

iGEM 2017 Shad Valley – Day 1: A Primer on Synthetic Biology iGEM

Presentation Outline

- Overview of Synthetic Biology
- Perspectives of Synthetic Biology
- Genetics Lab Theory Overview
- Introduction to iGEM and your Shad Valley Waterloo Experience
- Connection between Genetic Manipulation and Synthetic Biology

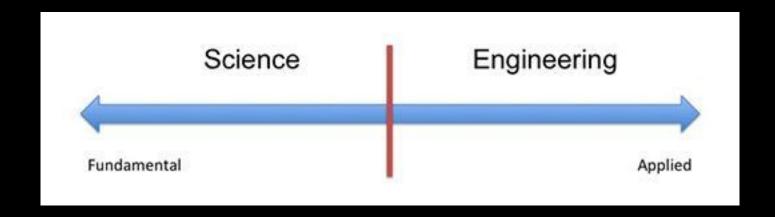


What is Synthetic Biology?





Perspectives: Science & Engineering





Genetics Theory Overview

- Central Dogma of DNA
- Plasmids and Restriction Enzymes
- BioBricks: The Plasmid Standard
- Cell Transformation & Competent Cells
- Plating Cells & The Observation of Colonies



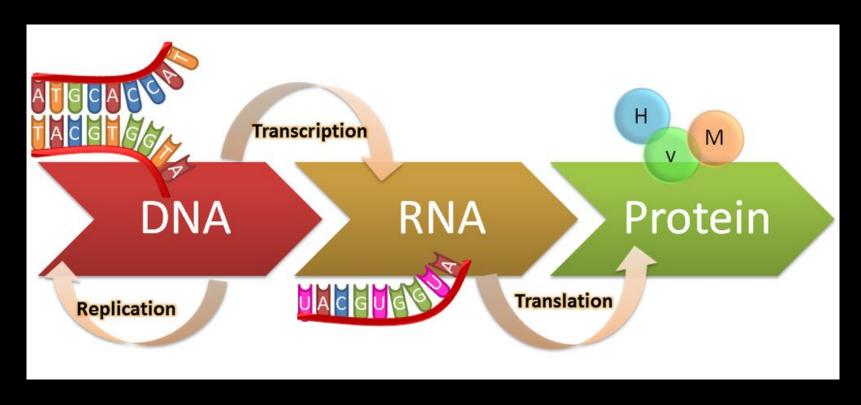
ATCG

Nitrogen Bases in DNA



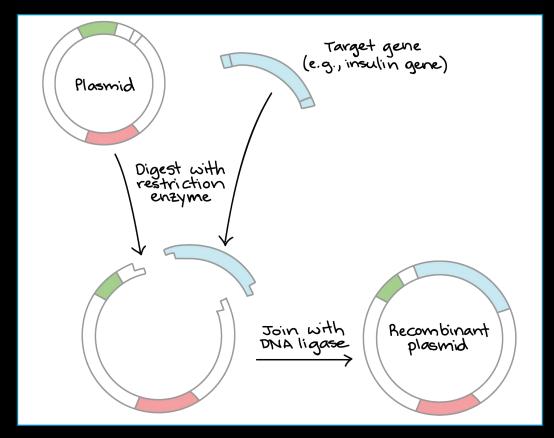


The Central Dogma of DNA



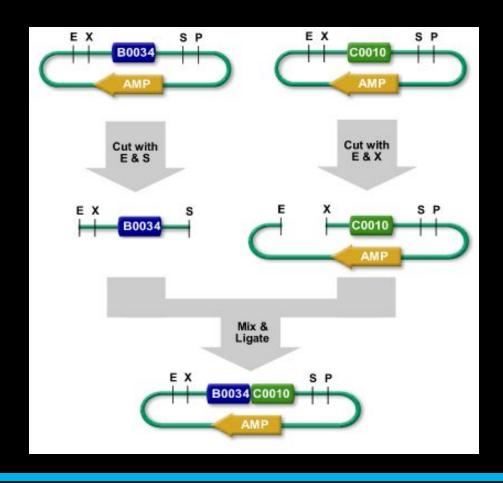


Plasmids & Restriction Enzymes



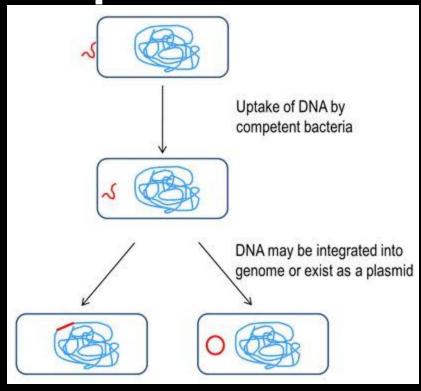


BioBricks: The Plasmid Standard



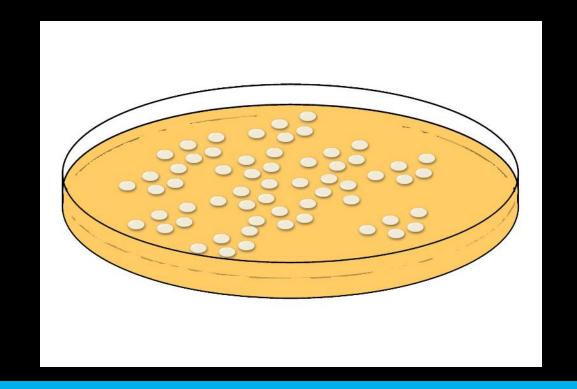


Cell Transformation & Competent Cells





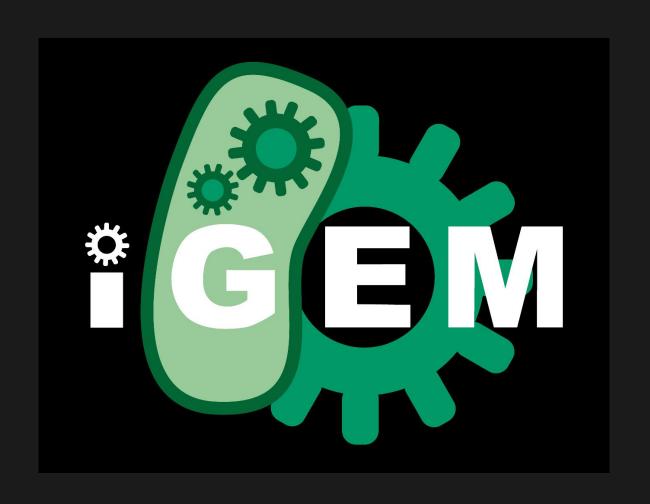
Plating Cells & The Observation of Colonies





What is iGEM?

- iGEM
- Teams from all over the world
- Compete in 9 tracks
- Our Team
- Our Project



Your Experiment @ Shad Valley Waterloo

Day 1: Pipetting, Streak Plating, Lab Tour

Day 2: Mobile Genetic Elements Talk & Mini-prep

Day 3: Electrophoresis, Gels, and Ethics in Synthetic Biology



The Link Between Genetic Manipulation & Synthetic Biology



Lab Safety

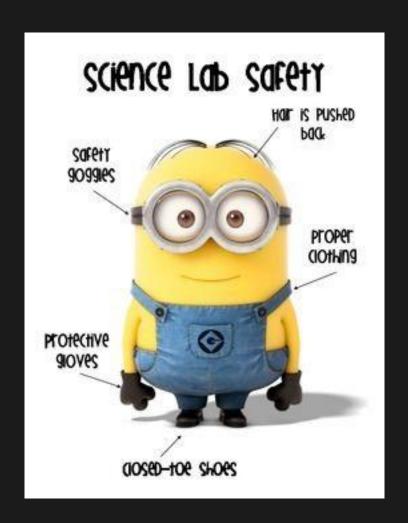
BEFORE WE CAN DO ANYTHING...



Why is lab safety Important?

Handling of Microbes

- Possible Pathogens
- Handling of Chemical Solutions
- Working with an Open Flame
- Report broken glassware immediately





Safety Equipment

Personal Protective Equipment (PPE)

- Lab Coats
- Closed-Toe Shoes
- Gloves
- Safety Glasses
