

# WIKI Part 2: Treat cells for $\gamma$ H2AX assay

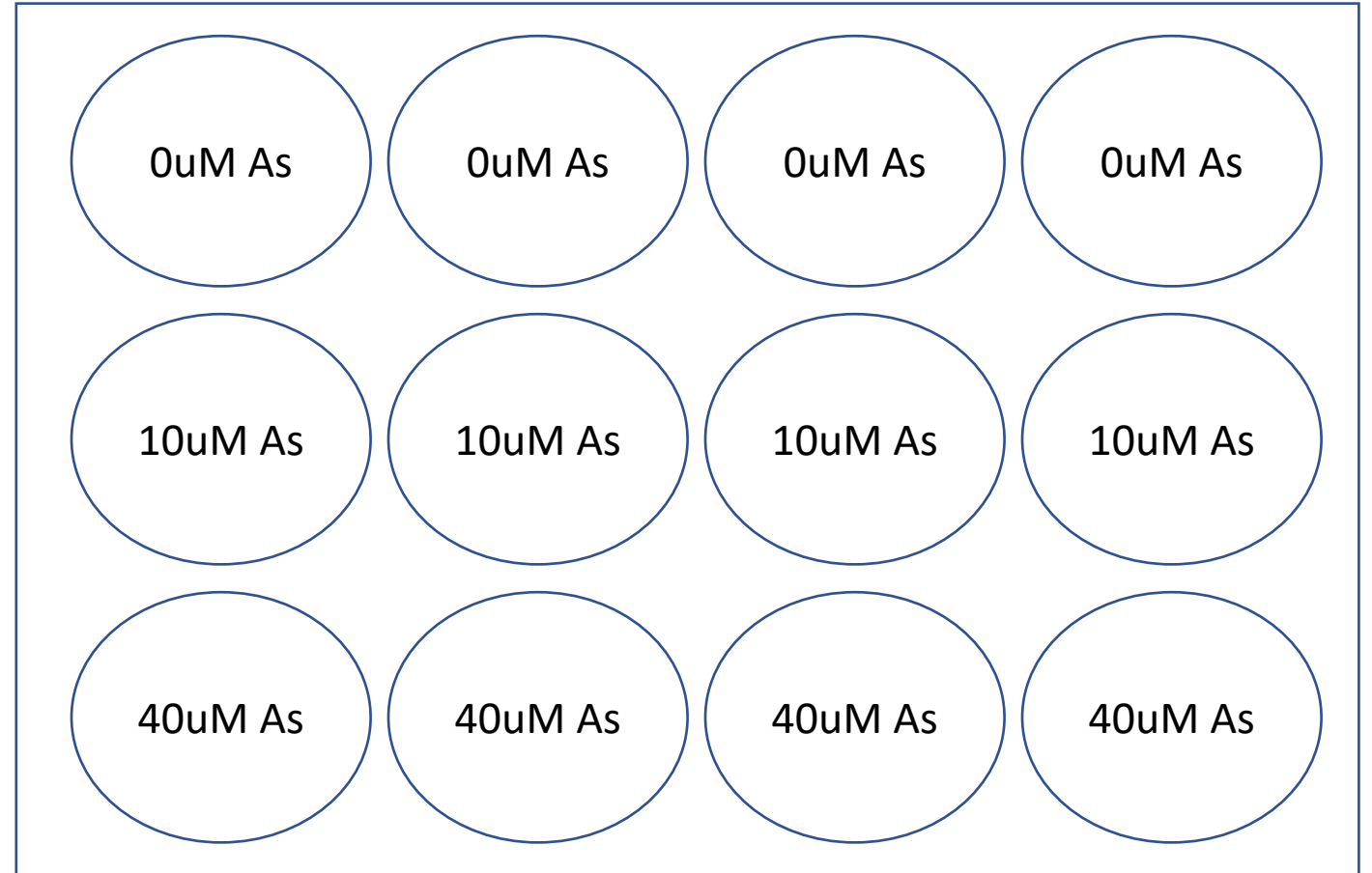
## Expose cells to As

Before going into TC,

- Calculate the dilution of As stock needed to prepare a final concentration of 10  $\mu$ M and 40  $\mu$ M in 5 mL of media.

