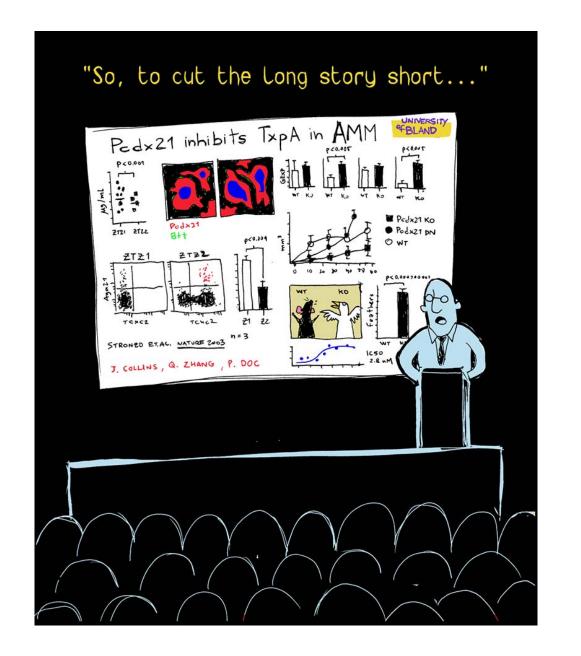
How to give a talk and how to be an active audience member



Overall organization (How to construct your talk)

INTRODUCTION

YOUR CONTRIBUTION

These two sections are equally important. Spend as much time working on your introduction as on your own work.

CONCLUSION

General Tips

(Things to think about as you construct your talk)

1) Keep your audience in mind

- Their knowledge
- Their familiarity with your field of study, your lab and your project

As much as you can, treat your audience like precautious 5 year olds. They are extremely intelligent, but ignorant.

(What is the difference between intelligence and ignorance?)

2) Divide your talk into sub-sections.

Have organizational slides.

3) One slide per minute.

If you are taking too long on one particular slide, add an extra slide!

4) Animations are okay, but don't over-animate (this is sometimes called Power Pointless!)

5) Use as few words, and as little text as possible in order to succinctly convey the information.

This takes practice and a lot of planning.

6) Deliver bite-size packs of information at a time.

This is a good place to use animations to help focus the audience's attention.

7) Walk through your slides

Guide your audience through each piece of relevant data. Every bar, cell ... every relevant data point! Remember that your audience is IGNORANT (but intelligent).

8) Practice with grad students that are not in your lab.

*Audience:

- Did you get it?
- Do you understand the problem/question being addressed?
- Are you missing the relevant background information?
- Are you confused and/or distracted by something in the presentation
- When critiquing the presenter:
 - Try to convey your ability to follow the reasoning behind the experiments
 - Don't try to give TOO MUCH ADVICE, you are not Napoleon! Your goal is to make the talk more intelligible!

How to Organize Each Section

<u>INTRODUCTION</u>: As important as your contribution.

You need to bring the audience up to speed. This is where you educate the precautious 5 year olds in the audience so that they can understand what is coming up in the section where you describe your contribution.

- -What is known?
- -What are the outstanding problems?
- -What is the relevance to human health, our general understanding of biology etc.?
- -Introduce questions that your work will go on to (or attempt to) address.

Be careful to bring up ALL relevant information, ideas, unorthodox techniques etc. Avoid giving IRELEVANT facts. You do not want to overload people with extraneous information.

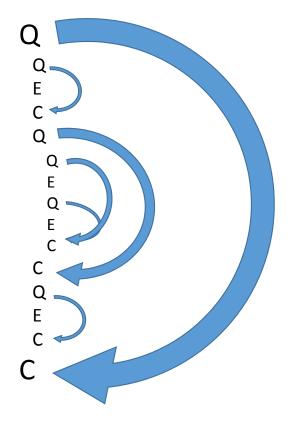
As you work on the "YOUR CONTRIBUTION" section, you should always ask yourself: did I provide the relevant information in the INTRODUCTION so that they can make sense of what is going on?

YOUR CONTRIBUTION

Start off with big question "Q"*
(ideally this flows naturally from your into)
Subdivide this question into smaller questions
How will you get an answer? (experiment, "E")
What did you learn? (conclusion, "C")

All questions need conclusions.

Sometimes many questions give rise to a single conclusion



"Is Rtn5 required to for cell viability?" can be replaced with "Rtn5 is Required for Cell Viability"

^{*} In a slide title, questions can be stated as conclusions

YOUR CONTRIBUTION

Present your work as a series of problems or questions.

Present each experiment as attempt to give an answer.