- Long hair tied up
- Loose clothing tucked away





Workspace

- Always sanitize with ethanol
- Remove materials that are irrelevant to the experiment
- Maintain aseptic area with Bunsen burner
- Electronic devices are prohibited in the laboratory
- NO FOOD OR DRINKS!
- Store belongings in the provided space
- Keep test tubes upright



Aseptic Technique

What is it?

Use of a flame to clean and maintain area near flame devoid unwanted contaminants.





Clean Up!

- All waste in proper bins
- Pipette tips
- Nitrile Gloves
- Used Test tubes
- Sticks



Clear Bench and wipe down paper with acetone/ethanol



In Case of Emergency

- Eye Wash Stations
- Shower Stations
- Fire Extinguisher

If you are unsure where any of these stations are during an emergency, please ask one of our lab members for some assistance.



Thank You





iGEM 2017 Shad Valley – Day 2: Genetics Foundation and Miniprep





Here's What We're Doing Today...

- Learning about the foundation of genetics!
- 2. Review the procedure for today's mini-prep!
- 3. Perform your first mini-prep!





Protein

- -complicated 3D structures composed of one or more polypeptides
- -polypeptides are molecules made of many amino acids





Nucleic Acid

- -direct the growth and development of living organism
- -determine the characteristics and functions of a cell
- -two types of nucleic acids, **DNA** and **RNA**
- -made of nucleotides, which are composed of a phosphate group, a sugar, and a nitrogenous base



DNA

- DNA is heredity material located in the cell nucleus

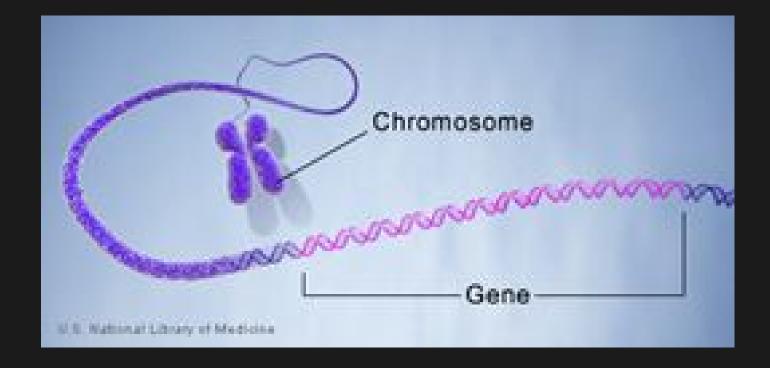




Chromosomes and Genes

Chromosomes: structures made of both nucleic acid and proteins that helps ensure DNA is accurately copied and distributed

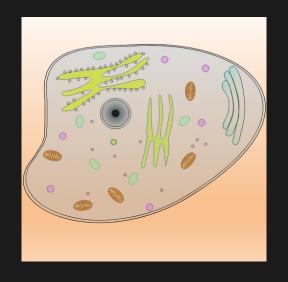
Genes: the part of the DNA that helps the cell make a protein

