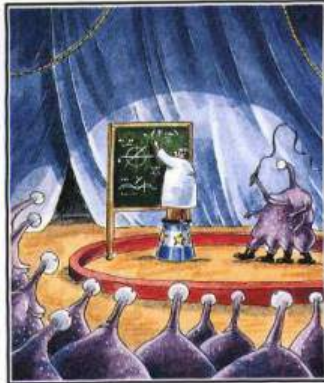


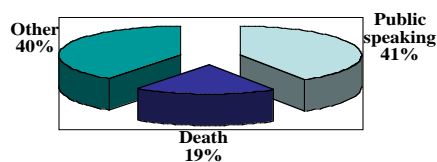
Giving technical presentations: Do's, Don'ts, and Why Nots?



Abducted by an alien circus company,
Professor Doyle is forced to write calculus
equations in center ring.

Neal Lerner, Northeastern Writing Program
n.lerner@neu.edu; 617-373-2451

First thing to do to prepare an oral presentation is to overcome fear



On October 7, 1973, the *Sunday Times* in London published a survey asking 3,000 Americans: "What is your greatest fear?"
The result: 41% of respondents answered "speaking in public."

An oral presentation to evaluate

<http://www.youtube.com/watch?v=tyzKrQv9zFI&feature=related>

Five Canons of Classical Rhetoric or the art of persuasive speech

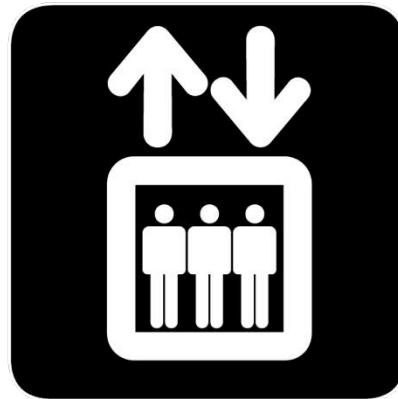
1. *Invention*: Generating content and honing your message
2. *Arrangement*: Organizing the material
3. *Style*: Using “proper words in proper places” (J. Swift).
4. *Memory*: Speaking without notes.
5. *Delivery*: Controlling voice, gesture, expression--and graphics!



Invention: What's your essential message?

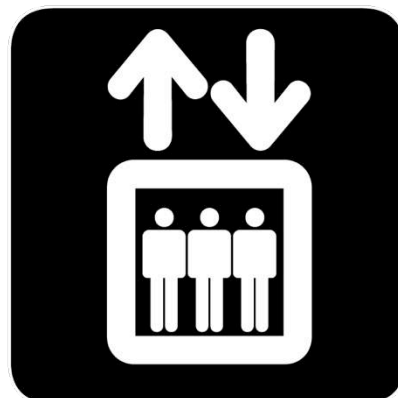
Prepare an **elevator speech** or a concise summary of your project in spoken language.

Practice your elevator speech.



Invention: What's your essential message?

Exercise: Give a two-minute version of your research to as many people in the room as possible.



Arrangement: Control the story that you want to tell about your content (in three parts).

1. Develop a general goal.

- Inform? Persuade? Brainstorm?

2. Develop a precise objective.

- e.g., After my presentation, the listeners will be able to identify my three major conclusions and their implications.

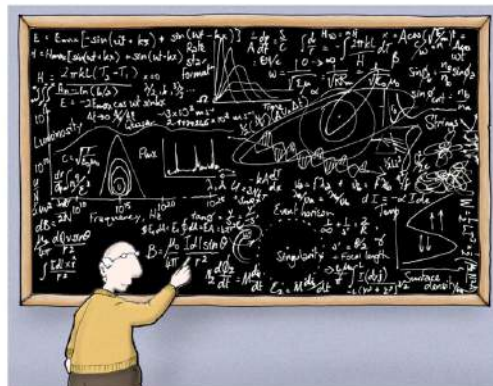
3. Consider the questions your presentation will answer for your audience.



Arrangement Step 1: Organize your data & assemble a storyboard

Start with figures:

- Assemble hard copies of your figures in a “storyboard.”
- Figure out the major technical theme of your presentation.
- Assess how each figure contributes to the major theme.
- REVISE figures to focus on the major theme (develop figures that summarize that major theme).



<http://www.nearingzero.net/index2.html>

<h3>Effect of Temperature of Velocity of Propagation</h3> <p>Dennis M. Freeman 6.021J November 10, 2004</p>	<h3>Outline</h3> <ul style="list-style-type: none"> • Background • Methods • Results • Discussion
<h3>Methods</h3> <ul style="list-style-type: none"> • Apply current stimulus in "Propagated Action Potential" simulation of HH model. • Measure velocity of propagated action potential • Repeat for temperatures from 0 to 50 C. • Plot results 	<h3>Results</h3> <ul style="list-style-type: none"> • Velocity increased with temperature for range from 0 to 30 C. • Velocity decreased with temperature for range from 30 to 35 C. • Action potentials did not propagate for temperatures above 36 C.
<h3>Results (continued)</h3> <ul style="list-style-type: none"> • Why did high temperatures kill the action potentials? <ul style="list-style-type: none"> - Stimulation normally generates wave of activation (due to sodium currents) and wave of inactivation (due to potassium currents). - Temperature increases speed of both waves, but rate of increase is faster for inactivation 	<h3>Conclusions</h3> <ul style="list-style-type: none"> • Velocity increased with temperature for range from 0 to 30 C. • Velocity decreased with temperature for range from 30 to 35 C. • Action potentials did not propagate for temperatures above 36 C.

