (David, Ronan)

**Atlas Training** - 5 min (Suvin)

**Principles, Ethics, Practices HW** - 10 min (Sebastian)

**Small Group Discussion** - 10 min (Students)

**Student Website Live Demo** - 25 min (Greg)

**Discourse Introduction** - 10 min (Derek)

**Homework 01 Requirements Review** - 2 min (Ronan)

**Lab Safety Training** - 30 min, Bldg 68 (All)

# Semester Preview



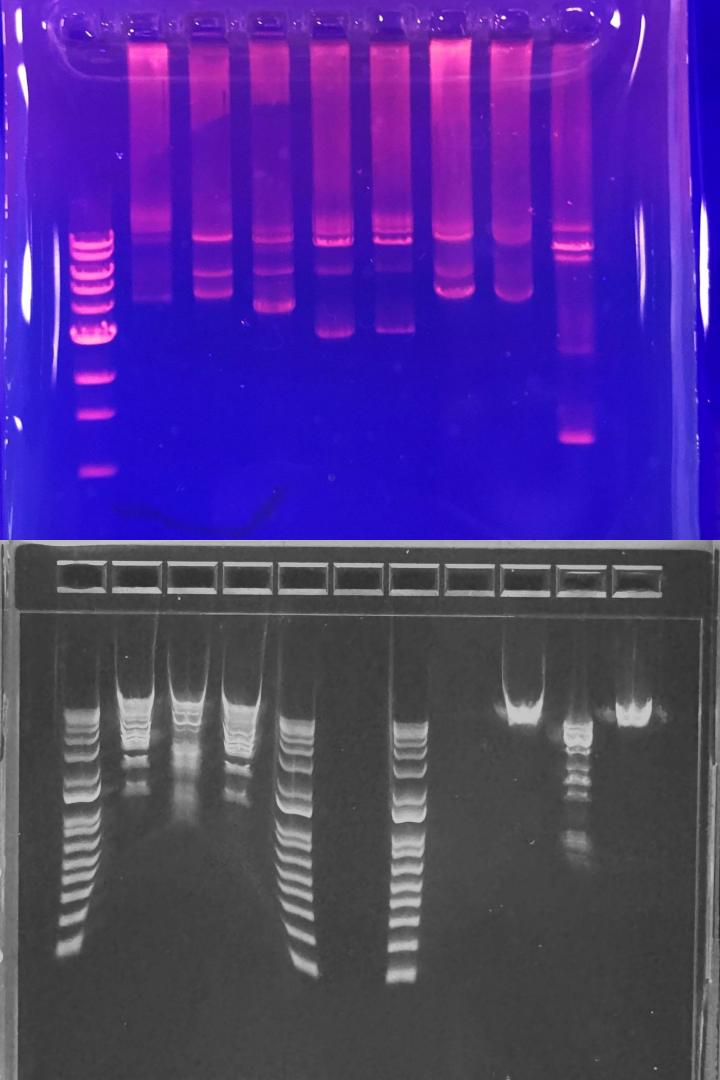
# Week 01

## Safety & Liquid Handling Fundamentals

Introduction to laboratory liquid handling tools and ethical considerations in synthetic biology

### *Lab Skills*

- Biological and chemical lab safety
- Establish student website portfolios
- Optional: Liquid handling fundamentals (e.g. pipetting)



# Week 02

## DNA Manipulation

Edit DNA using restriction enzyme digestion to generate patterns visualized on gel electrophoresis

### *Lab Skills*

- Benchling for DNA digests
- Restriction enzymes
- DNA gel electrophoresis
- Fluorescence imaging



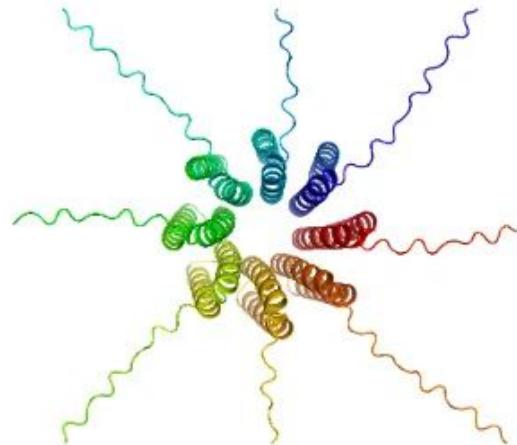
# Week 03

## Lab Automation & Liquid Handling Robots

Program the Opentrons liquid handling robots to generate pixel arrays of fluorescent bacteria

### Lab Skills

- Python and Jupyter notebooks
- Opentrons robot programming
- Bacterial culture and glycerol stocks
- Agar plate preparation and autoclaving
- Fluorescence imaging



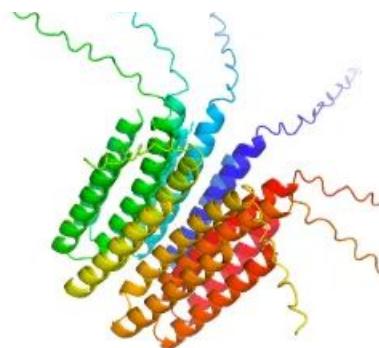
# Weeks 04-05

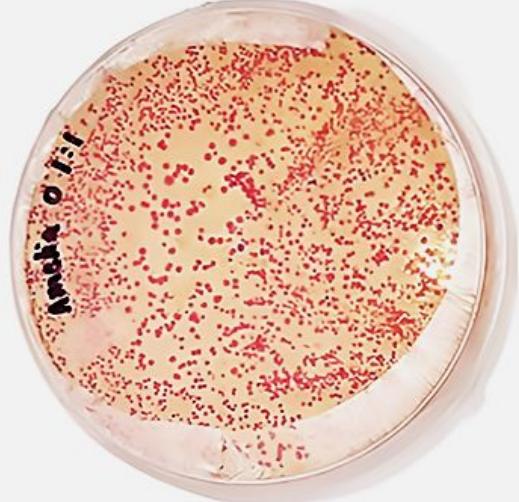
## Protein Designs Parts 1 & 2

Use AlphaFold, ESMFold, and Rosetta to study and model protein folding in silico

### *Lab Skills*

- Deep learning-based structure prediction
- RMSD calculation and structural alignment
- Interface and binding-site prediction
- Protein design and optimization
- Multiple sequence alignment (MSA)
- MS2 bacteriophage lysis/coat protein mutations