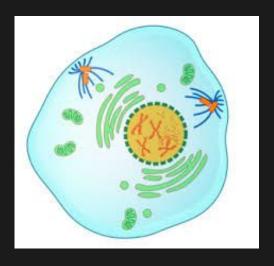




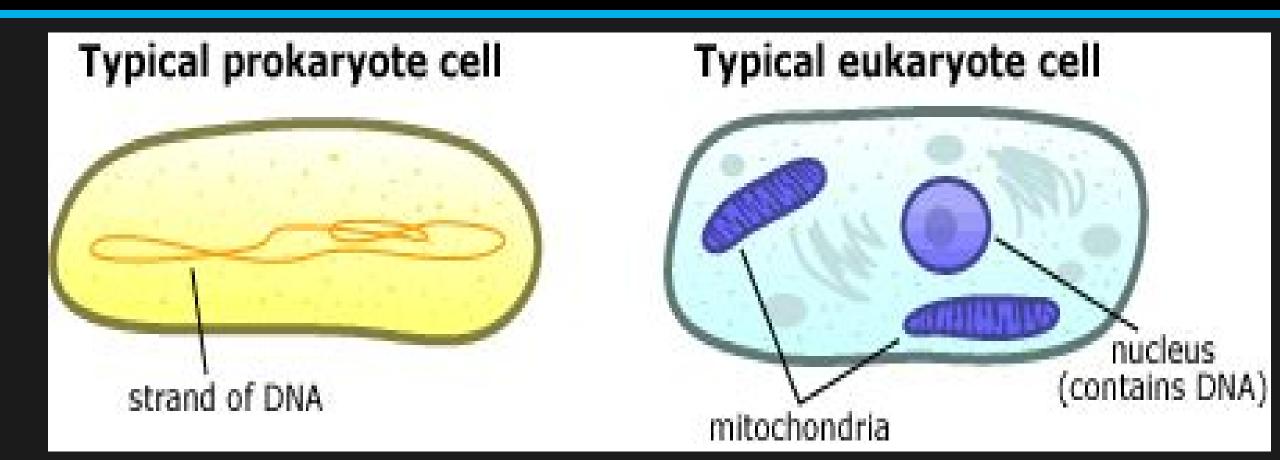
Cells

- -the basic biological unit of all living organisms
- -there are two types of cells: **prokaryotes** and **eukaryotes**





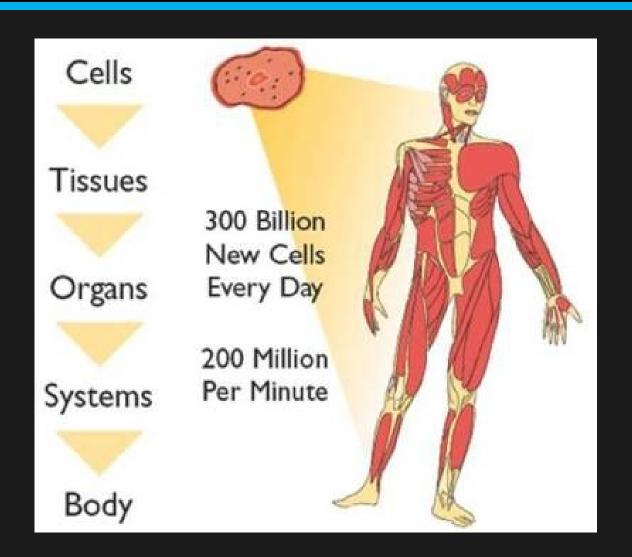






Which Are We Made Of?

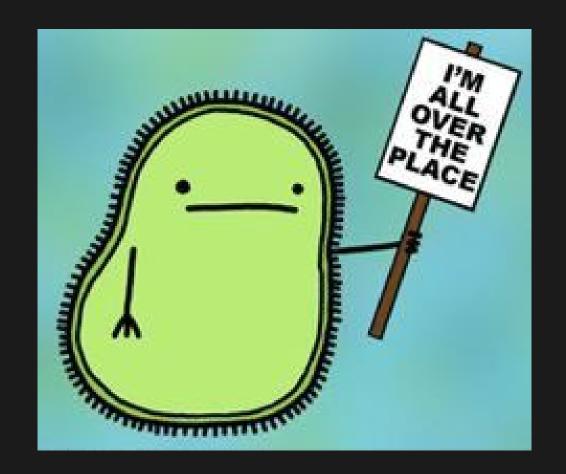
- -humans are made of around 37.2 trillion cells
- -in our bodies, there are hundreds of different functions
- -our cells are eukaryotic





Bacteria

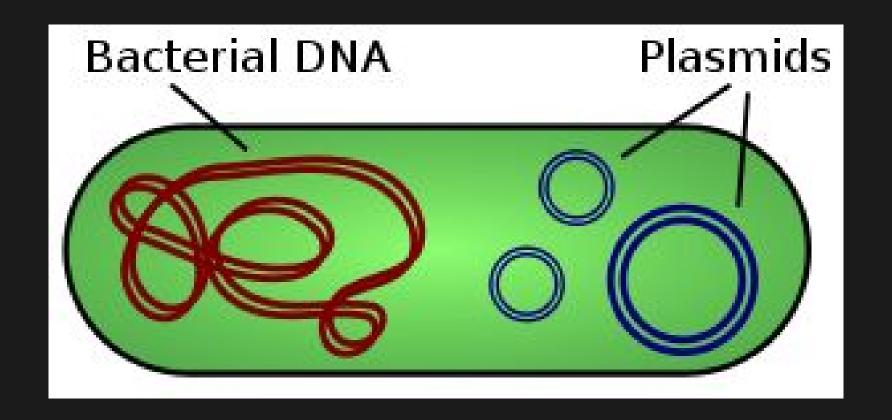
- -prokaryotic, functional organisms
- -they're almost everywhere, but not all of them are dangerous!
- -bacteria were one of the first lifeforms on earth
- -in one drop of water, there are about one million bacteria cells





Plasmid

- -circular, double stranded DNA molecules often found in bacteria
- -often used to manipulate genes











What Are We Doing?