<h3>Effect of Temperature of Velocity of Propagation</h3> <p>Dennis M. Freeman 6.021J November 10, 2004</p>	<h3>Outline</h3> <ul style="list-style-type: none"> • Background • Methods • Results • Discussion
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<h3>Results (continued)</h3> <ul style="list-style-type: none"> • Why did high temperatures kill the action potentials? <ul style="list-style-type: none"> - Stimulation normally generates wave of activation (due to sodium currents) and wave of inactivation (due to potassium currents). - Temperature increases speed of both waves, but rate of increase is faster for inactivation 	<h3>Conclusions</h3> <ul style="list-style-type: none"> • Velocity increased with temperature for range from 0 to 30 C. • Velocity decreased with temperature for range from 30 to 35 C. • Action potentials did not propagate for temperatures above 36 C.

No Outline Slide needed

Your audience expects IMRaD format

T+IMRaD = Talk Structure

- **Title** establishes the focus
- **Introduction** provides general field or context.
- **Methods** follows a particularized path.
- **Results** describe specific findings.
- **Discussion** moves from specific findings to wider implications.

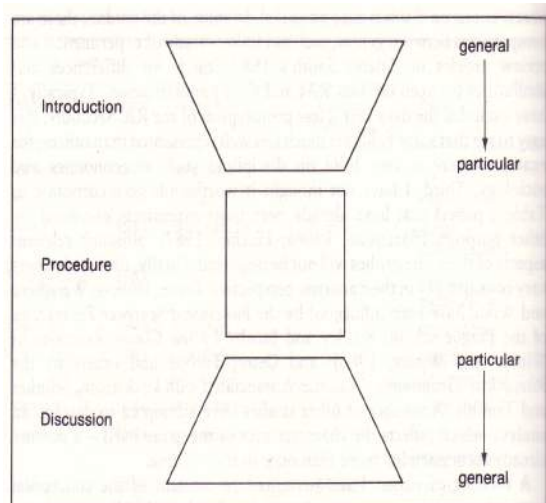


Figure 7 Overall organization of the research paper (Hill et al., 1982).

Titles are mini-abstracts

- Informative
- Specific
- **Understandable at a glance**
- Your name and partner's name
- Date
- Institutional Affiliation

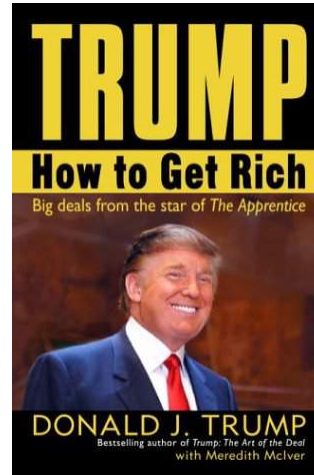
Thermal Blockage of Propagated Action Potentials in the Hodgkin-Huxley Model

Dennis M. Freeman and Aek Thitimon

6.021J: Quantitative Physiology
Massachusetts Institute of Technology
November 21, 2004

Titles are mini-abstracts

- Informative
- Specific
- **Understandable at a glance**
- Your name and partner's name
- Date
- Institutional Affiliation



Introduction provides context & purpose of research

- If you wish, you may have a background slide.

Purpose slide

- Explains the ***purpose of the project*** (aims?)
 - Relate to Results + Discussion points
- Show logic of ideas in words or text
- Meaningful graphic OK; bullet points OK

The “ideal” introduction follows . . .

Create a Research Space

1. Re-establish significance of research field.
2. Situate actual research in these terms.
3. Show how this niche will be occupied and defended.

Swales (1990)

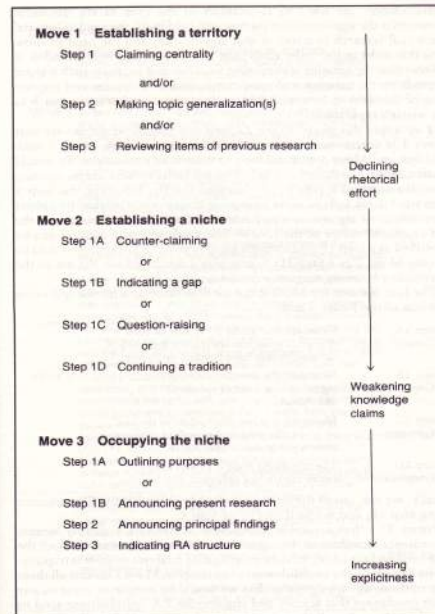


Figure 10 A CARS model for article introductions

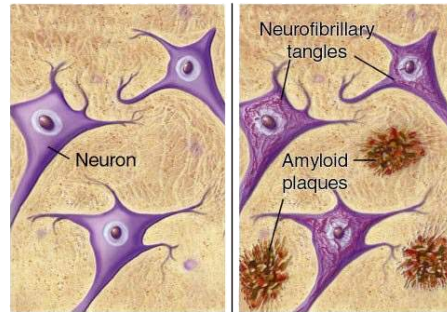
- Sample Introductions:
 - 1 slide
 - 2 slide
 - Multi-slide