- Stock concentration of As is 10 mM.

- In TC, aspirate the spent media from your coverslips and replace it with media containing arsenic according to the chart to the side (1ml for each well)

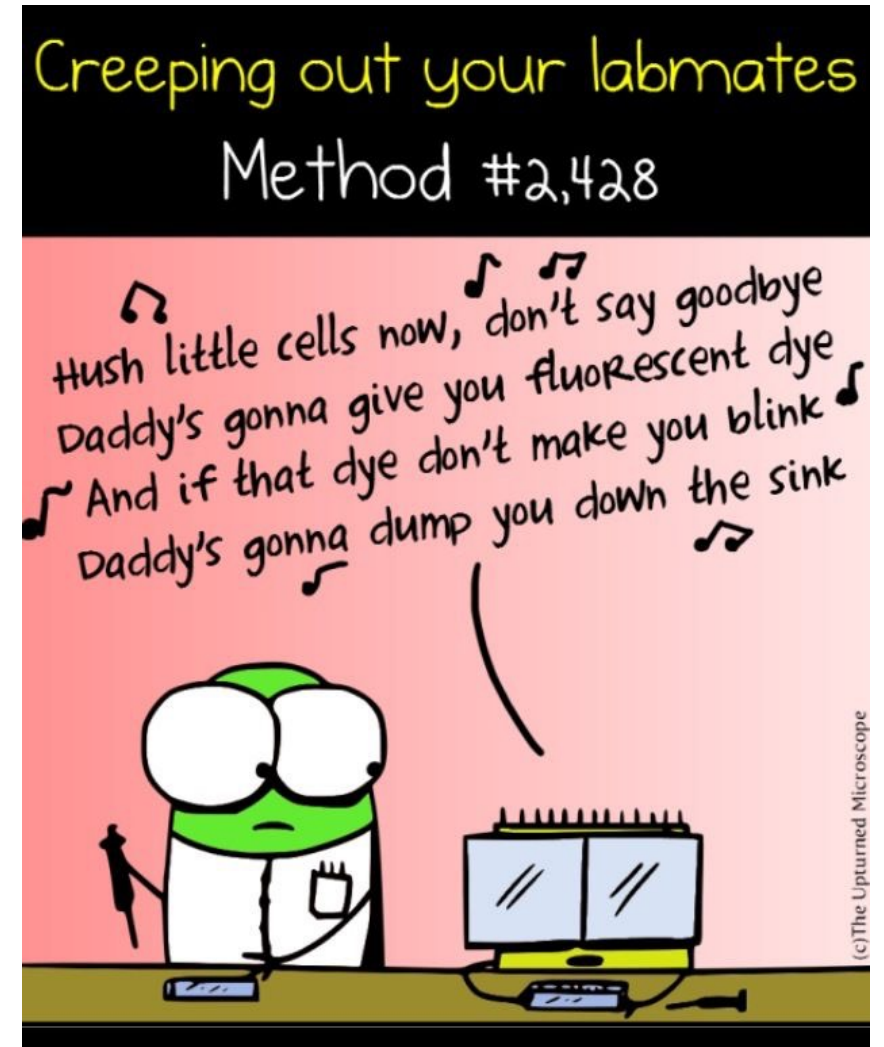


# M1D2: Prepare and treat cells for $\gamma$ -H2AX experiment

1. Prelab
2. Treat and fix primary MEF cells for H2AX assay
3. Work on Background and Motivation section

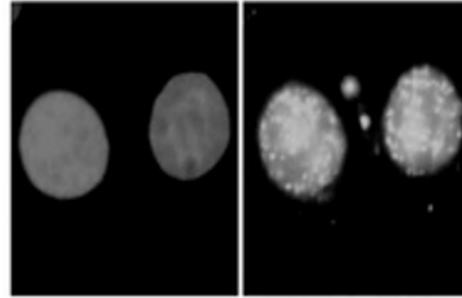
## Admin Notes:

- Sign EHS training roster for lab-specific training
- If you haven't already, respond to Noreen's office hours email



# Mod1 Overview

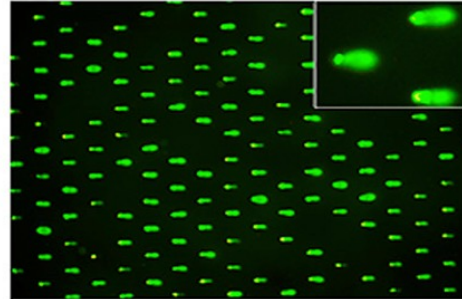
Last lab:



## 1. Use repair foci experiment to measure DNA breaks

- Examine effect of  $H_2O_2$  +/- As on double strand DNA breaks by measuring  $\gamma$ H2AX foci formation

This lab:



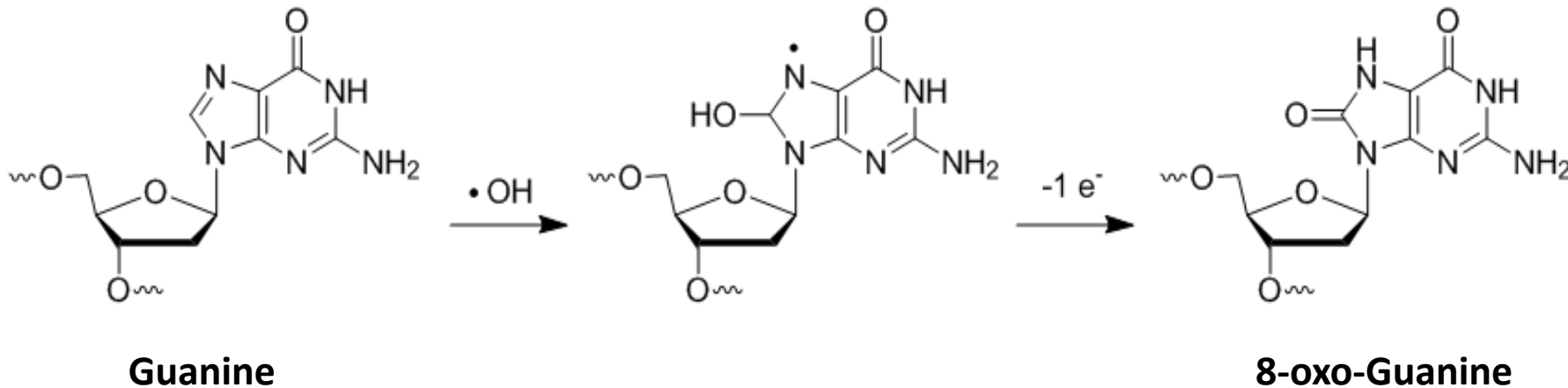
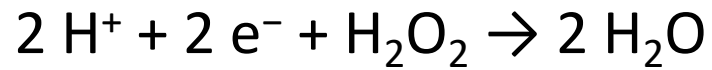
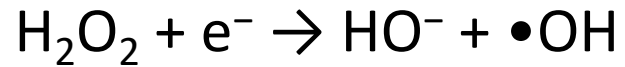
## 2. Use high-throughput genome damage assay to measure DNA damage

- Measure effects of  $H_2O_2$  +/- As on DNA damage by measuring DNA migration in agarose matrix