The goal of a miniprep is to isolate plasmid DNA from a bacterium to use the plasmid for other things!

(Tomorrow we will separate the DNA by size through electrophoresis)

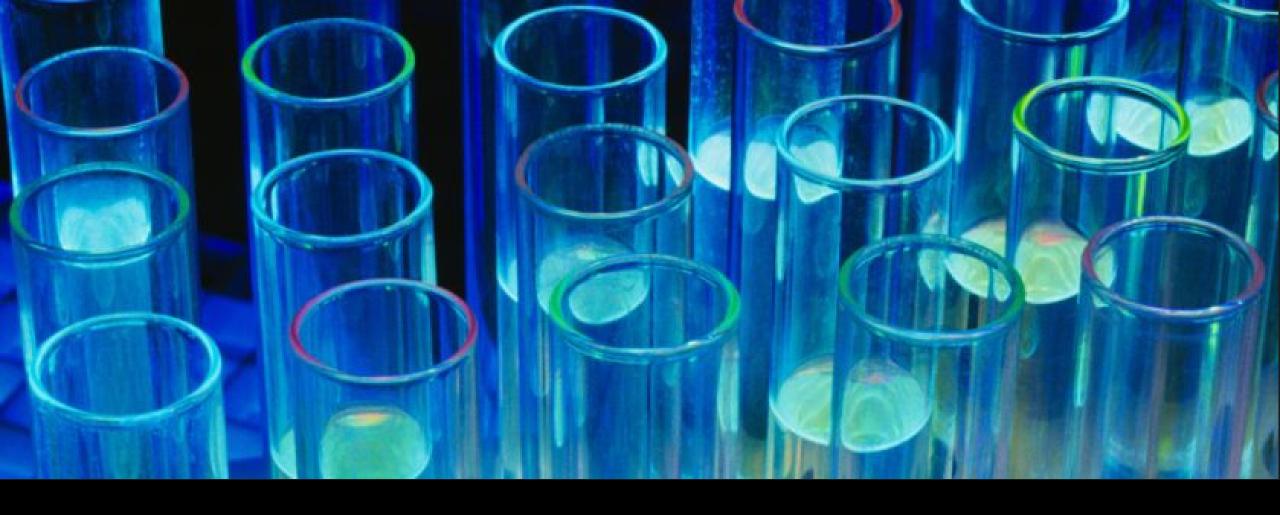


Video Time!

https://www.youtube.com/watch?v=7Xgy5_i6iOc



Good luck, and go knock your socks off!