MGH/MIT/HMS Athinoula A. Martinos Center
For Biomedical Imaging

Alzheimer's Disease Difficult to Detect in Early Stages

- **Problem:** Lack of inexpensive, effective imaging techniques for Amyloid- β (Alzheimer's cause)
- Alzheimer's normally diagnosed in latter stages, when symptoms are severe
- **Overall Project Goal:** to enable non-invasive, effective amyloid burden imaging techniques
- **Summer Aim:** to calibrate instrumentation and automate imaging system



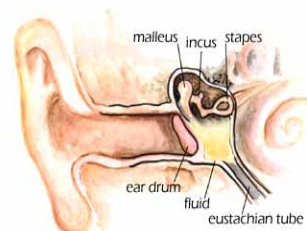
Figure¹ depicting both the accumulation of amyloid- β as plaques in neural tissue and neurofibrillary decay

Courtesy of F. Hammond

¹ - American Health Assistance Foundation

Otitis Media (OM)

- inflammation of the middle ear space, usually associated with fluid backup
- different types: acute, chronic, with effusion
 - different types of fluids: purulent, mucoid, serous

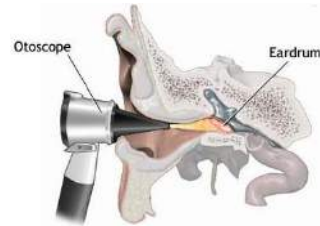


- 30 million physician visits and \$5 billion spent on diagnosis/treatment annually

Courtesy of S. Hon

The Problem:

- current standard: otoscopy
- misdiagnosed up to 50% of the time
- leads to inappropriate prescription of antibiotics



<http://www.uvm.edu/cms/luse/otoscope-examination.jpg>

The Long-Term Goal:

- develop an optical device that enables accurate diagnosis of otitis media, leading to more effective treatment

Courtesy of S. Hon



Reversible Phosphorylation is an Important Regulatory Mechanism



Edwin Krebs



Edmond Fischer

Krebs, E. G. and Fischer, E. H. The phosphorylase b to a converting enzyme of rabbit skeletal muscle. *Biochim Biophys Acta*. 150-7 (1956).

Reversible phosphorylation can modulate a protein' s:

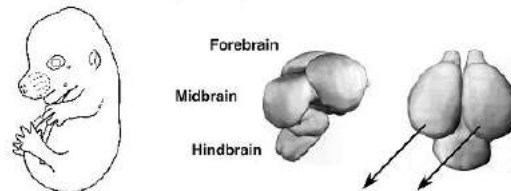
- Enzymatic activity
- Interaction with other proteins
- Stability or its Destruction
- Subcellular Localization

Courtesy of R. Carey

Global Tyrosine Phosphorylation Extraction

- Emerging technologies to identify phosphorylation sites for more than a single protein
- Technology used for first time in primary animal tissue,
- 239 phosphorylation sites extracted in one experiment

Primary Embryonic Forebrain



"Phosphoproteomic analysis of the Developing Mouse Brain", Bryan Ballif, ppt presentation

Courtesy of R. Carey

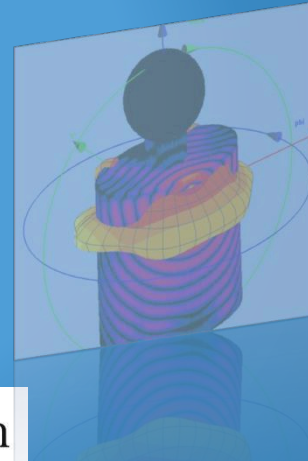
Project Goals

- Develop a method to calculate a conservation score for phosphorylated residues
- Automate method for high throughput analysis
- Find biologically meaningful correlations in the resulting conservation scores

Courtesy of R. Carey

Multi-path 2-Port Channel Characterization for Galvanic Coupled Intra-body Communication

Meenupriya Swaminathan
 Joan Sebastia Pujol
 Gunar Schirner
 Kaushik R. Chowdhury



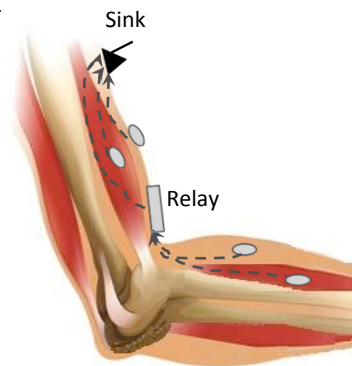
Courtesy of M. Swaminathan



Northeastern

Intra-body Network (IBN)

- IBN allows implanted sensor to actuator communication
- High frequency signals (Eg. RF signals)
 - Get absorbed by wet tissues
 - Requires high transmission power
 - Propagates through air and not secured



✓ Need low power consuming communication paradigm that is suitable for signal propagation through tissues.