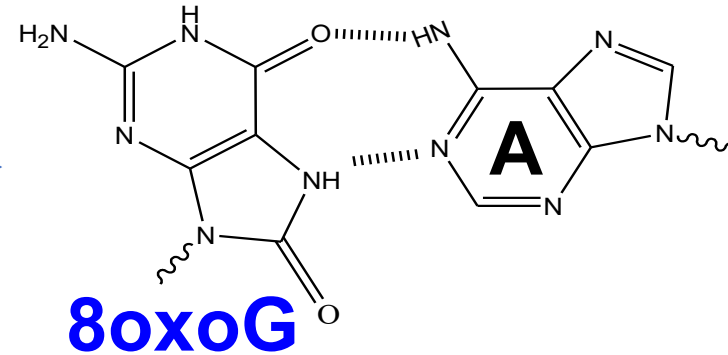
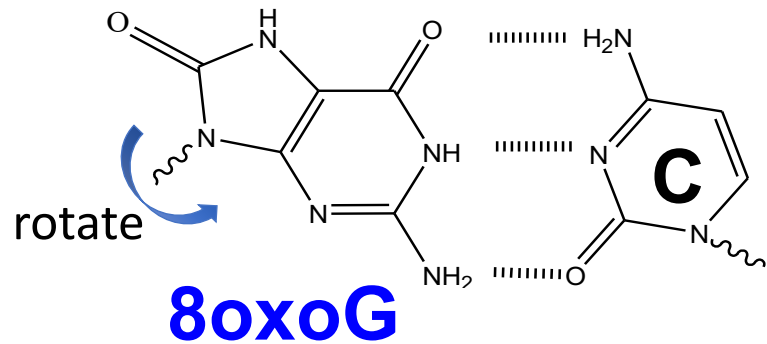
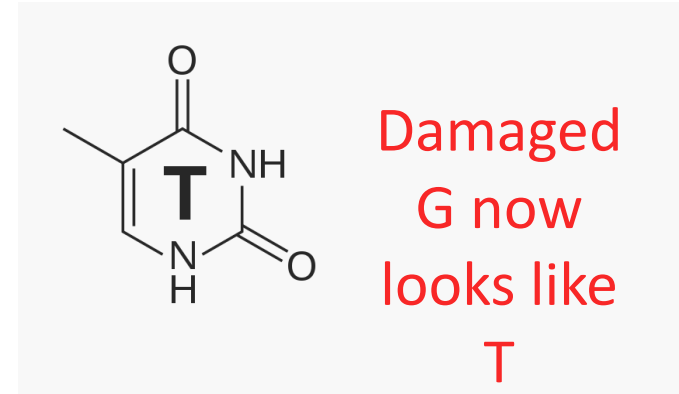
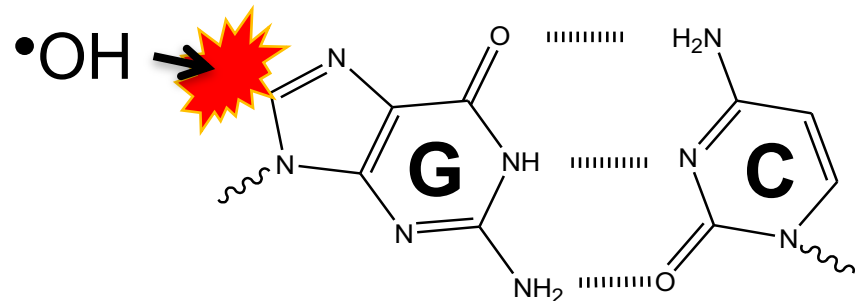
Next lab:

# How does H<sub>2</sub>O<sub>2</sub> damage DNA?

By producing Reactive Oxygen Species



# How does H<sub>2</sub>O<sub>2</sub> damage DNA?

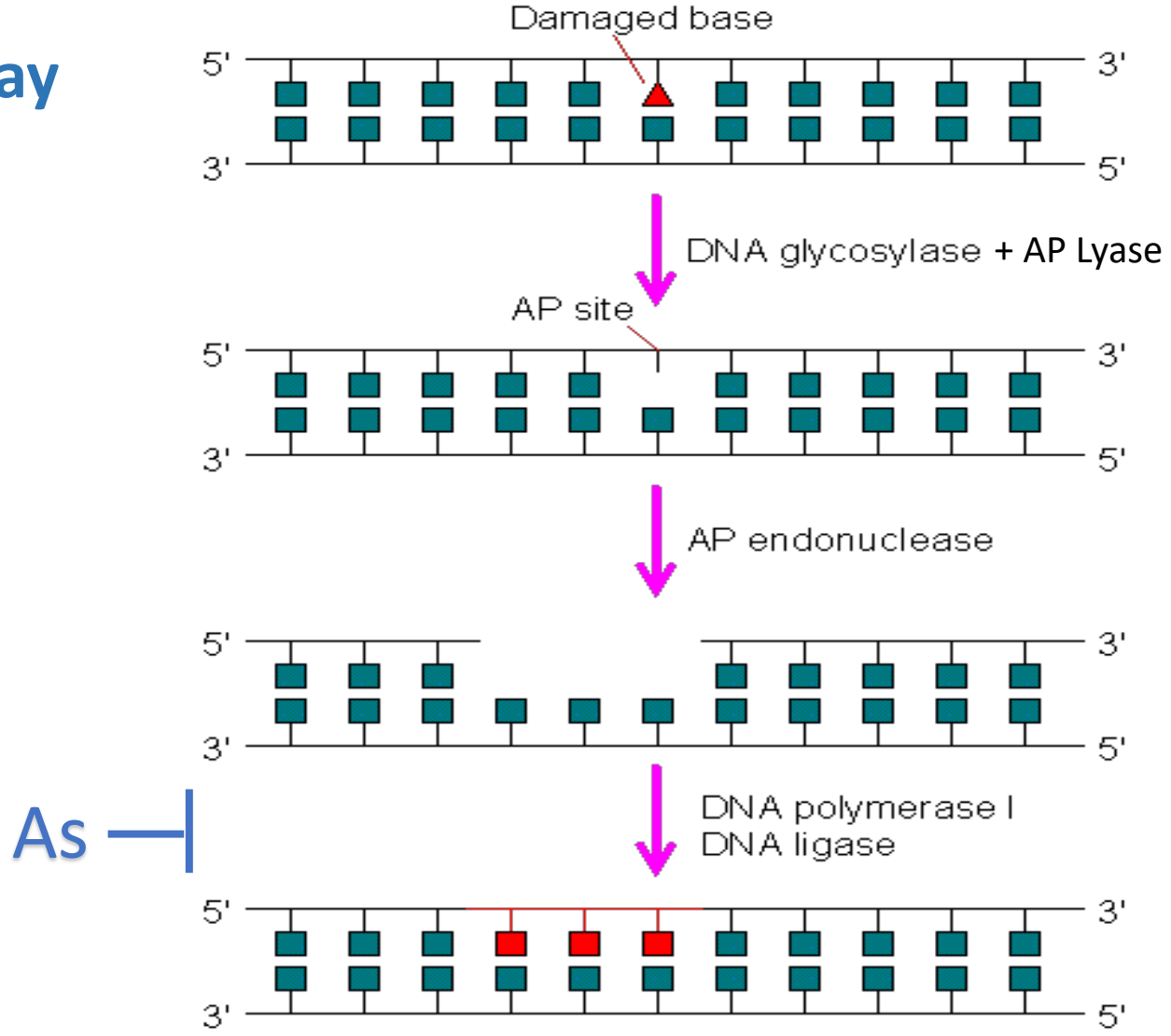


Mutation if replicated



# How do cells repair oxidative DNA Damage?

## The BER pathway



As —|

# How do we look at DNA damage in intact cells?

## Look for $\gamma$ -H2AX foci

H<sub>2</sub>O<sub>2</sub> can cause damage to DNA, resulting in a damaged base



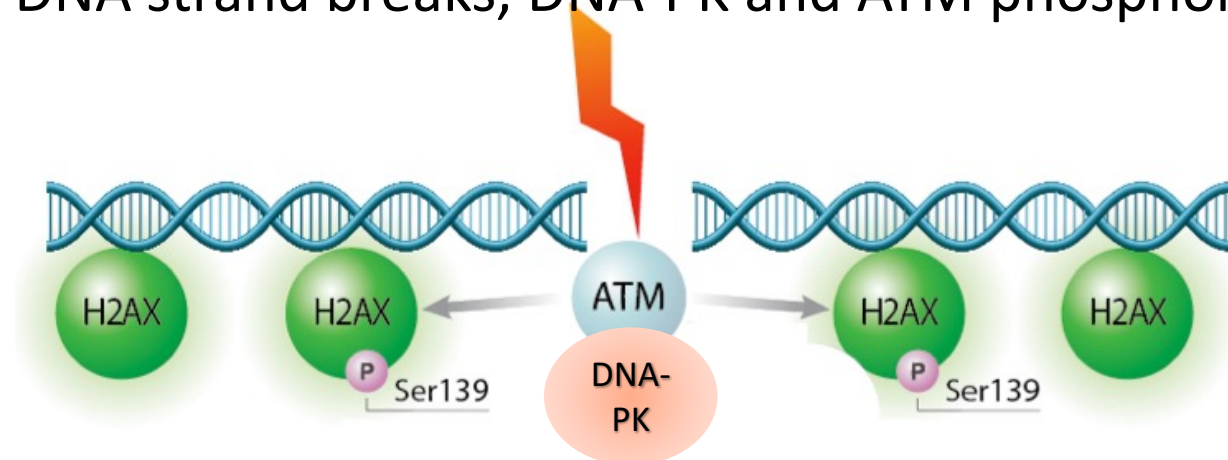
That damage causes a single strand break as cell tries to repair the DNA