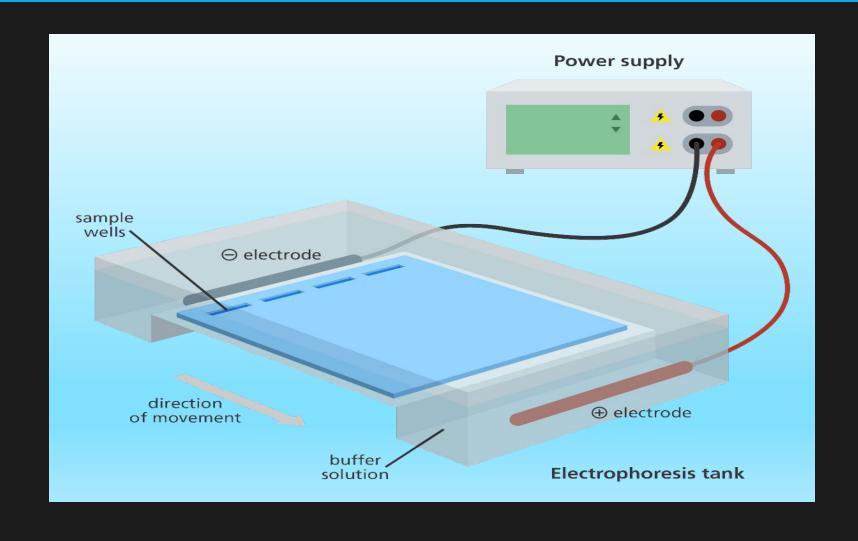
Gel Electrophoresis

Using electricity to separate DNA fragments



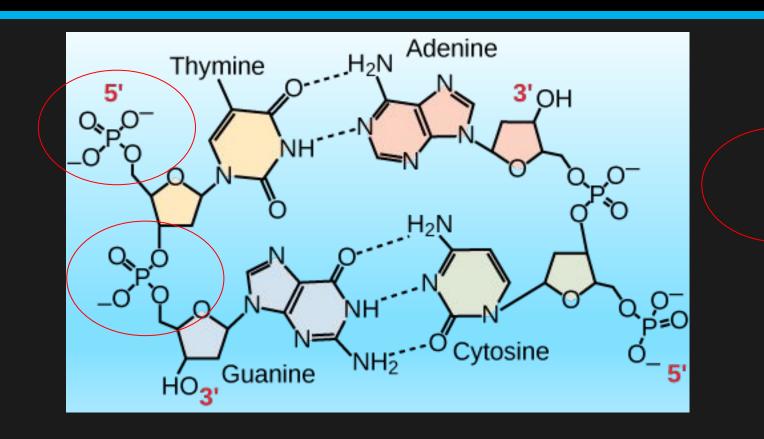


Agarose Gel Electrophoresis





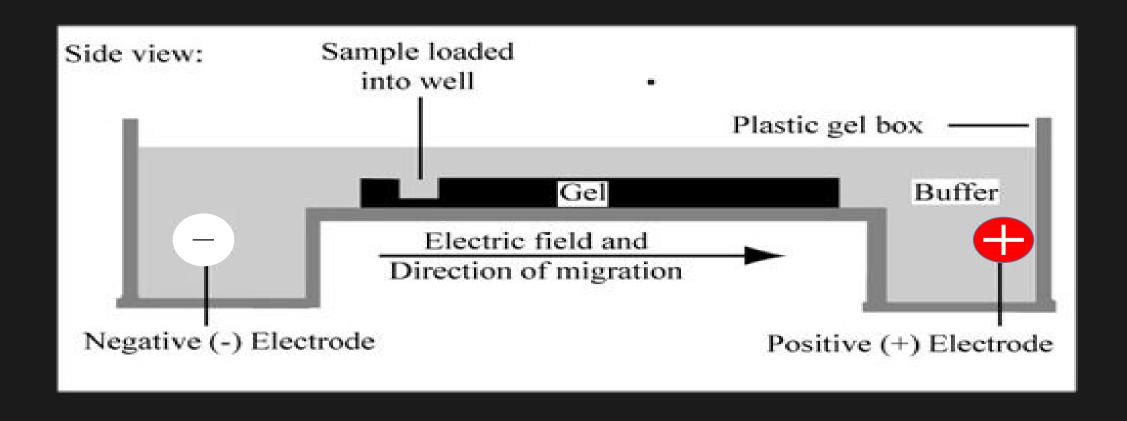
How do we sort DNA?



What's the charge on DNA?

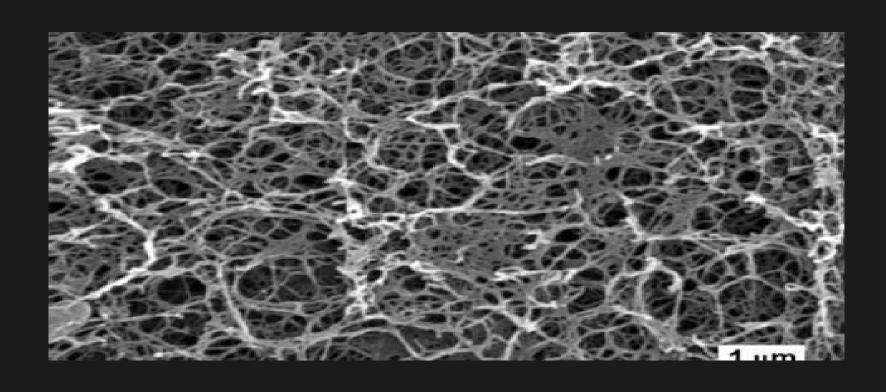


Negatively charged DNA moves to positive end





Agarose gel





Imagine Pulling a Toboggan Across a Field





What Will Travel Faster





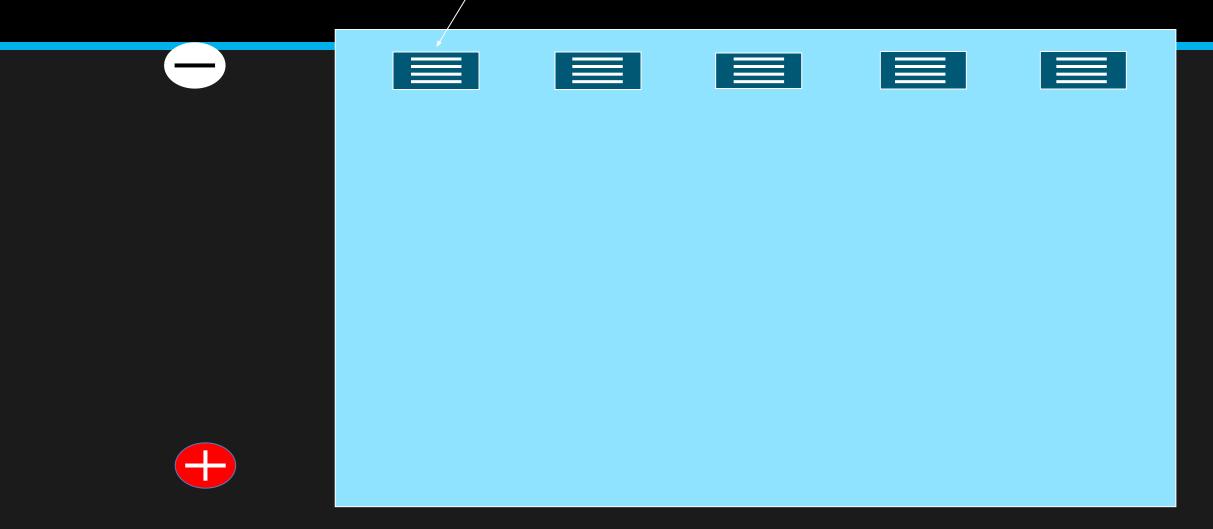




FRICTION!