Courtesy of M. Swaminathan

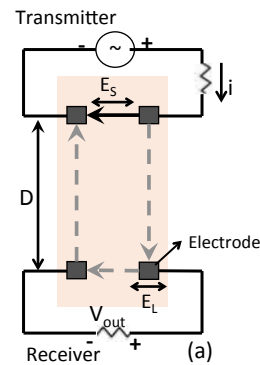
January/29/2015

Multi-path 2-Port Channel Characterization for Galvanic Coupled Intra-body Communication

24

Galvanic Coupling (GC)

- A pair of electrodes directly couple weak electrical signals to tissue
- The difference in voltage propagates through tissue and is detected by a pair of receiver electrodes



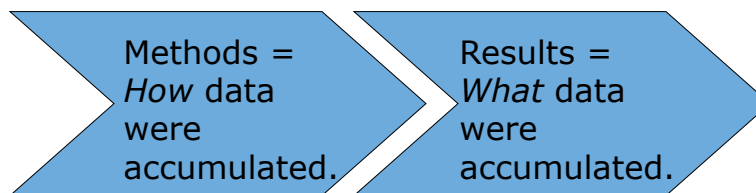
Courtesy of M. Swaminathan

January/
29/2015

Multi-path 2-Port Channel Characterization for Galvanic Coupled Intra-body Communication 25

Methods describes FINAL APPROACH you took in your experiment.

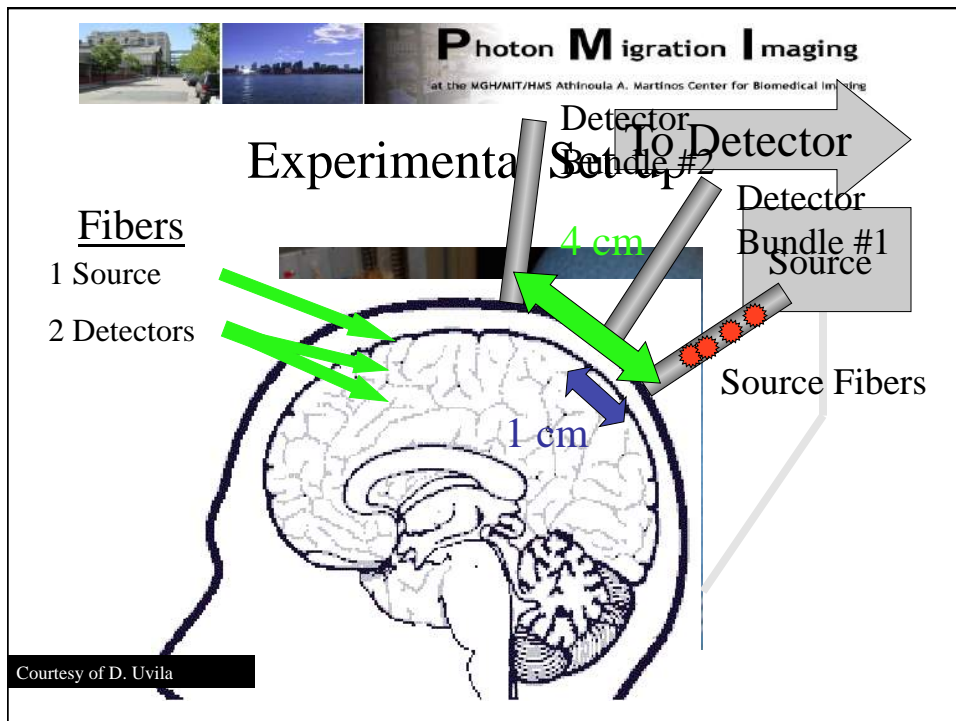
The Methods Section is cool to you but often dull to the audience.




- Approximately 1-2 slides (may be supplemental material)
- Distill Methods to key procedures
- Equations should be extremely simple and friendly
- Visual representations of methods are easiest to comprehend


Methods

- Apply current stimulus in “Propagated Action Potential” simulation of HH model.
- Measure velocity of propagated action potential
- Repeat for temperatures from 0 to 50 C.
- Plot results





BRIGHAM AND
WOMEN'S HOSPITAL
Specializing in care for a lifetime



HARVARD
MEDICAL SCHOOL
TEACHING AFFILIATE

Extracting the Sequence

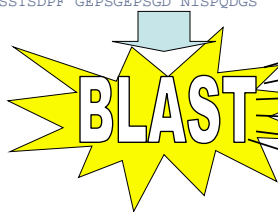

Phosphorylation Data: Gene Phos-Tyrosine

Dab1 KKEGVY*DVPKS

Entire Sequence:

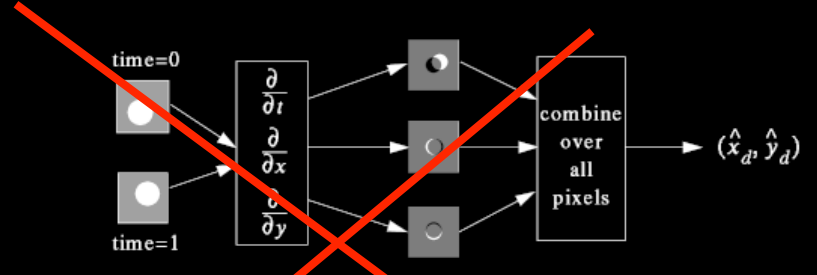
```

MSTETELQVA VKTSAKKDSR KKGQDRSEAT LIKRFKGEV RYKAKLIGID EVSAARGDNL CQDSMMKLG VVAGARSKGE
HKQKIFLTIS FGGIKIFDEK T GALQHHHAV HEISYIAKDI TDHRAFQYVC GKEGNHRFVA INTAQAAEPV ILDDADLFQL
IYELKQREEL EKKAQKDKQC EQAVYQTILE EDVEDPVYQY IVFEAGHEPI RDPETEENIY QVPTSQRKKEG VYDVPK SQPN
SQPLEDFESR FAAATPNRNL SMDFDELLEA TKVSAVTQLE LFGDMSTPPD ITSPTPATP GDAFLPSSSQ TLPGSADVFG
SMSFGTAAVP SGYVAMGAVL PSFWGQQLV QQIAMGAQP PVAQVIPGAQ PIAWGQPLF PATQQAWPTV AGQFPFAAFM
PTQTVMLAA AMFQCPLTPL ATVPGTNSA RSSQSDKPR QKMGKESFKD FQMVQPPVP SRKPDQPSLT CTSEAFSSVF
NKVGAQDTD DCDDFDISQL NLTPVTSTTP STNSPPTPAP RQSSPSKSSA SHVSDPTADD IFEEGFESPS KSEEQEPADG
SQASSTSDPF GEPSGEPGD NISFQDGS
    
```