# Week 06

## The Chromophore Color Cloning Quest: DNA Design & Assembly

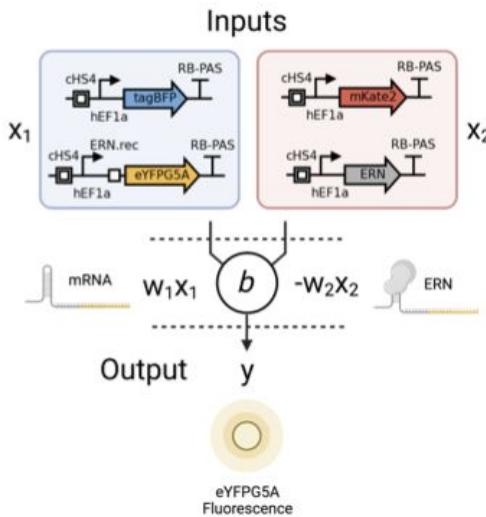
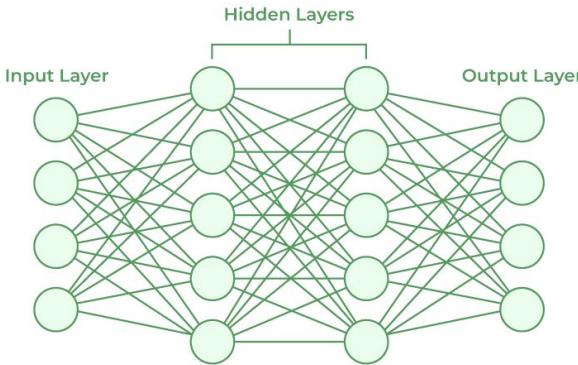
Change the color of the amilCP chromoprotein using PCR mutagenesis

### *Lab Skills*

- DNA primer design (Benchling)
- Polymerase Chain Reaction (PCR)
- DNA assembly (Gibson Assembly)
- DNA transformations
- Bacterial agar plating

# Week 07

## Neuromorphic Circuits Lab



### Lab Skills

- Neuromorphic circuit design and simulation
- DNA assembly (Gibson Assembly)
- Liquid-handling robot mediated assembly
- Transfection and flow cytometry for circuit testing

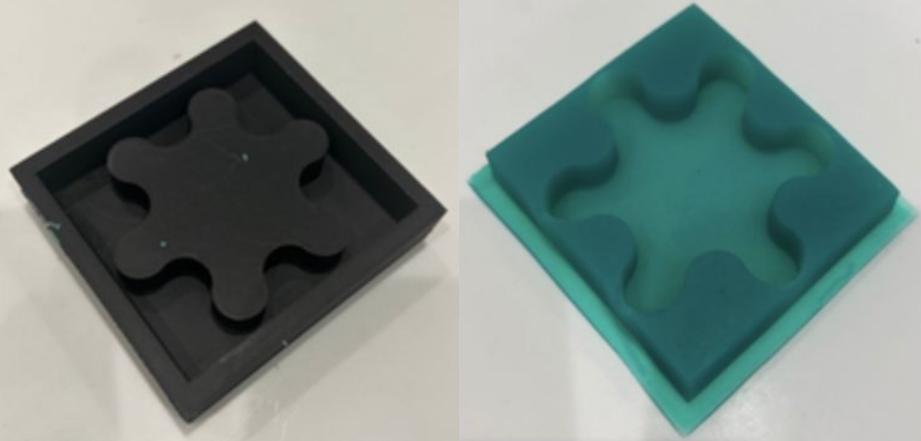
# Week 07.5

## Living Materials Lab

Design and fabricate a 3D living material artifact made from mycelium

### *Lab Skills*

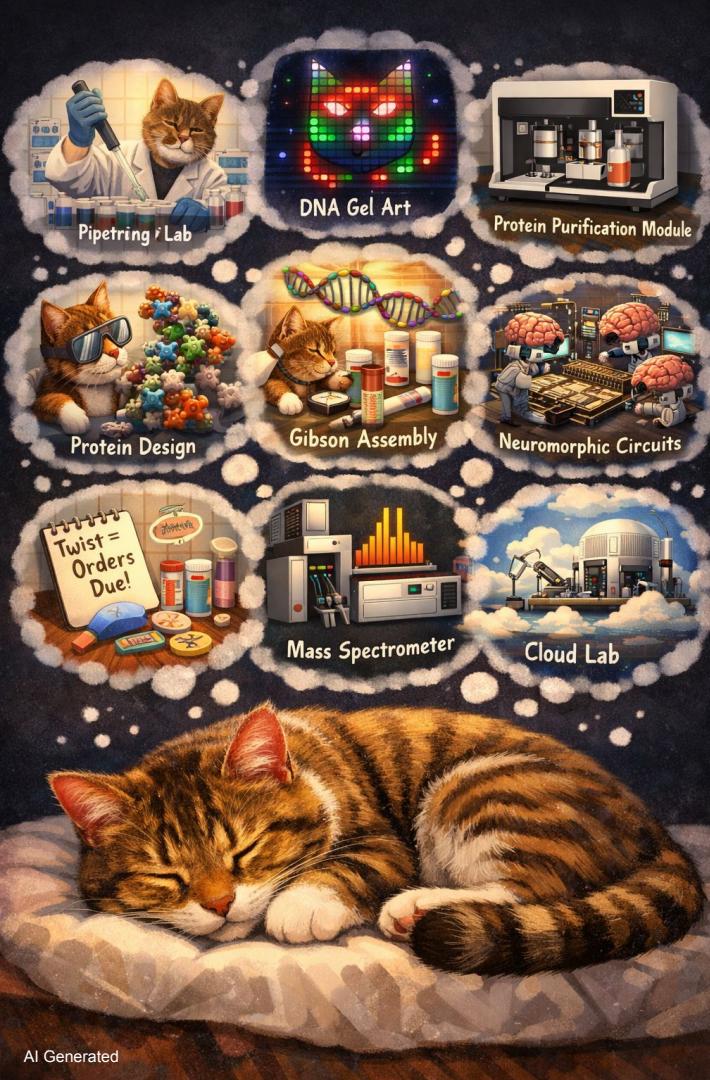
- Computer-aided design
- 3D printing
- Mycelium based living materials