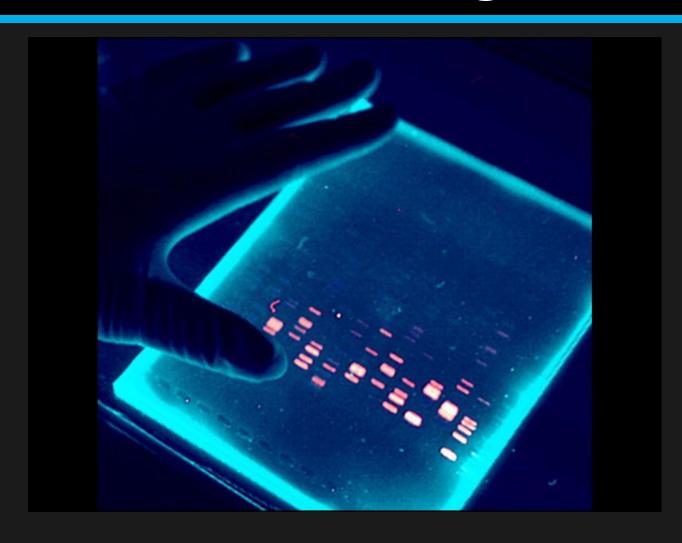




GEL ELECTROPHORESIS

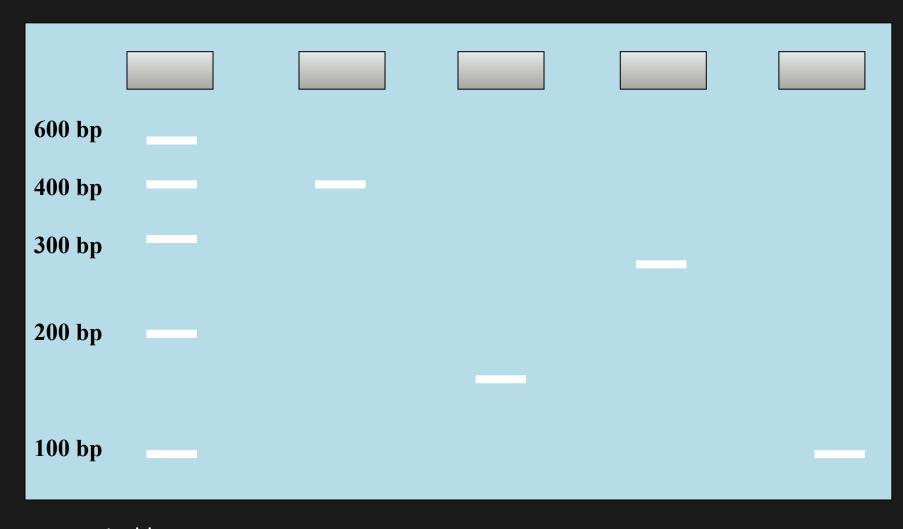


Visualizing a Gel





Determining the Length of a DNA Fragment



Ladder



BIOTECH BOOT CAMP

Melissa Prickaert

Ingalls Lab on Slack

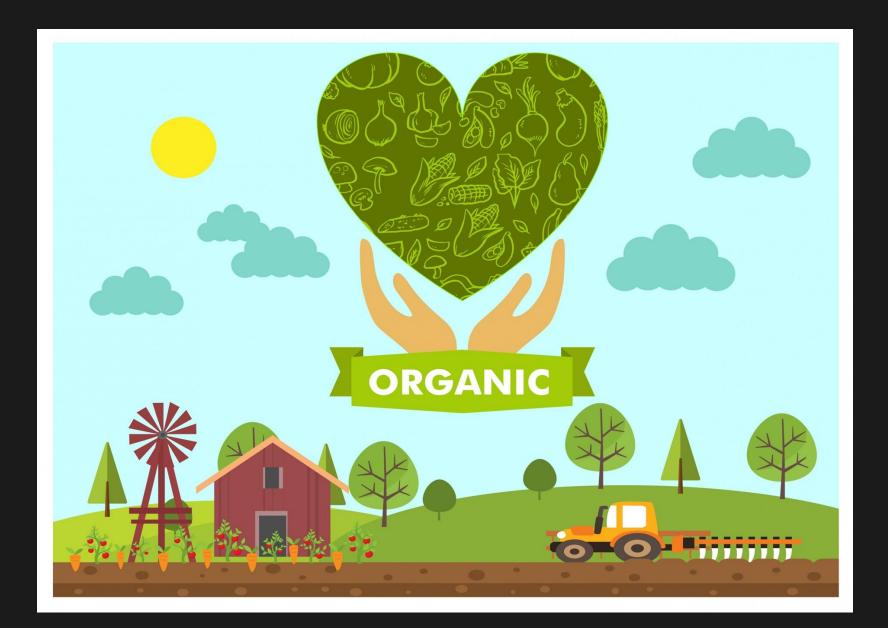




Initial Thoughts



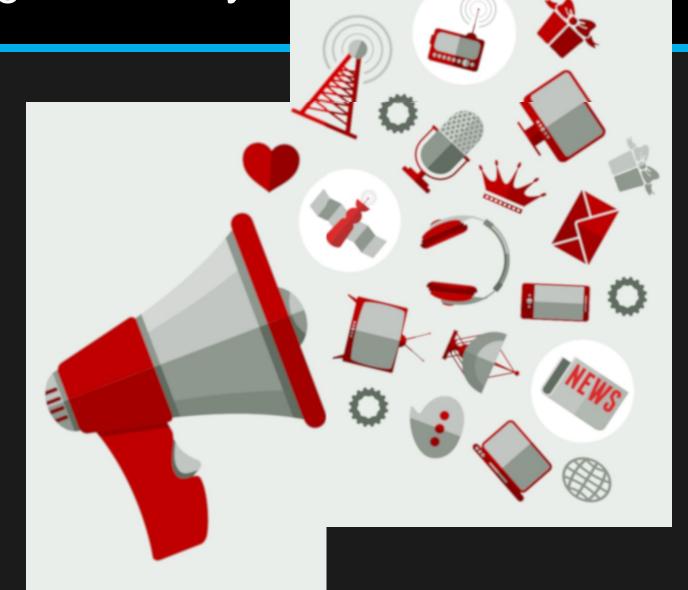






Creating Your Story



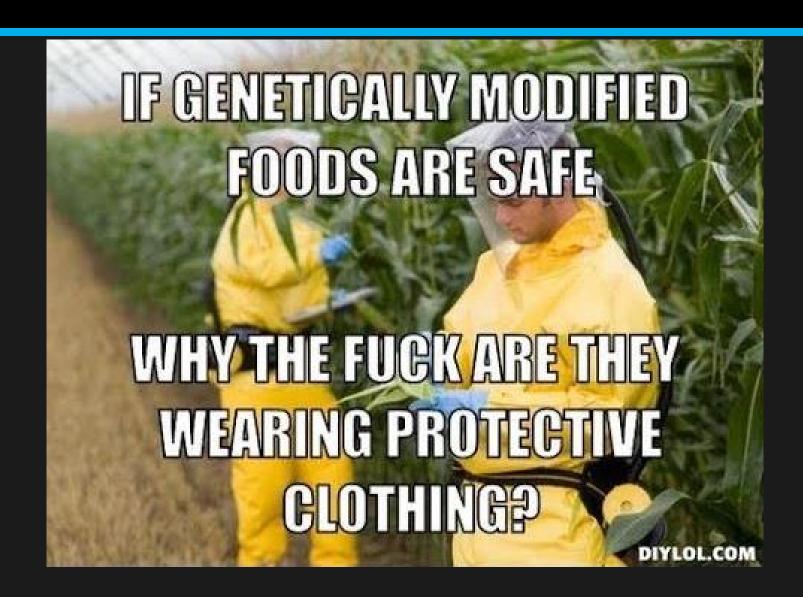


BEFORE SOMEONE ELSE DOES







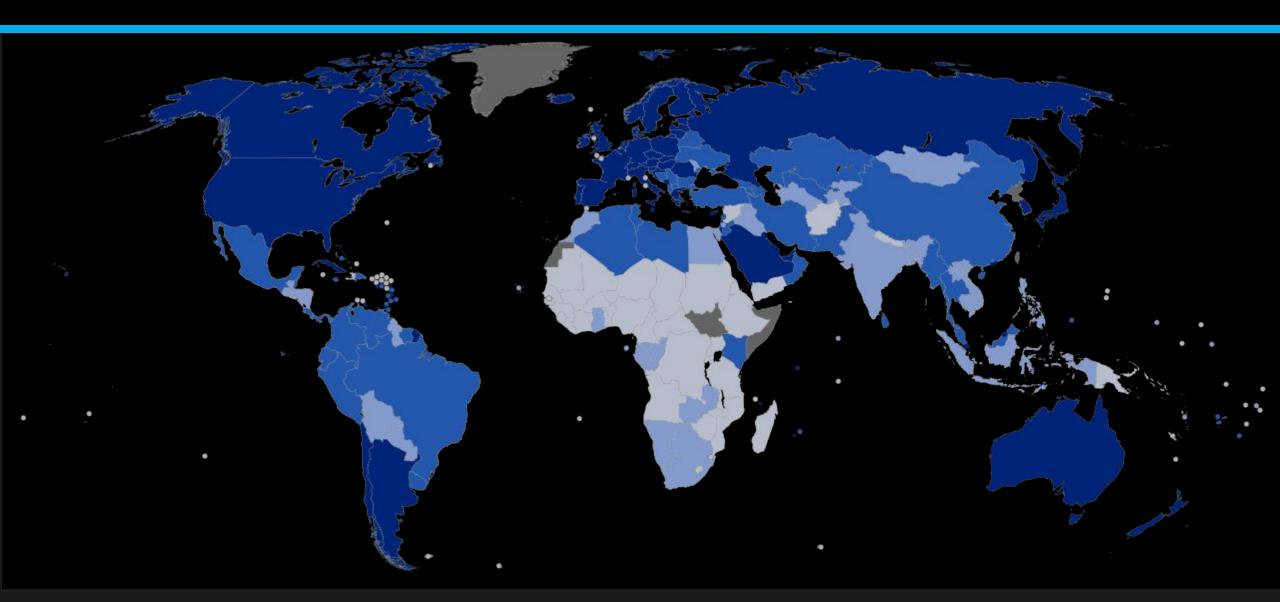


Different Stakes + Different Tactics











IRL/GB Salad dressing.

Ingredients: water, vegetable oils contains geneticly modified soyabean oil), sugar, vinegar, modified starch, wheat starch, salt, mustard (water, mustard seed, vinegar, salt, spices, herbs), egg yolk, thickener [E412], acids (E330), preservatives (E202), colours (E160a), antioxidant (E385).

Produced in: The Netherlands. Store in a cool, dry place. Shake before use.









Banana Wilt



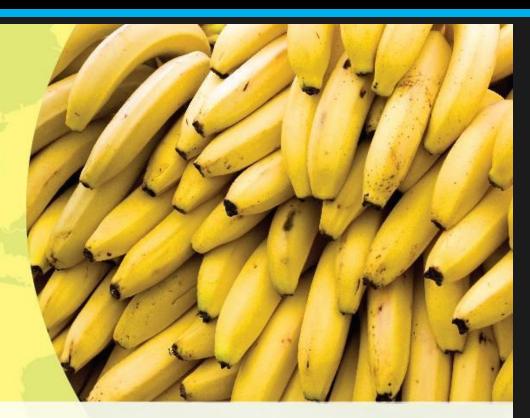








Banana bacterial wilt is devastating one of East Africa's staple crops.