Dos:

- Show the ESSENTIAL data, including KEY controls (but don't show everything!!!!! Think of your audience.)
- Remember, an experiment is changing <u>one</u> variable and presenting an outcome emphasize the variable that you are changing Example: CONTROL TREATMENT vs. EXPERIMENTAL TREATMENT (e.g. WITH REVERSE TRANSCRIPTION vs. NO REVERSE TRANSCRIPTION)
- Use bullet points to help organize ideas and cut down on text
- Use large clear fonts (don't use comic sans)
- As much as possible, show schematics/cartoons instead of text
- Make the data intelligible without your input (titles, axis labels, panel labels)
- Show statistical significance if possible

YOUR CONTRIBUTION

Don'ts:

- Show extraneous data/experiments
- Fill the page with text or long sentences
- Present too much data on one slide
- Present distracting slides/graphics
- Show too much raw data in your talk (show summary tables/graphics when appropriate)
- Display info that is different from what you are saying (this distracts and confuses the audience)
- Use too many acronyms, abbreviations and jargon (explain all non-standard acronyms and abbreviations)

DON'T LOSE YOUR AUDIENCE!!

My Bullet Point-filled Slide

- Instead of dumping paragraphs of text on slides that no one can possibly read while still paying attention to the speaker, consider reducing points to three or four words max
- Then the presenter can fill in extra details verbally (the visible text summarizes)
- Or, even better, replace the text altogether with visual elements such as pictures

YOUR CONTRIBUTION - Data presentation (blots, microscopy, photos ...)

- Show high resolution images
- For microscopy, <u>try never to show exclusively color images!!!!!!!</u> (i.e. always use grey scale, with an additional color overlay being optional)
- Use the image as a tool to display your result
- (i.e. if you can't see the result clearly, alter the image or choose another image)
 - Make the image as big as possible
 - Crop the image to show what is essential
 - Make all your panels uniform
 - Increase the brightness/contrast if imaging data is key to your talk, show up early and test your slides on the screen
 - Label your images/panels
- Have quantitative data in hand (you don't necessarily have to show it but you may want to have it on an additional slide at the end of your talk)
- Do not alter the X/Y ratio of the image (do not show squashed or stretched images!!!!!!)
- Include statistical tests where necessary

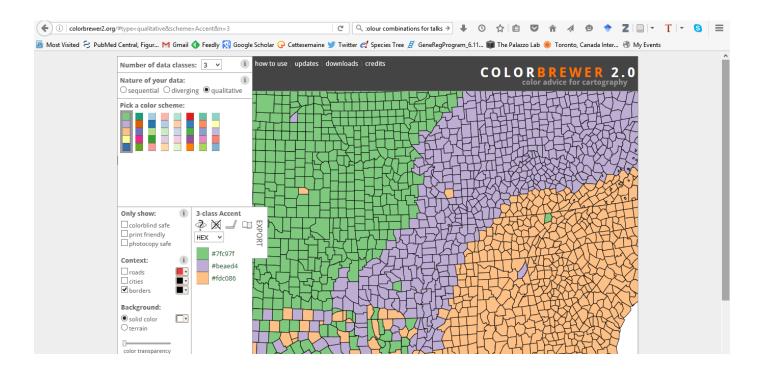
Order of priorities:

- 1) Convey information this is what your data needs to do!
- 2) Beauty if it looks nice, that's great, but this is not as important as point 1

The problem with colour:

- For immunofluorescence, your data is harder to see
- For figures and immunofluorescence you should also consider colour blindness

If you have to use color, choose the right combination: www.colorbrewer2.org



Other miscellaneous tips

- If you want to make sure that you say something that you may forget, show it in a small text box in your slide

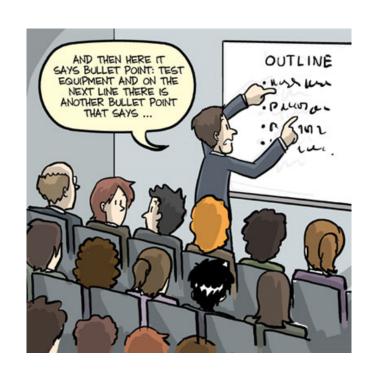
 Example: "We also confirmed this result by expressing [OTHER DOMINANT NEGATIVE PROTEIN]."
- If you show published data, include the citation somewhere in the slide

Example: Cui et al., PLoS Bio 2012

- If you show other people's data, acknowledge them in the slide (i.e. write their name at the bottom right corner)
- Don't be afraid to steal images from other labs' publications, but make sure to cite them in the slide
- State what organism/cell line you are using
- Emphasize key pieces of data with boxes or other graphics
- Use animations to display each bite-size piece of data at a time
- Add extra data at the end of your presentation. Although you wont show this data, you can flip to it during the Q&A.