Courtesy of R. Carey

Gradient-Based Estimation



Constant brightness: $E(x, y, t) = E(x + x_d, y + y_d, t + t_d)$

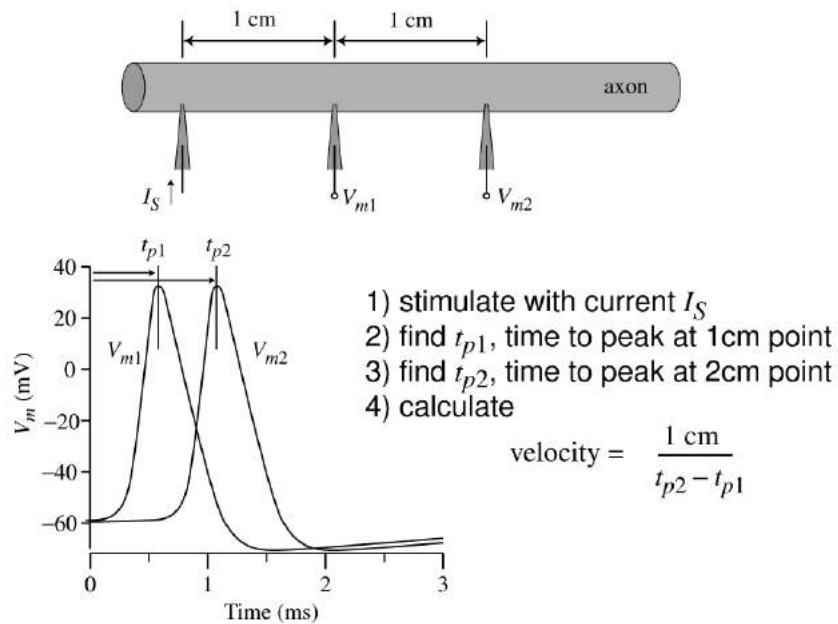
Expand in Taylor's series: $= E(x, y, t) + E_x x_d + E_y y_d + E_t t_d + \dots$

Set $t_d = 1$ frame: $E_x x_d + E_y y_d + E_t + \dots = 0$

Sampled version: $G(i, j, k)$ $G_x x_d + G_y y_d + G_t + \dots = 0$

Least squares: $(\hat{x}_d, \hat{y}_d) = \arg \min_{x'_d, y'_d} \sum_{i, j} (G_x x'_d + G_y y'_d + G_t + \dots)^2$

Methods: Calculating Velocity of Propagation



Results section describes the major findings of experiment.

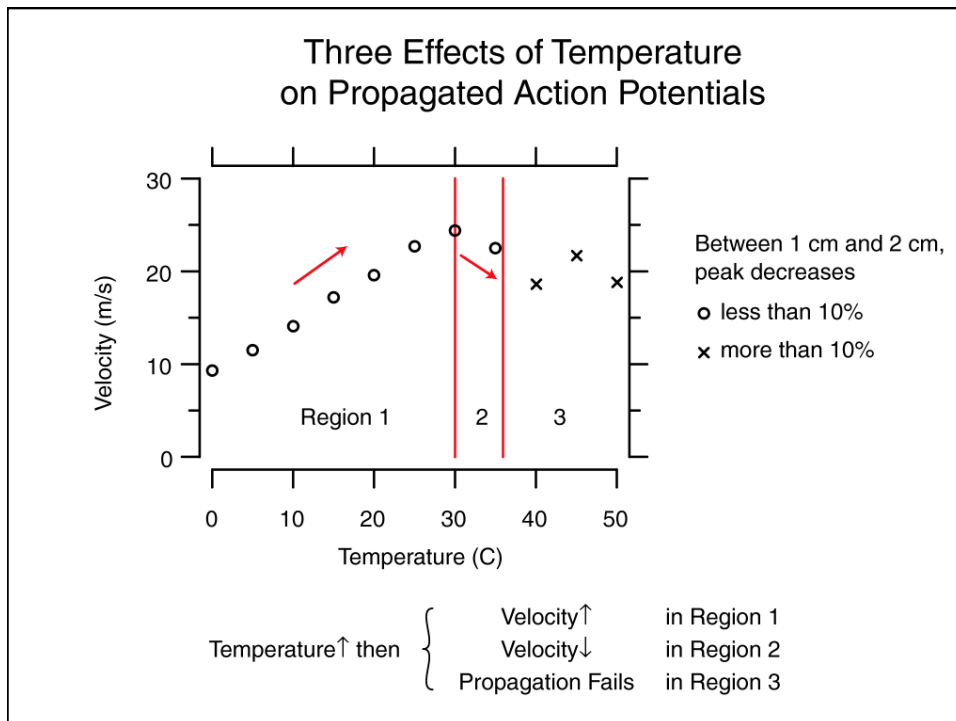
- Approximately 2-3 relevant figures
- Distill information about each figure into 2-3 bullet points
- Include key words in figures to remind yourself (and audience) of each bullet point