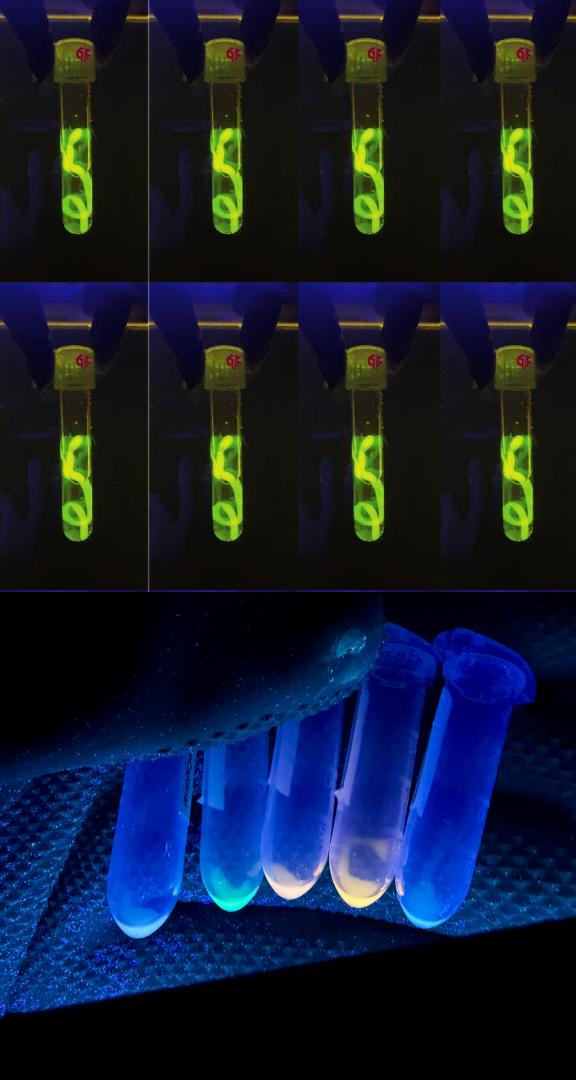


# Week 08

## MIT Spring Break

You'll rest and dream about your final projects





# Week 09

## Cell-Free Protein Expression

Cell-free protein synthesis to produce fluorescent proteins, removing the need for living organisms

### *Lab Skills*

- Cell-free protein synthesis reactions
- Reagent optimization
- Standard curve preparation
- Quantitative fluorescence measurements
- Data analysis
- Protein purification



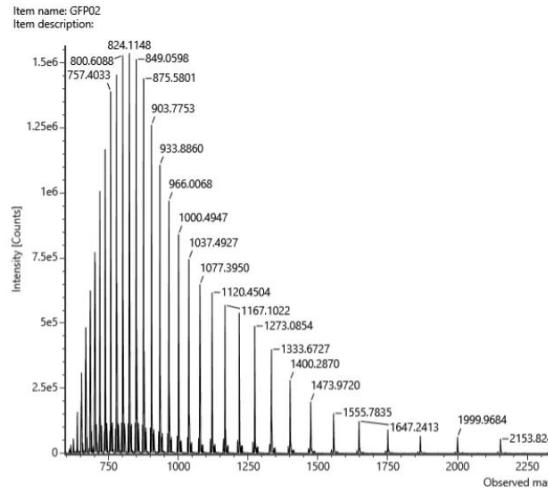
# Week 10

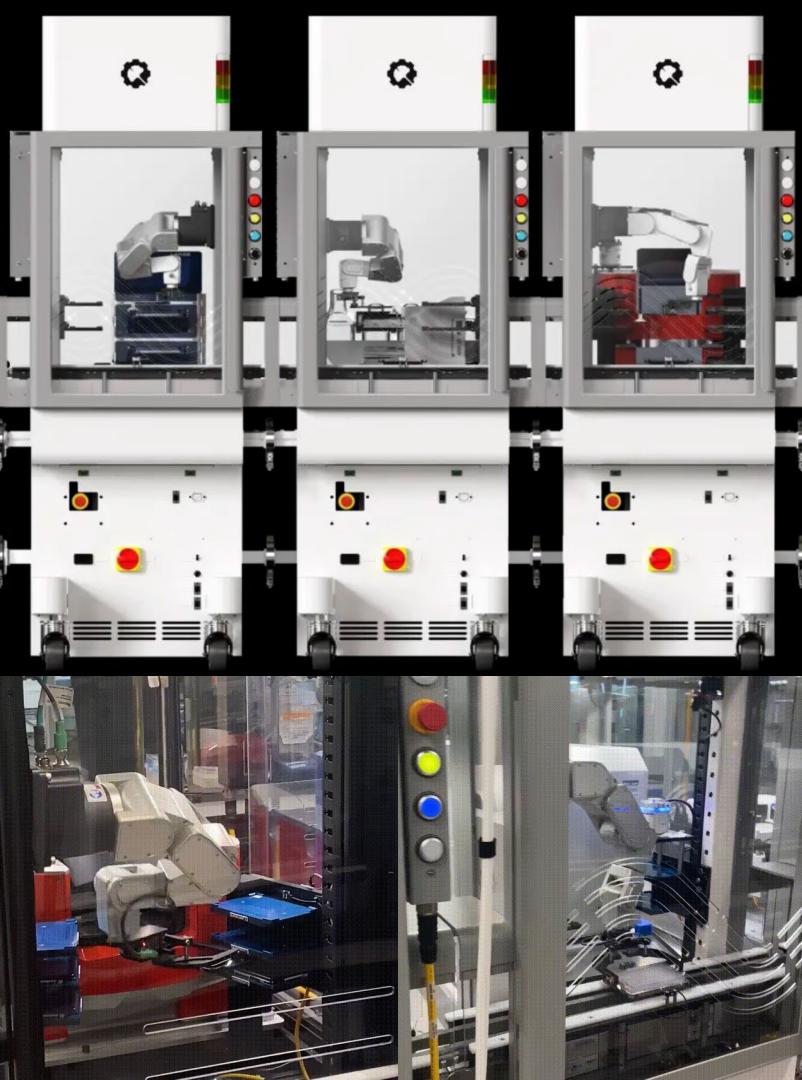
## Advanced Imaging & Measurement Technology

Work with purified and unpurified cell-free proteins under denaturing conditions to measure molecular weights at Waters (Immerse Cambridge)

### Lab Skills

- LC-MS charge states and m/z to calculate molecular weight
- Trypsin digestion to cleave proteins into predictable peptides
- Peptide mapping & mass accuracy