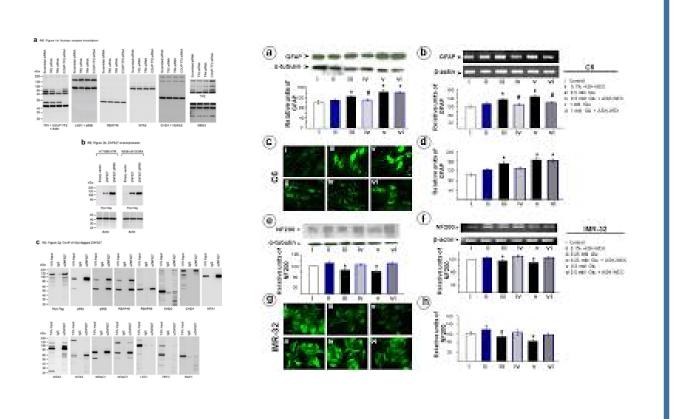
Tips for speaking.

- Do not read off of your slides
- Reduce the number of filler words (so, ah, uh, um, like, right, you know ...)
- Make eye contact with the audience
- Be engaged
- Try to explain, not recite
- Avoid sounding scripted
- Stay succinct (cut to the chase!)
- When using words like "it", "that", "this" etc. ... is it clear what you are referring to?
- Clearly point to each piece of data, do not whip your laser pointer around the slide!



Examples:

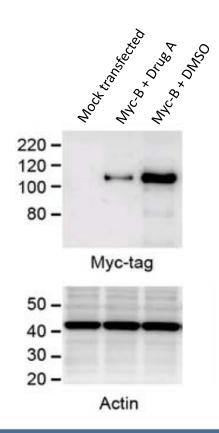
Look at my cool data!



- 1) Show the ESSENTIAL data, including KEY controls
- 2) Present bite-size data



Drug A reduces expression of protein B



Speaker:

We wondered whether drug A affected protein B. So, we took cells and we transfected them. We then ran the lysate on a gel and blotted it for myc. The drug affected protein B [points to lane 2 of the top gel]. If we then looked at actin levels [points to the second blot], untransfected cells had as much protein as cells treated with drug A or with cells treated with control medium. So, um, from this experiment we concluded that Drug A affected the levels of protein B, just like our hypothesis stated.

"Running" a gel is slang, not the worst offence. but should be kept to a minimum.

This is too vague. Use the active voice.

How was it affected?

What cell type?

With what?

Get rid of this

How was the protein

Control? Is that lane 1

What is labeled with myc? Speaker:

affected?

or 3???

We wondered whether protein B was affected by drug A. So, we took cells and we transfected them. We then ran the lysate on a gel and blotted it for myc. The drug affected protein B [points to lane 2 of the top gel]. If we then looked at actin levels [points to the second blot], untransfected cells had as much protein as cells treated with drug A or with cells treated with control medium. So, um, from this experiment we concluded that Drug A affected the levels of protein B, just like our hypothesis stated.

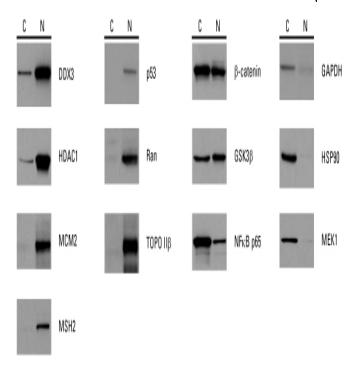
Too wordy! This is just a loading control

Get rid of this

Unnecessary

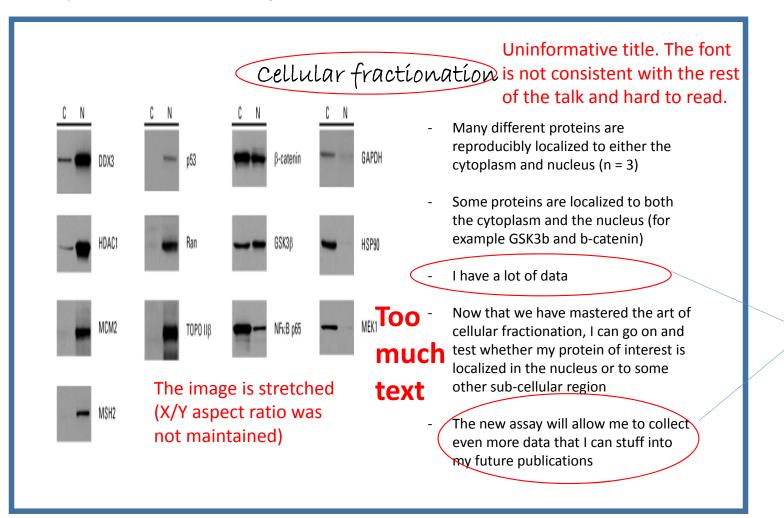
To determine whether drug A affects the expression of protein B, we transfect a myc-tagged version of this protein in COS-7 cells that were treated with drug [points to lane 2] or vehicle [points to lane 3]. As you can see, Drug A inhibits production of myc-tagged B. No signal was seen in untransfected cells [points to lane 1]. Actin was used a s a loading control [points to the second blot].