<http://www.nearingzero.net/index2.html>

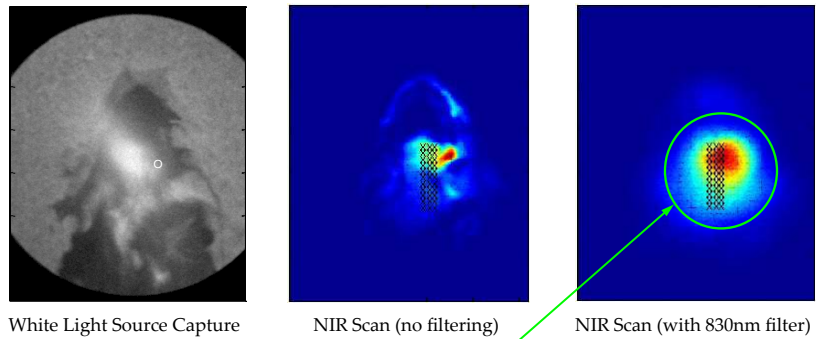
Results

- Velocity increased with temperature for range from 0 to 30 C.
- Velocity decreased with temperature for range from 30 to 35 C.
- Action potentials did not propagate for temperatures above 36 C.



Phantom Test – Reconstruction

Acquired 2D image of mouse specimen, showing accumulation of ICG in the head

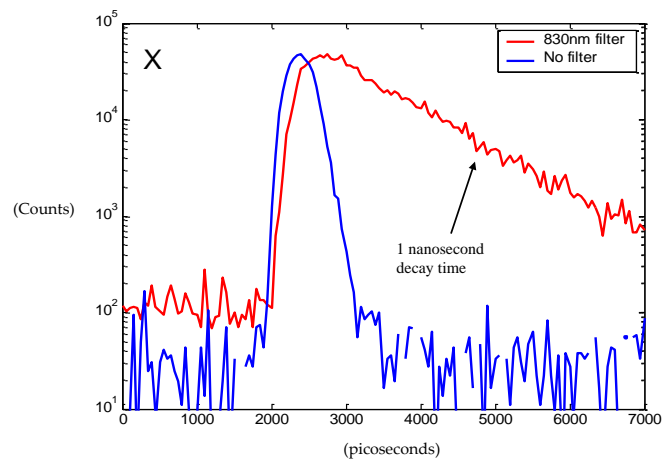


Fluorescence shows accumulation of ICG in brain/CSF

Courtesy of F. Hammond

Temporal Point Spread Function

TPSF of NIR pulse detection through mouse skull



Courtesy of F. Hammond

The Discussion offers your interpretations of your findings.



- Approximately 1-2 slides
- Explain **limitations**, questions left unanswered, major experimental constraints, lack of correlation, negative results. It's not all NOISE!
- How do your results **relate to the goals** of the study? (See Introduction)
- How do your results **relate to the background information** obtained in outside readings?

Summer 2005 Conclusions

Ductile Tissue Phantom

- Currently in calibration trials for TOBI at MGH

The future..

- Optically different phantoms: Truly mimic tissue
- Mechanically develop to be softer



Breast Spectroscopy System

- Developed and working: patient trials have begun!

The future...

- More patient trials and data analysis to capture breast tissue makeup

Courtesy of G. Lewis

Summary

- Created a library of decorrelation time constants for different tissue types:
 - Fatty-Fibrous, Glandular, Smooth Muscle, Fibrous
- Found that the rate of speckle decorrelation is dependent on tissue composition

Courtesy of A. Elliott

Future Directions

- Expand speckle library by increasing number of sampled tissues
- Utilize ability to characterize tissue using LSA to build a guidance probe with potential uses in needle biopsies

Courtesy of A. Elliott

Arrangement Step 2: Edit the Slide Show

- Edit slides for coherence
- Check for irrelevant bullets, plots.
- Check for balance and coherency in storyboard
- Spell-check and proofread

Style: Consider your speaker's persona

- Think in terms of talking to people
- Look at your audience
- Observe their reactions
- Adjust your style accordingly
- Make your enthusiasm for your work infectious
- Even shy people can be very effective public speakers!



Memory: Memorizing your talk word-for-word is not necessary



However, use notes, note cards, or a printed version of your PPT slides to keep yourself organized and on track.

Delivery: The rain in Spain falls mainly on the plane.

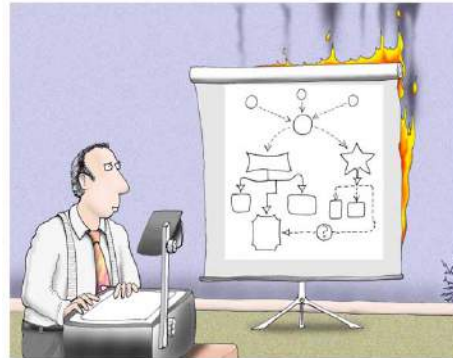


- Avoid filler words and distracting sounds (um, like, you know, okay).
- Don't hide from your audience.
- Talk to your audience not to the screen or poster.
- Don't fix your gaze on only one or two audience members.
- Don't read your visual aids verbatim.
- Practice, practice, practice.