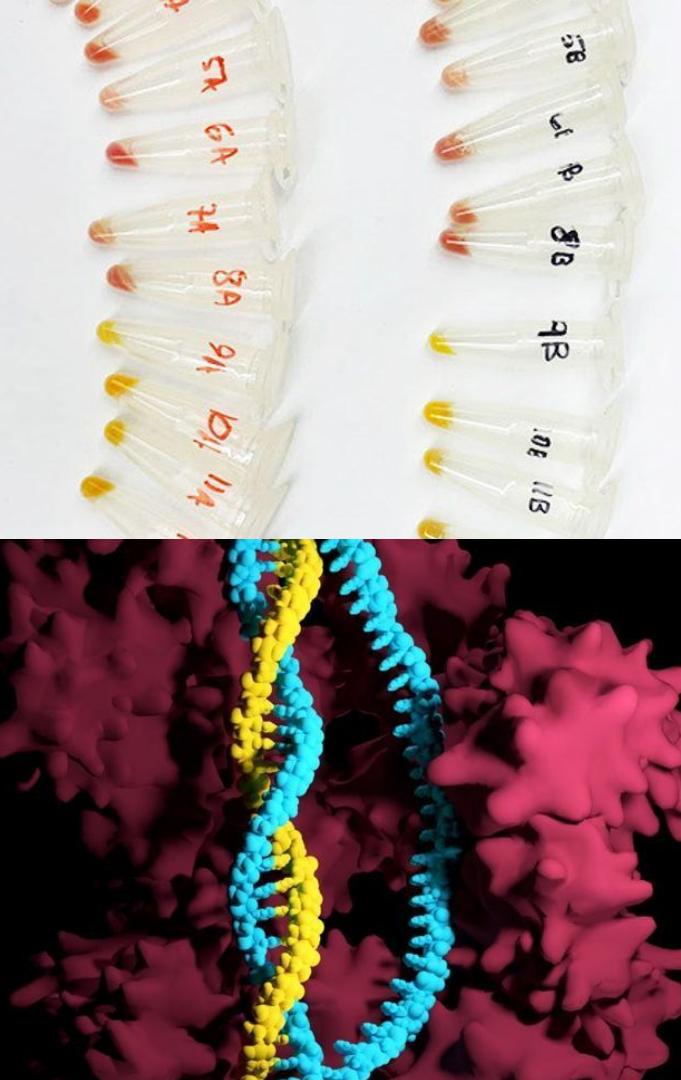
# Week 11

## Cloud Laboratories

Program the Ginkgo Bioworks cloud laboratory to conduct autonomous experiments

### *Lab Skills*

- Remote protocol creation and deployment
- Experiment standardization & reproducibility
- High-throughput cell-free assay development



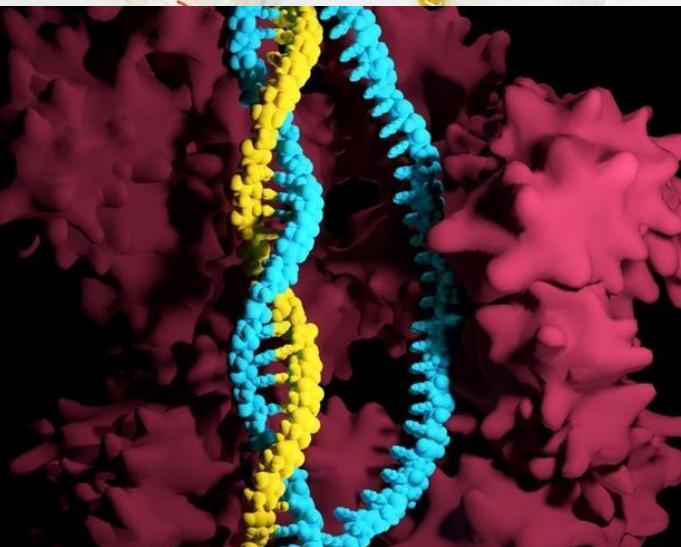
# Week 12

## CRISPRa Metabolic Engineering

Use CRISPRa for combinatorial pathway programming of biopterin bioproduction

### *Lab Skills*

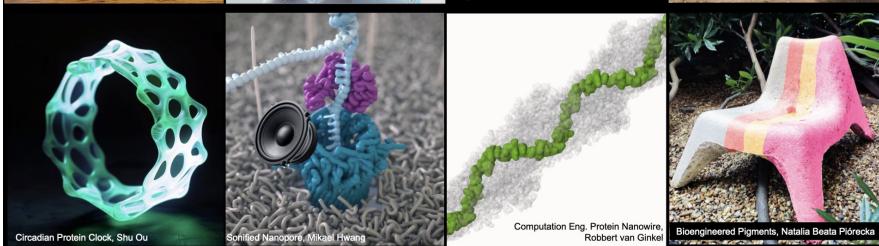
- CRISPR fundamentals
- Dual plasmid transformation
- Analyzing growth patterns
- Quantitative spectrophotometric measurements



# Week 13

## Final Projects

Conduct lab work for your final project  
utilizing the skills and techniques developed  
throughout the semester



# Homework 01 Requirements

# Homework I Requirements

Homework I (All MIT/Harvard students and Committed Listeners) Due: **Due: Feb 10** (Before lecture)

- Set up your [student webpage](#)
- Propose a biological engineering application, then design and compare multiple [governance/policy strategies](#) to ensure it is developed and used ethically and safely
- Before next week lecture on “DNA Read, Write, and Edit”:
  - Answer pre-lecture questions from Dr. Jacobson, Dr. LeProust, Dr. George Church
  - Review the associated papers referenced in the [week 2 slides](#)

Lab Safety Training (MIT/Harvard Students)

- Complete Lab Specific Training in Person. **Due: Feb 3** (we will complete as a group after this recitation)
- Complete Safety Training in Atlas **Due: Feb 4** (required for enrollment)
  - Navigate to [atlas.mit.edu](#) and on the right-hand side, click “Learning Center”
  - Head to the Course Catalog and find the following two courses:
    - General Biosafety for Researchers (EHS00260w)
    - Managing Hazardous Waste (EHS00501w)



**atlas.mit.edu**

1. Go to Learning Center
2. Click on the Course Catalog Tab:
  - General Biosafety for Researchers (EHS00260W)
  - Managing Hazardous Waste (EHS00501W)



Important information: Please see [MIT's Response to the Challenge of Covid-19](#) to understand how MIT is closely monitoring and responding to the Covid-19 situation.



## MENU

edit ↗

[Time and Vacation Entry](#)[Buying](#)[My Reimbursements](#)[Service Requests](#)**Learning Center**[Event Planning](#)[FULL MENU](#)

## My Training Needs

## My Courses

## Course Catalog

## My Profile



## General Biosafety for Researchers

TRAINING NEED

[My Training Needs](#) / [My Training](#)

You successfully completed this training need.