Multiple single strand breaks cause double strand breaks



At DNA strand breaks, DNA-PK and ATM phosphorylate the histone H2AX



We can identify the frequency of these breaks as a measure of DNA damage

# Treatment conditions for this experiment

- Goal: Identify potential additive/synergistic effect pretreatment with As has on H<sub>2</sub>O<sub>2</sub> induced DNA damage
  - Treat cells with As for 2 hours, then treat cells with H<sub>2</sub>O<sub>2</sub> for 30 minutes

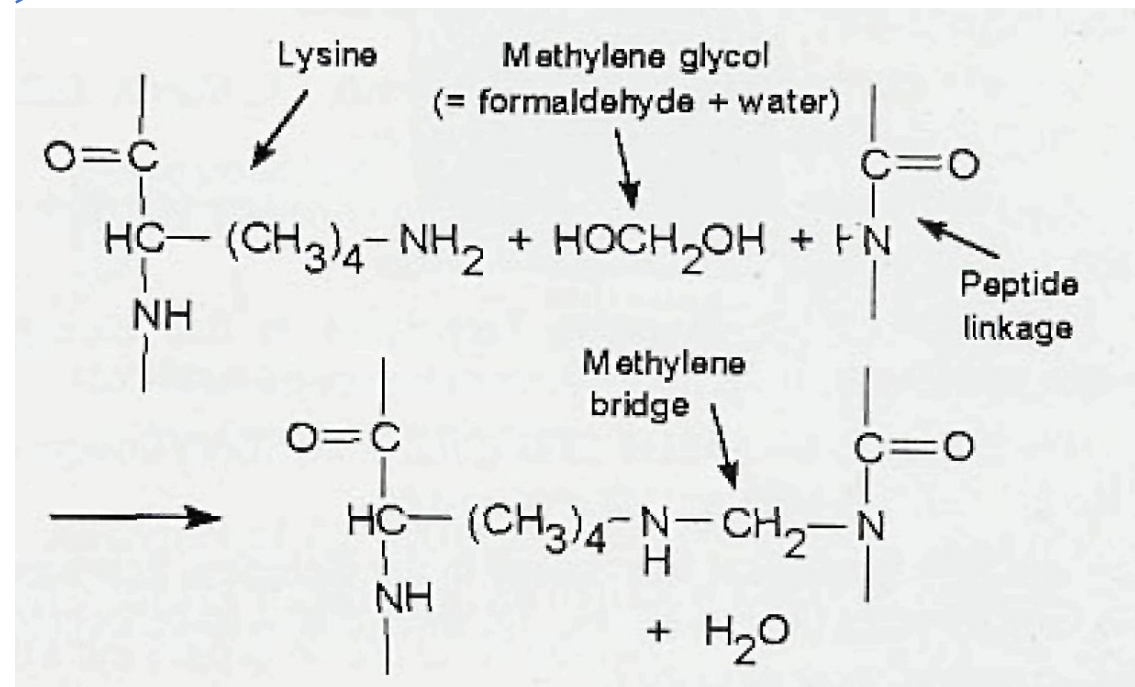
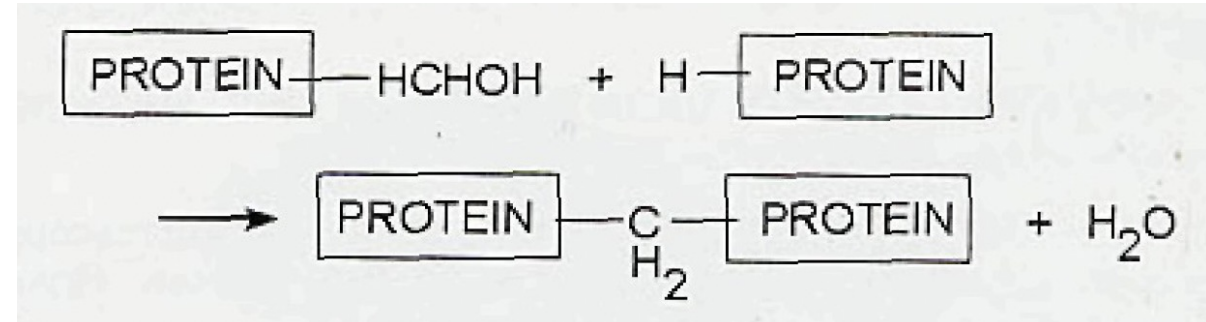
Experimental Condition