The key to effective delivery is to practice!

- Stick to the time limit.
- Practice speaking slowly. Breathe.
- Work around your nervous habits.
- Use visuals as cues, not notes.
- Know how to use the equipment.

- Have a printed copy of your presentation for backup.
- If you get lost, stop and regroup. Your audience wants you to succeed.



Chapter 3. Andrew's scheme backfires.

<http://www.nearingzero.net/index2.html>

Prepare for Q&A

Big Picture

- Q&A is usually the best part of the talk!
- Usually questions are easier than you expect.
- Stay polite & composed. You're the expert.
- The moderator should police the audience.

Q&A Protocol

1. Invite questions from different parts of the room.
2. Listen to the ENTIRE question.
3. Make sure everyone hears the question
4. Identify the type of question.
5. Answer concisely.
6. End Q&A on good note.
7. Don't offer to answer via email unless you mean it.
8. Prepare backup slides.
9. Thank the audience.

What to do about bad questions

If . . .

Then . . .

A question contains many parts

Acknowledge many parts

- Answer 1 of the questions
- Answer more than 1 if time

There is no question

Spin into a question

The question is vague

“As I understand your question . . .”

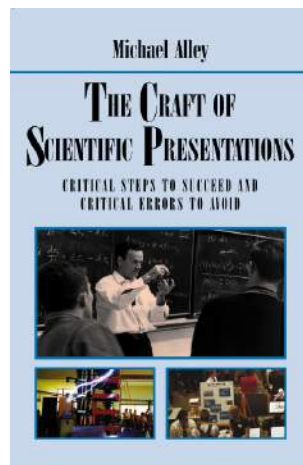
The question is combative

“I disagree.” + explain

“Thank you for your concerns.”

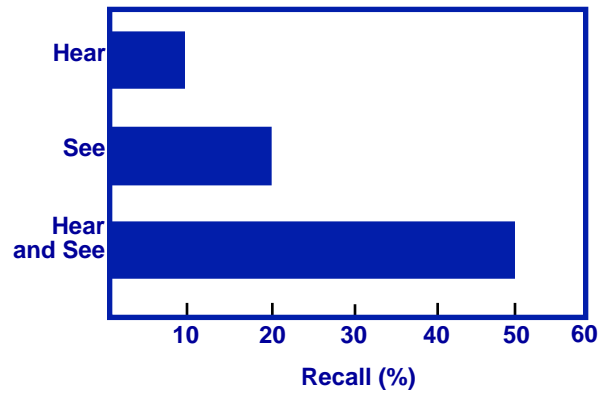
- “Perhaps we can talk after the Q&A”

Michael Alley has sensible advice on how to use PowerPoint.



<http://www.writing.engr.psu.edu/>

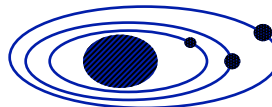
Audiences remember more when you use well-designed slides



Include slides that accent important details

Images

Neptune has three moons



Results

The world is warming

Four warmest years of century → 1988
1987
1983
1981



Exclude details that the audience does not need or cannot remember

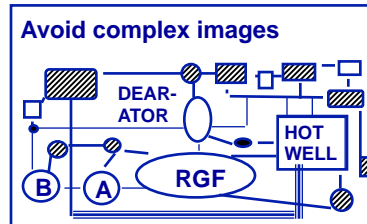
Avoid filler information

Roentgen discovered x-rays in 1895. He found that a cathode-ray tube produced fluorescence in a distant platinum-barium-cyanide screen.

Avoid long lists

- Corrosion
- Acid rain
- Toxic materials
- Pulsed combustion
- Energetic materials
- Pyrogenic materials
- Smog

Avoid complex images



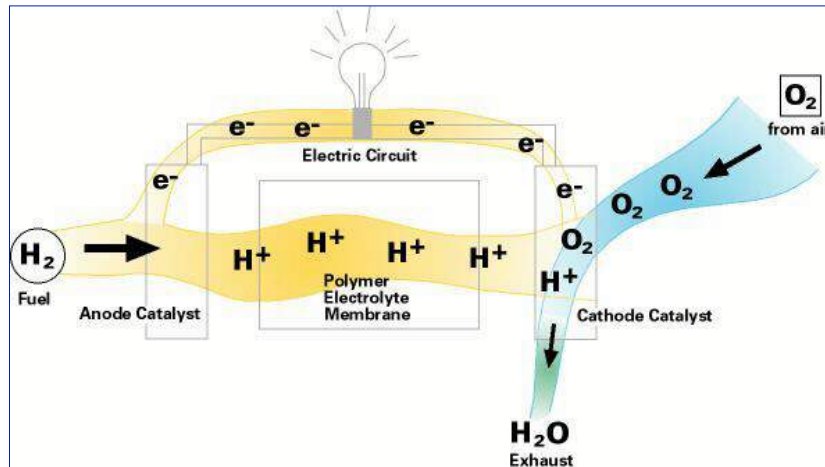
This sentence headline introduces the first topic (28 points, left justified, no more than two lines)