## REFERENCE CODE

EHS00260

## PRIORITY

Required

## STATUS

COMPLETED

## EXPIRATION DATE

No Expiration

## REASON FOR REQUIREMENT

Activity: Use biological materials (including microorganisms, plants, animals, recombinant or synthetic DNA/RNA, etc.), requiring BL1 or BL2 containment.

## ASSIGNMENT FULFILLMENT OPTIONS (You must complete ONE of the following)

**General Biosafety for Researchers (Web-Based)**

WEB-BASED

Course Code: EHS00260w

**General Biosafety for Researchers CR (Classroom)**

CLASSROOM

Course Code: EHS00260c

## NO CLASSES CURRENTLY SCHEDULED

Click through for more information and/or to Prebook for the next session

# Principles, Ethics, Practices HW

**YOU WERE SO PREOCCUPIED WITH WHETHER OR NOT YOU COULD**

**YOU DIDN'T STOP TO THINK IF YOU SHOULD**



# 1

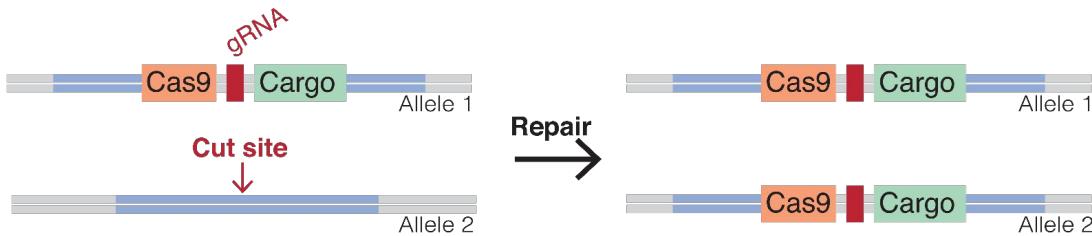
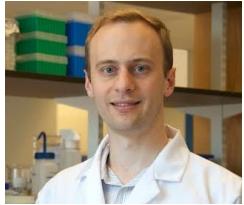
## **Describe a biological engineering application or tool you want to develop and why.**

An idea for your HTGAA final project

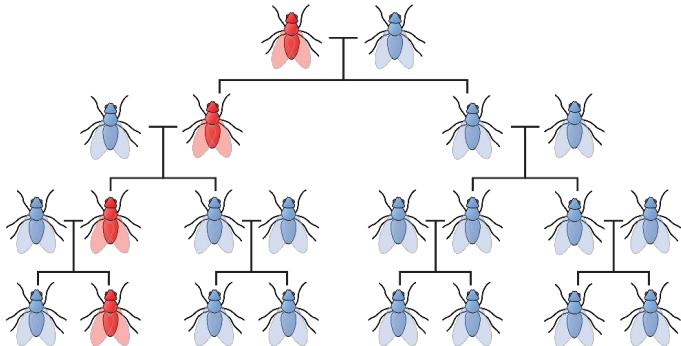
Your current research

A topic you are curious about

# Gene drives

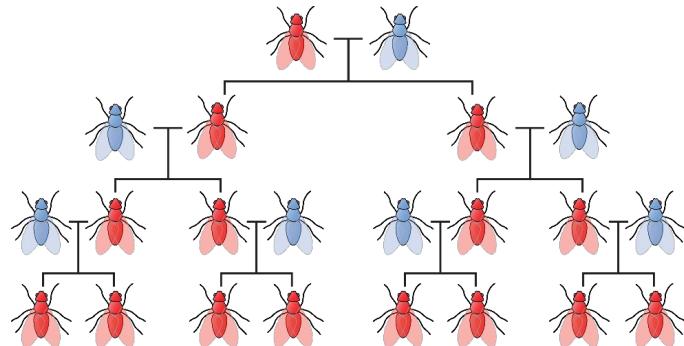


## Normal inheritance



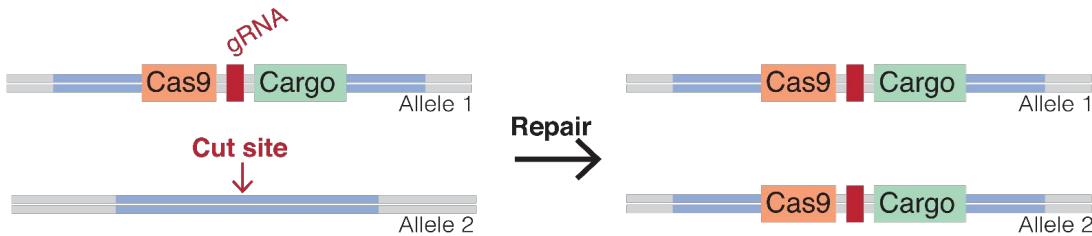
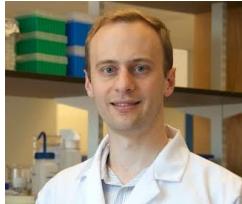
Altered gene does not spread

## Gene drive inheritance



Altered gene is always inherited

# Gene drives

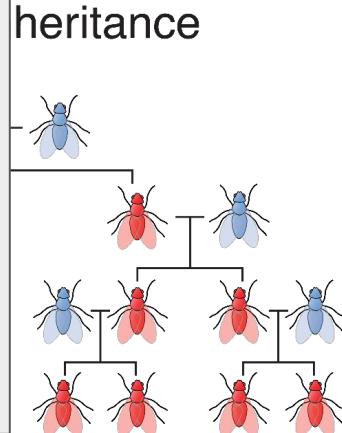


**Science**  
JOURNALS AAAS

## Regulating gene drives

Regulatory gaps must be filled before gene drives could be used in the wild

KENNETH A. OYE, KEVIN ESVELT, EVAN APPLETON, FLAMINIA CATTERUCCIA, GEORGE CHURCH, TODD KUIKEN, SHLOMIYA BAR-YAM LIGHTFOOT, JULIE McNAMARA, ANDREA SMIDLER, AND JAMES P. COLLINS [Authors Info & Affiliations](#)



Altered gene does not spread

Altered gene is always inherited

# “Virus hunting”



# 2

**Describe one or more governance policy goals related to ensuring this application contributes to an ethical future & prevents harm.**

Consider policy goals:

- ensure safety & security
- promote constructive uses
- promote equity & autonomy

# 3

## **Describe at least three different potential governance actions by considering the purpose, design, assumptions, and risks of failures & “success”**

Draw upon your existing knowledge and a little additional digging, and feel free to use analogies to other domains such as 3D printing, drones, financial system, etc.