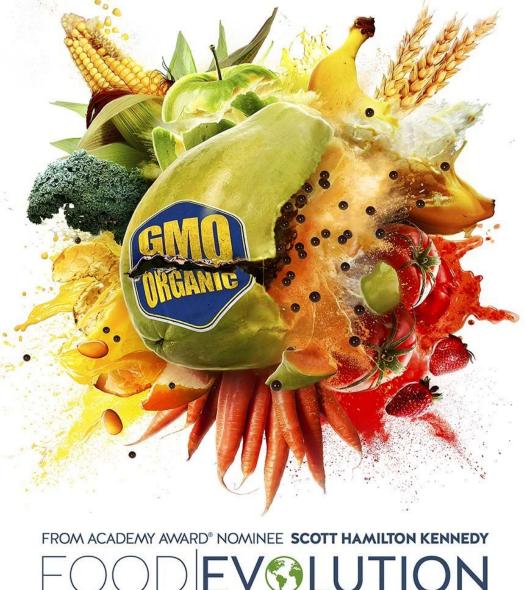


Meet Leena Tripathi, a scientist who's working on a gene from a sweet pepper that could help save millions of bananas.

LEARN WHY THIS RESEARCH IS SO A-PEELING AT GMOANSWERS.COM.



FEAST ON **FACTS**







https://www.foodevolutionmovie.com/









FROM ACADEMY AWARD® NOMINEE SCOTT HAMILTON KENNEDY



NARRATED BY **NEIL DEGRASSE TYSON**

"Choose a healthy Future for our Kids and the Planet"



"good" organics, "bad" conventional agriculture



By adding just one more organic product to your grocery cart each week you are supporting a healthier future for all of us. onlyorganic.org #onlyorganic



"I don't believe that these marketing strategies reflect the ethics of real organic farmers, certainly none that I've met"



"They're [alternative food companies] building their businesses trying to portray me and my family as bad guys — for simply doing what's best for our farm, community, family and customers."



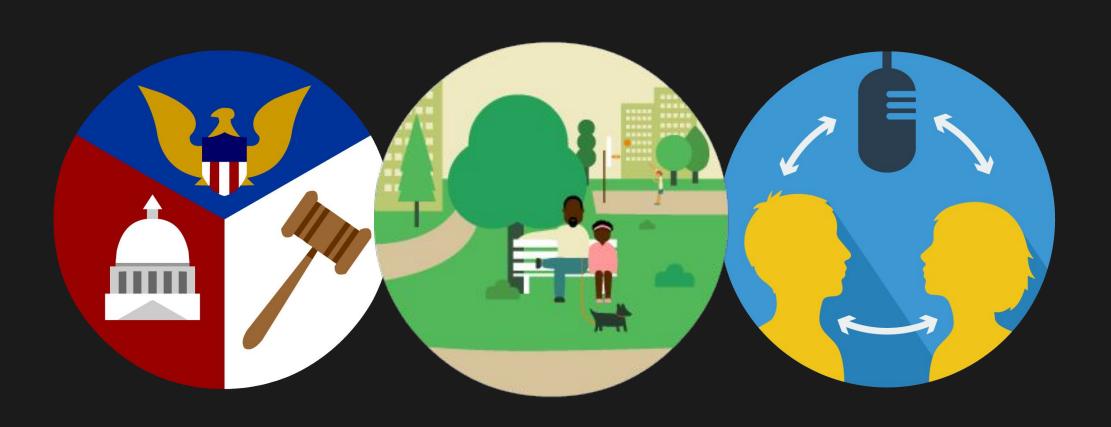
Understand Your Audience





Lawmakers, The public, and newsmakers





GMO RESEARCH, REVIEW AND REGULATION | How Does a GMO Get to Market?

On average, GMOs take



and \$130 million



of R&D BEFORE

coming to market

The regulatory process alone can take 5 to 7 years

REGULATORY SCIENCE

75+ different studies are conducted to demonstrate each new GMO is:



Safe to grow

- Crop grows the same as non-GM varieties
- Crop exhibits expected characteristics (e.g., insect resistance)

Safe for the environment and beneficial insects



Safe to eat

- · Same nutrients as non-GM crops
- . No new dietary allergens



REGULATORY REVIEW

More than 90 government

bodies² globally

review and approve

GMOs. In many countries,

multiple agencies are involved in the regulation of GMOs.

GMOs have been grown or imported by **70 countries**³ since 1996.



U.S. REGULATORY AGENCY REVIEWS



Safe to



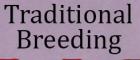
Safe for the



Safe to eat









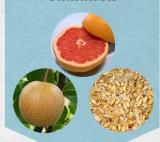
Crossing plants and selecting offspring

Almost All Crops

Mutagenesis



Exposing seeds to chemicals or radiation



RNA Interference



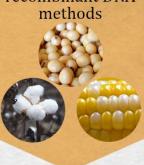
Switching off selected genes with RNA



Transgenics

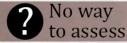


Inserting selected genes using recombinant DNA methods



Number of Genes Affected

10K - >300K



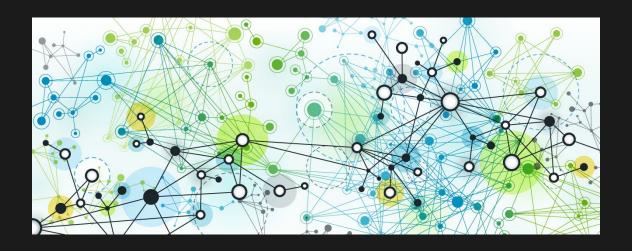
1-2

1-4

Desired gene(s) inserted with other genetic material. No safety testing requirements.

Random changes in genome, usually unpredictable. No safety testing requirements. Targeted gene(s) switched off or 'silenced'. Safety testing required. Desired gene(s) inserted only at known locations. Safety testing required.

Criticism: Very One Sided Cohort



Thoughts + Questions?