Cellular fractionation



- Many different proteins are reproducibly localized to either the cytoplasm and nucleus (n = 3)
- Some proteins are localized to both the cytoplasm and the nucleus (for example GSK3b and b-catenin)
- I have a lot of data
- Now that we have mastered the art of cellular fractionation, I can go on and test whether my protein of interest is localized in the nucleus or to some other sub-cellular region
- The new assay will allow me to collect even more data that I can stuff into my future publications

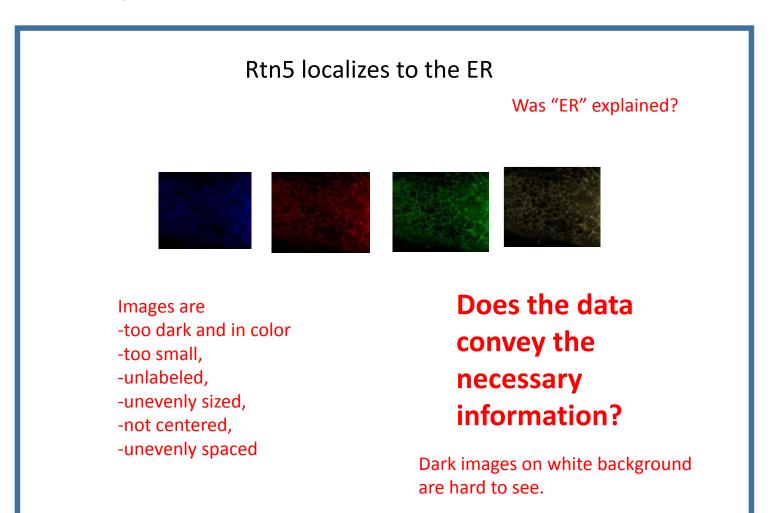
- 1) Don't fill the page with text or long sentences
- 2) Do not show squashed or stretched images

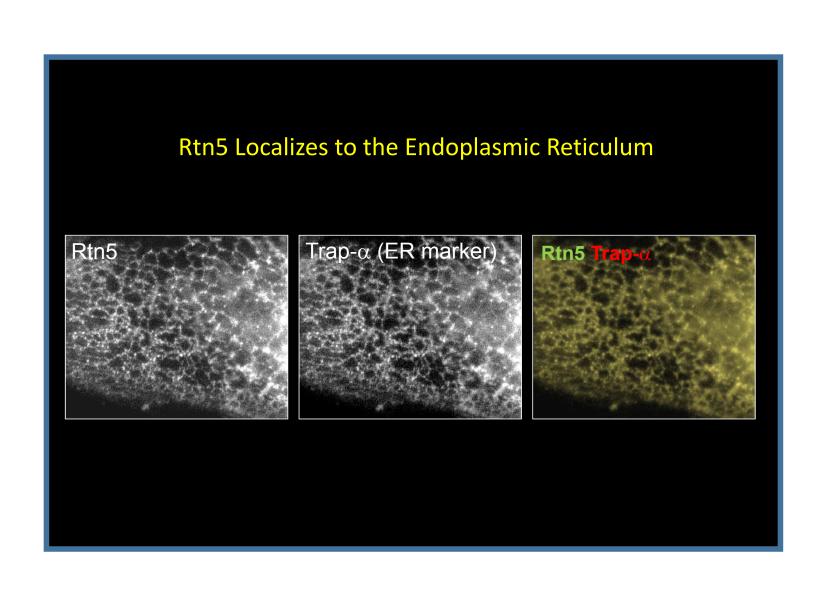


Irrelevant information

Rtn5 localizes to the ER

1) The data does not convey the information.





Last part:

CONCLUSION

- Short (NOT TOO WORDY!)
- Summarizes the most salient points.
 - What is the take home message?
- Should refer back to your introduction
 - How does your work add to our knowledge?
 - Relevance to human health, everyday life ...

Future Directions (can come before or after the Conclusion)

- -Again, use as few words as possible
- -Only state major questions/experiments. The audience does not want to hear a long litany of things to do.

(As a thinking exercise, but not necessarily stated in your talk, you should ask whether these experiments <u>help to address **major** questions</u> raised in the intro or by your work.)

Thankyou slide:

This can come at the beginning or end of the talk.

- Lab mates
- Collaborators
- Others who have helped you
- Funding agencies

This is the one slide that you don't have to worry about being too wordy or crammed. People will get offended if you don't acknowledge them.