**Purpose:** What is done now and what changes are you proposing?

**Design:** What is needed to make it “work”? Consider the actor(s) involved - who must opt in, fund, approve, or implement?

**Assumptions:** What could you have wrong? Incorrect assumptions? Uncertainties?

**Risks of Failures & Success:** How might this fail, including any unintended consequences of the “success” of your proposed actions?

# 4

**Score each of your governance actions against  
your rubric of policy goals.**

You can score from 1-3, 1 as best, or n/a

## Governance Actions

Your context:  Does the option	Option 1:	Option 2:	Option 3:
Enhance Biosecurity		<b>Scores</b>	
<ul style="list-style-type: none"> <li>● By preventing incidents</li> <li>● By helping respond</li> </ul>			
Foster Lab Safety <ul style="list-style-type: none"> <li>● By preventing incident</li> <li>● By helping respond</li> </ul>			
Protect the environment <ul style="list-style-type: none"> <li>● By preventing incidents</li> <li>● By helping respond</li> </ul>			
Other considerations <ul style="list-style-type: none"> <li>● Minimizing costs and burdens to stakeholders</li> <li>● Feasibility?</li> <li>● Not impede research</li> <li>● Promote constructive applications</li> </ul>			

Policy Goals

# 5

**Based on scores, describe which governance option or combination of options, you would prioritize, and why.**

Outline any trade-offs you considered as well as assumptions and uncertainties.

Think about your audience - very local (MIT, Cambridge Mayoral Office), to national (President or Head of a Federal Agency), to international (United Nations Office of the Secretary-General)

Lenni's week 1 homework

## STAKEHOLDERS & ACTIONS

R	I	L	C
RESEARCHERS/ ACADEMICS	INDUSTRIAL RECYCLING COMPANIES	LAW ENFORCEMENT / REGULATORS	CITIZEN / USERS
<p>R1. Establish best practices to handle bacteria and enzyme</p> <p>R2. Compile database and share climate specific strategies</p> <p>R3. Develop strategies to evaluate and track bacteria in the wilderness and built environment</p>	<p>I1. Undergo annual review with researchers and regulators</p> <p>I2. Compile manual for enzyme use</p> <p>I3. Develop framework for commercial distribution of enzymes for approval</p>	<p>L1. Ensure that researchers and industrial parties register their bacterial and enzymic quantities</p> <p>L2. Ensure educational on best practices are held for the general public</p> <p>L3. Carry out routine environmental inspection of the deployed areas</p>	<p>C1. Be educated about current best practices</p> <p>C2. Contribute to monitoring and responding efforts for bacteria deployed in the environment</p> <p>C3. Register DIY home enzymic 'composting' into monitored</p>

 <p>ENVIRONMENTAL PROTECTION</p> <p><i>If the waste-ingesting bacteria is deployed in the wilderness, how can we ensure that it will not negatively affect other living organisms?</i></p>	<p>How do we evaluate climate-specific waste-eating bacteria strains in relation to its engineered quality?</p>
 <p>ENHANCE BIOSECURITY</p> <p><i>Who will be regulating the research and development globally in research institutes, industrial and domestic use?</i></p> <p><i>How do we promote an equitable and safe use in specific climate and environments, preventing misuse?</i></p>	<p><b>AREAS OF CONCERN: GENETICALLY ENGINEERED BACTERIA FOR ENZYMES TO BREAKDOWN WASTES</b></p> <p><i>How do we prevent incidents harmful to researchers?</i></p> <p><i>How do we promote and not impede research developments?</i></p>
 <p>BUILT ENVIRONMENT PROTECTION</p> <p><i>How do we control waste-eating bacterial enzymes to not degrade existing and in-use buildings (vs feeding on urban derelicts)?</i></p>	

Amelia Gan - local deployment of bacteria that produce plastic-ingesting enzyme (PETase)



#### BIOSECURITY CONCERN

- > If engineered bacteria can be used to solidify the ground can it be used to cause instability?
- > Can genetically engineered bacterial soils negatively affect soil fertility?



#### EQUITY CONCERN

- > Who has the ability to use the engineered bacterial soils?
- > Who regulates the use of engineered bacterial soils at different scales/ in different countries?



#### ENVIRONMENTAL CONCERN

- > How is microbial activity in soils affected in the immediate and surrounding contexts?
- > How do we evaluate the cons/pros when you consider non-human needs? Will non-human organisms benefit or be harmed?

## GENETICALLY ENGINEERED BACTERIA TO ALTER SOILS

	RESEARCHERS	MANUFACTURERS	INDUSTRY	ORGANIZATIONS
<b>ENHANCE BIOSECURITY</b>				
Monitoring	○	○	○	○
Response		○	●	
<b>ENSURE EQUITABLE USE</b>				
Monitoring	○	●	○	○
Response	○	●	○	○
<b>PROTECT ENVIRONMENT</b>				
Monitoring	○	○	●	○
Response	○	○	○	●