Image(s)
about
first topic

Subordinate point (keep point to no more than two lines)

Logo

Fuel cells are devices for energy conversion



[Breakthrough Technologies Institute/Fuel Cells 2000]



This sentence headline introduces the 2nd topic (28 points, left justified, no more than two lines)

First point (keep points to no more than two lines)

Second point (parallel in structure to the others)

Third point (parallel in structure to the others)



Logo

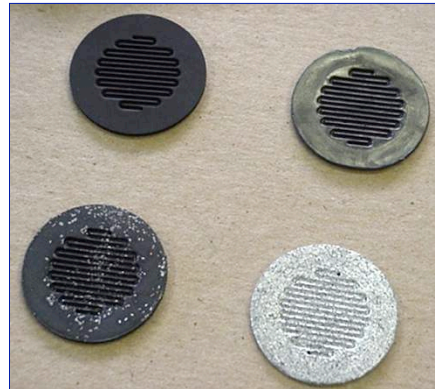
Composite materials are ideal for bipolar plates

Advantages

- Easy to shape
- Light in weight
- Resistant to corrosion

Disadvantages

- Low conductivity
- High cost (at present)



This sentence headline introduces the third topic (28 points, left justified, no more than two lines)

Image
about
third topic

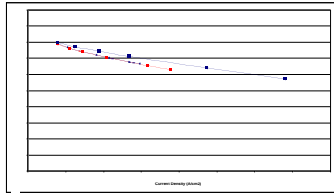
First point (keep points to no more than two lines)

Second point (parallel in structure to the others)

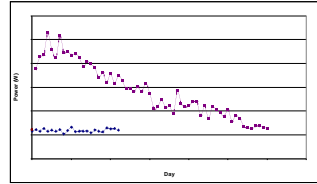
Third point (parallel in structure to the others)

Logo

Three methods exist for evaluating bipolar plates



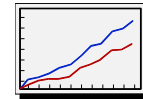
Polarization Curves



Power Curves



Visual Inspection



In summary,... (here, you state your most important conclusion of the work)

Supporting point (no more than two lines)

Another supporting point (parallel to the first)

A third supporting point (parallel to the first)

Image that supports conclusion

Questions?

Logo

**In summary, composite bipolar plates
show promise for fuel cells**

**Composite materials function well,
while under operating conditions**

Minimal corrosion was observed

**Conductivity difficulties need to be
addressed**



Questions?



An oral presentation to evaluate

[http://www.youtube.com/watch?
v=4lu9DVSWK4g](http://www.youtube.com/watch?v=4lu9DVSWK4g)

Oral presentation evaluation criteria

Communication—Presenter exhibits proper:

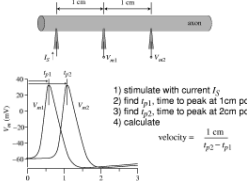
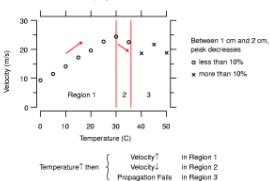
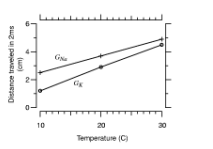
- eye contact
- voice
- gestures
- stance
- respect for viewers
- use of time

Content—Presenter exhibits:

- knowledge of subject
- concise explanation/summary of content
- effective answers to questions

An exercise

- From your one-page project description, create a six-slide storyboard of your presentation.
- Share your description and your storyboard with a colleague for feedback.

<p>Thermal Blockage of Propagated Action Potentials in the Hodgkin-Huxley Model</p> <p>Dennis M. Freeman</p> <p>H-H Project: 6.021J November 10, 2004</p>	<p>Introduction</p> <p>Question: Will action potentials propagate faster at higher temperatures?</p> <p>Pro: Rates of many chemical reactions increase with increasing temperature. Therefore it seems reasonable that the electro-chemical reactions underlying neural conduction would occur more rapidly at higher temperatures.</p> <p>Con: However, excessive heat leads to stroke, which represents profound neurological failure.</p> <p>Goal: Develop a mathematical model for effects of temperature based on the Hodgkin-Huxley model of neural excitation.</p>
<p>Methods: Calculating Velocity of Propagation</p>  <p>1) stimulate with current I_{stim}</p> <p>2) find t_{p1}, time to peak at 1cm point</p> <p>3) find t_{p2}, time to peak at 2cm point</p> <p>4) calculate</p> $\text{velocity} = \frac{1 \text{ cm}}{t_{p2} - t_{p1}}$	<p>Three Effects of Temperature on Propagated Action Potentials</p>  <p>Between 1 cm and 2 cm, peak decreases</p> <ul style="list-style-type: none"> o less than 10% x more than 10% <p>Temperature ↑ then { Velocity ↑ in Region 1 Velocity ↓ in Region 2 Propagation Fails in Region 3</p>
<p>Increasing Temperature Speeds Sodium and Potassium Conductances</p>  <p>... but rate of increase greater for potassium!</p>	<p>Summary</p> <ul style="list-style-type: none"> Increasing temperature increases velocity of propagation ... but only for a range of low temperatures. Increasing temperature above a critical temperature blocks the propagation of action potentials. Thermal block results because inactivation processes increase faster with temperature than do activation processes.

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Questions?