Control Conditions



# Fixing cells with paraformaldehyde

- Formaldehyde is a chemical crosslinker
  - Paraformaldehyde is the polymer version—depolymerize into formaldehyde with heat or basic conditions
- Commonly used in cell and tissue microscopy
  - Crosslinking of proteins will “fix” samples in place for static imaging
  - Can also trap nucleic acids, etc... between crosslinked proteins



# In lab today

1. Expose As-treated cells to  $\text{H}_2\text{O}_2$  and fix them for IF staining
2. Talk about purpose and structure of background and motivation during As incubation

**DO NOT** use the aspirator to remove cell media with Arsenic.  
It needs to be collected as hazardous waste!

# M1D3HW

1. Work with your lab partner to write methods for M1D1 and M1D2
2. Schedule appointment with BE Comm Fellow before M1D5

# Pro tips for writing a methods section

Include enough **information to replicate** the experiment

- List manufacturer's name (Company)
- Be **concise and clear** in your description

Use **subsections** with descriptive titles

- Put in logical order, rather than chronological order
- Begin with topic sentence to introduce purpose / goal of each experimental procedure

Use clear and **concise full sentences**

- NO tables or lists, all information should be provided in full sentences and paragraphs
- Write in passive voice and use past tense

Use the most **flexible units**

- Write concentrations (when known) rather than volumes

Eliminate 20.109 specific details

- Example "labeled Row A, Row B..."
- Do not include details about tubes and water!
- Assume reader has some biology experience
- Include parts of the protocol that the teaching faculty completed, but do not say "completed by teaching faculty."

# How can you improve this example?

“Cells were grown in 12 mL of DMEM supplemented with FBS. We spun down the cells and counted them with a hemocytometer. Flasks were incubated in 37 C incubator.”

# Revised example...

## Maintaining primary MEF cells

Primary Mouse Embryonic Fibroblasts (MEF) cells (gift of Engelward Laboratory, MIT) were grown in Dulbecco's... (DMEM) (Manufacturer) supplemented with 20% fetal bovine serum (FBS) (Manufacturer) and 100 U / mL of penicillin and streptomycin (Manufacturer). To harvest, cells were trypsinized, collected in fresh media, and counted using 10% (v/v) trypan blue and a hemocytometer. Cultures were maintained at 37 C, 5% CO<sub>2</sub>, and 95% relative humidity.



What is the **purpose** of the Background & Motivation section?

What ***specific information*** should be included in the Background & Motivation section?