#### KEY

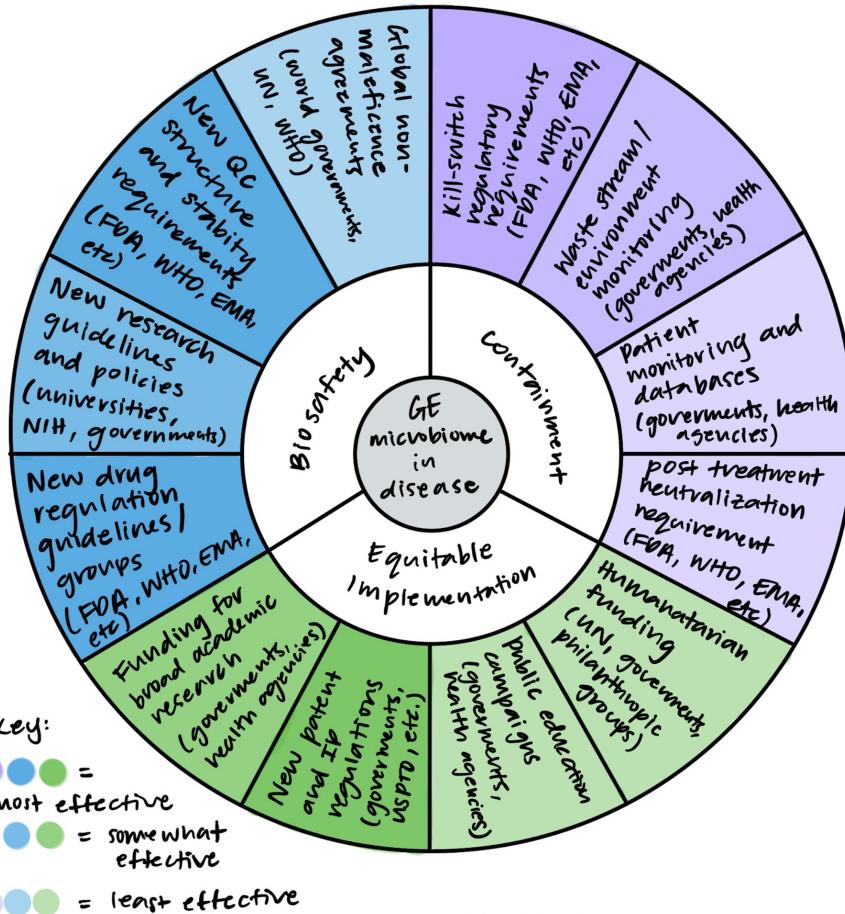
● Most Effective ○ Moderately Effective ○ Minimally Effective

Researchers include academic (university) researchers along with researchers at community labs.

Manufacturers include producers of the genetically engineered bacteria. They would be the source for the bacteria.

Industry includes registered users and can vary in scale from design/construction companies, to indiv. farmers or ceramicist.

Organizations include committees and advisory boards that establish and enforce legislation at various scales.



Jenna Houle - engineered bacteria for the detection and treatment of cholera

## Weekly Assignments

- 1) Outline any ethical concerns that arose, especially any that were new to you.
- 2) Propose any governance actions you think might be appropriate to address those issues.
- 3) Include the governance actions on your class page for that week.

## **Final Project Assignments**

As part of your final project, design one or more strategies to ensure that your project, and what it enables, contributes to growing an ethical biological future.

# Small Group Discussion

**Introduce yourself to the person on your left & right!**

*Share a synthetic biology project idea & related governance actions*

***JOIN A ZOOM BREAKOUT ROOM PLEASE!***

# Student Website Live Demo



# HTGAA Pages: How To Edit (Almost) Anything



webpages/content at main · 20 · + edit.hgaa.org/2026a-recitation-demo/webpages/src/branch/main/content

main Add file webpages / content History

Search code... Exact

Spring 2026 Recitation De... 1912e7828e Initial commit two days ago

..

homework Initial commit two days ago

labs Initial commit two days ago

projects Initial commit two days ago

\_index.md Initial commit two days ago

README.md Initial commit two days ago

README.md

## Your HTGAA site

This is the top level of your HTGAA site - this directory contains your homepage and directories inside it contain all of the other pages on your site. (We recommend you bookmark this page so you can come directly here to edit.)

Click [\\_index.md](#) here (or above) to edit the homepage of your HTGAA site, or click [homework](#), [labs](#), or [projects](#) to work on those. Follow the guidance in the READMEs in those locations, and refer to the [HTGAA Markdown Editing Guide](#) for tips on markdown, images, tools, and more.

Find your published pages at <https://pages.hgaa.org/2026a-recitation-demo/> once you start adding content here.

## Homework assignments

### To start a new week's homework assignment:

1. Right above the box which says `Search code...` click on `Add file` → `New file` and give it a name with the pattern `<week-name>/_index.md`, such as `week-01-hw-principles-and-practices/_index.md`. (For convenience, see some suggested `<week-name>`s below.)
2. Inside that new file, make the first few lines of your file contain something like this: (including the `---`)

```
---
```

```
title: 'Week 1 HW: Principles and Practices'
```

```
weight: 10
```

```
---
```



The weight values are used to order the weeks; a good default is  $10 * \text{week number}$  (so 10, 20, 30, ...).

3. Then write your homework assignment using **markdown!** (And for reference see the [HTGAA Editing Guide](#).)
4. Don't forget to scroll down and click the `Commit changes` button to save after making some additions or edits. After saving while you're still on that `_index.md`'s page you can click the pencil icon (next to the trash can on the far right) to go back into the editor.

To make it easier the first time around, Week 1 has already been started for you.

Remember, once you save you can view your published pages at <https://pages.htgaa.org/2026a-recitation-demo/>

### Extra customization:

1. To add **images** to your pages,
  - i. Upload the image: after you've saved your `_index.md`, click on the red `<week-name>` directory link a few lines down from the top.
  - ii. There, you can click on `Add file` → `Upload file` and either upload an image file you have locally or when it asks for a filename try giving it a URL to the image.
  - iii. Back inside your document (click on `_index.md` and edit it) you can include the image by using its filename in an **image link** (e.g. ``)
  - iv. Remember, if you put the word `feature` anywhere in an image's filename (e.g. `featured-woolly-mammoth.webp` or `david-george-joe.jpg-feature`) that image will automatically be placed inside the "card" for that page in a `{}% children type="card" %}}` (or `childrenof`) card listing (such as a list of all your homework, if you are specifying `type="card"`).
2. Page summaries are listed in cards or page listings; long ones are automatically generated, but you can create your own short description for those listings by adding between the `---` lines at the top of your markdown document a line such as:

```
description: 'whatever you want the summary to be'
```

webpages/content/homework x Editing Guide :: HTGAA Spring 2026 x +

pages.htgaa.org/2026a/course-pages/editing-guide/ Incognito Finish update

# HTGAA Spring 2026

Search...

Course Logistics

Weeks

Final Projects

Students

TAs

Global Nodes

Editing Guide

HTGAA Spring 2026 > Editing Guide

Markdown Tools New pages Images

# EDITING GUIDE

emojis icons cheat sheet full guide shortcodes mermaid

## Markdown

In your `_index.md` files you mainly write "Markdown", which is an easy way to add simple formatting to a document just by typing. If you just write text it will get displayed as text (without any special formatting). By adding simple punctuation you can mark portions of text as **bold**, *italic*, **highlighted**, **quoted**, lists, headings of a half dozen levels, inlined images, [web links](#), and more.

- You can copy-paste into your markdown any emoji (😊) or type its name (:grin:) from the [official emoji list](#)
- Icons can be included via `{}% icon <icon-name> %{}%` using `<icon-name>`s which you can [search](#) for in the [official icon directory](#) (be sure the "Free" box on the right, below the search bar, is selected). If the "Web Awesome" code has a `family` name listed (e.g. if you click the "Web Awesome" tab for [this icon](#) you see the family name is `brands`), enter that after the icon name, e.g. `{}% icon google brands %{}%`.
- See a [basic markdown cheat sheet](#)
- Or refer to a [more complete markdown guide](#) with many more advanced features (navigate using the Table of Contents by clicking the `table` icon at top on the left side)

## Tools

The markdown you write is processed using the [Hugo Relearn Theme](#).<sup>1</sup> For advanced layout of your pages you can use their many "[shortcodes](#)" – a few to call out are:

- For layout & formatting, use `cards` and `notices`; also `buttons` can provide colorful links.
- For organization, you can group related pages into directories and use `children` to automatically display titles & optional descriptions of all pages below in various formats. We also provide a custom version of it, `{}% childrenof path="/projects" ... %{}%` which takes all the same parameters as `children` but only lists

# **Student Website Editing**

# Start editing your HTGAA Website!

Good places to begin include:

- Your homepage: `content/_index.md`
- Your Week 1 Homework submission:

`content/homework/week-01-hw-principles-and-practices/_index.md`

# editing mechanics: directories, files, & headers

- Homework should go under content/homework, labs under content/labs
  - Inside those directories a README.md is displayed giving more detailed instructions
- Make a directory for each webpage (e.g. for one homework, for one lab, etc)
  - Put the content (markdown) in the file `_index.md` in that directory
  - Add images for the page into that directory
  - Within it you can have subdirectories with their own `_index.md` and image files if you like
- At the top of your `_index.md` files add a header which looks like this: (with the "---")

```
---
```

```
title: 'Week 1 HW: Principles and Practices'
```

```
weight: 10
```

```
---
```

- Scroll down and click the "Commit changes" button to save your edits

- Tips:

- Get content in first – just type text! – then add markdown for formatting later.
- Add/change a bit at a time; save ("Commit") often and view the results right away.
- Browse & copy markdown/layout tricks freely from anyone – BUT NOT CONTENT!

# help!

- Not seeing the change you expected? Wait 5 secs, reload, check Build Failure Log.
- Build is all-or-nothing – a breaking mistake on one page will prevent all your pages from being built/redeployed!
- Unexpected line break? Check for  $\geq 2$  trailing spaces on end of previous line.
- Icon not displaying? Be sure it's free, not pro. Does it need a second "family" name?
- Mysterious  or ? You have an image link `![]()` with an incorrect filename or path.
- On a markdown page in [edit.htgaa.org](https://edit.htgaa.org) is the pencil icon not clickable to edit / does it say "You must fork this repository..."? You're not logged in.
- Need some markdown tips? See the [HTGAA Editing Guide](#), which also links to a [Markdown Cheat Sheet](#) and an [extensive Markdown Guide](#).
  - This is linked from every one of your generated pages: "Resources" (lower left) -> "Editing Guide"
- **ASK FOR HELP!** in the [Editing/Publishing category on forum.htgaa.org](#).  
Don't spend days on a problem someone can help solve in minutes!

# Great Student Websites from previous years

Laura Maria Gonzalez (2021)

Rosalie Lin (2022)

Kevin Huang (2023)

Jocelyn Keyser (2023)

Alison Cabrera (2025)

Tanisha Shaw (2025)

# Discourse Introduction

# Welcome back!

Search



categories ▶ tags ▶

Latest

New (5)

Unread (2)

Hot

Categories

New Topic

Category

Topics

Latest

## General

Create topics here that don't fit into any other existing category.

Editing/Publishing Bootcamp 1 new

11

3 new

## TA Groups

Infrastructure Design Comms 1 new Social Media

27

2 unread  
1 new



Logging into edit.htgaa.org •

General feature-request

2

29m



BioBootcamp Presentation Slides and Survey! •

Bootcamp

1

1h

<https://forum.HTGAA.org>

# Homework 01 Requirements Review

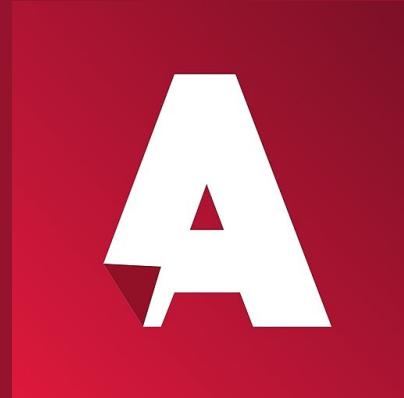
# Homework I Requirements

Homework I (All MIT/Harvard students and Committed Listeners) Due: **Due: Feb 10** (Before lecture)

- Set up your [student webpage](#)
- Propose a biological engineering application, then design and compare multiple [governance/policy strategies](#) to ensure it is developed and used ethically and safely
- Before next week lecture on “DNA Read, Write, and Edit”:
  - Answer pre-lecture questions from Dr. Jacobson, Dr. LeProust, Dr. George Church
  - Review the associated papers referenced in the [week 2 slides](#)

Lab Safety Training (MIT/Harvard Students)

- Complete Lab Specific Training in Person. **Due: Feb 3** (we will complete as a group after this recitation)
- Complete Safety Training in Atlas **Due: Feb 4** (required for enrollment)
  - Navigate to [atlas.mit.edu](#) and on the right-hand side, click “Learning Center”
  - Head to the Course Catalog and find the following two courses:
    - General Biosafety for Researchers (EHS00260w)
    - Managing Hazardous Waste (EHS00501w)



**atlas.mit.edu**

1. Go to Learning Center
2. Click on the Course Catalog Tab:
  - General Biosafety for Researchers (EHS00260W)
  - Managing Hazardous Waste (EHS00501W)

# Lab Safety Training

---

Walk over to building 68-089!

# Lab 1: Introduction to Wet Lab Liquid Handling

- **Understand Units and Conversions:** moles (mol), molarity (M), and conversions between nL,  $\mu$ L, mL, and L.
- **Gain Pipetting Proficiency:** Operate P20, P200, and P1000 pipettes accurately for volume transfers.
- **Perform Serial Dilutions:** Learn the stepwise dilution process to achieve specific solution concentrations.
- **Visualize Mixing Outcomes:** Use colors and absorbance measurements to observe concentration gradients.

$$C_1 V_1 = C_2 V_2$$

Starting concentration  
Starting volume  
Final concentration  
Final volume



# Recitation 01