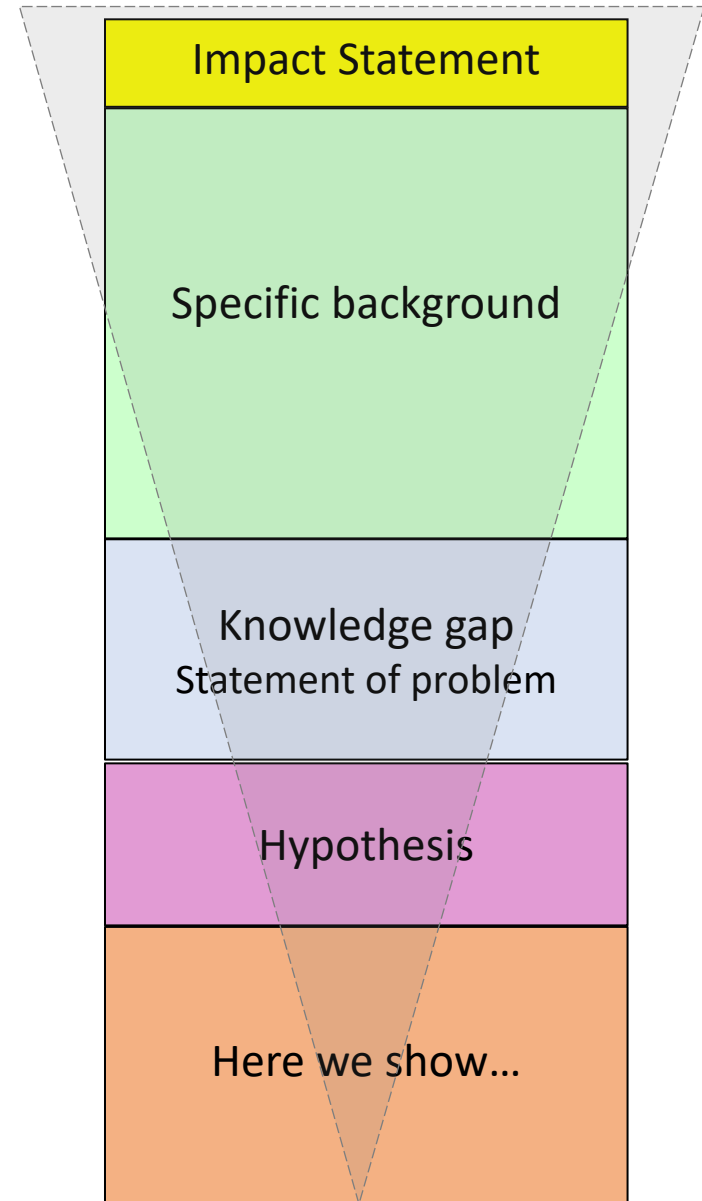
# Notes on Background & Motivation section...

- Anchor your research in a general topic that is important to a broad audience
  - Focus on describing what is currently known in the field
  - Reference the relevant research in the field
- Connect your research to the general topic
  - Minimum essential information
  - Introduce specific technologies necessary for understanding your specific project
- Address how you will expand on what is currently known
  - Include evidence of incompleteness of current understanding
  - Motivate your investigation
  - **Include a clear hypothesis / research goal**
- Provide a preview of your findings and the implications
  - Tie back to the initial general topic
  - Avoid including extensive methods details



# Notes on topic sentences...

- **Topic sentence = first sentence of each paragraph**
- Should 'funnel' from big picture topic to your specific research question / project
  - Provide only the background needed to understand research / problem / goal
  - Clearly state what is not currently known
  - Address how you will fill knowledge gap
  - Provide preview of your results
- Include references!!



# How should you introduce your story?

1<sup>st</sup> paragraph: what is the big picture / problem?

2<sup>nd</sup> paragraph: what is currently known?

3<sup>rd</sup> (or 4<sup>th</sup>) paragraph: what is your research question?

4<sup>th</sup> (or 3<sup>rd</sup>) paragraph: how will you address your question?

5<sup>th</sup> paragraph: here we show...

