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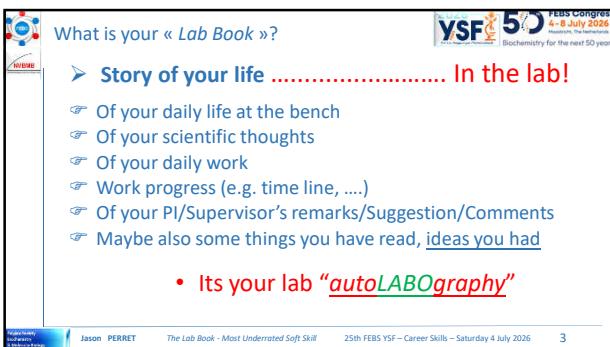
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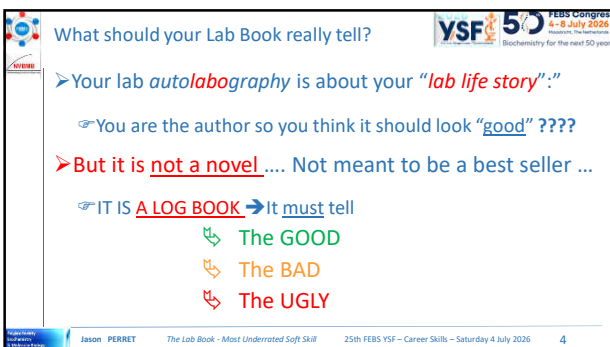
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The Bottom Line?

Its the *Bad* & the *Ugly*  
That are going to make you do the  
Good science

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So what is your Lab Book's purpose?

- Follow up of experiments => Experimental Flow => Research Finality
- Traceability of Experiment and Sample
- Results & Interpretations
- Scientific Integrity
- Proof of experiments and results and dates
- Contribute to the labs group research advancement - Discussions
- Write up your articles
- Write up your thesis

WHYYYYYYY ????

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BECAUSEEEEEEEEEEEEEEEEE !!!!!

Do you really think  
that you will remember tomorrow  
what you did today?

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What should be in your Lab Book?

- Planning**
  - Purpose/Goal/Question Asked of the Experiment(s)
  - What you plan to do
  - How you will do it
  - What you will do
- Action**
  - What you actually did
- Outcome**
  - What were the outcomes – RESULTS
  - What it means to you – the interpretation
  - What to do next

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What you actually do????!!!

- This is where The **Bad & the Ugly** ..... and also the **Good** are consigned
- Here is where you describe what you do

everything you do - everything that happens

- The **changes** made (to protocols or during the experimental procedures)
- The **temperatures** (what is a "room temperature"??)
- The **timing** – incubation times (what is an "overnight incubation"??)
- OBSERVATIONS MADE** e.g.: un expected change in color or transparency, a precipitation, a drop lost, a drop to much, no sure of exact volume, mistake in timing, temperature, sample,...

**WHYYYYYYYYY ????????**

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**BECAUSEEEEEEEEEEEEEEE!!!!**

Murphy's Law is especially true in the lab

**"If anything can go wrong it will !!!"**

*Luckily "Richard Zeckhauser" said:*

*"Sometimes things that should not work, work nevertheless"*

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What should your Lab Book allow you to do?

- Discuss results with your supervisor or PI – at Lab Meeting
  - ☞ Plan next experiment(s)
  - ☞ Know what was done
  - ☞ How it was done
  - ☞ Results generated
  - ☞ Communicate be able to describe it to colleagues and PI/Supervisor
  - ☞ Write it up
- **AND** ..... →  
 AND ..... →  
 AND ..... →

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Your Lab Book should allow you and others

- ☞ **TO ANSWER QUESTIONS KNOWLEDGEABLY!!!**
- ☞ **And always allow others to:**
  - ✓ Understand what was done
  - ✓ To Reproduce and/or Pick Up where you left off when you leave the lab for new adventures!!

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ANSWER THE QUESTIONS KNOWLEDGEABLY ????

Let us look at a typical scenario!

PI: "How much antibody did you use on your blot? Or PCR product did you put on gel? Or the amount of primers used (final concentration)? Or antibiotic used for selection? Or was the serum heat inactivated?"

The Young & Bold & Growing Scientist: "20 µl"

PI: That does not mean anything!!! 20 µl of what? What was the dilution of your antibody or volume of your PCR reaction or the concentration of your primer stock and dilution factor? ....."

The Young & Bold & Growing Scientist: looks in her/his lab book - Oups "Euh .. I didn't write that down".

**It was last week and she/he does not remember .... Of course**

PI: 🤔 or maybe even 🤡 because it is lost experiment – it is **YOUR** time & PIs money 🤡

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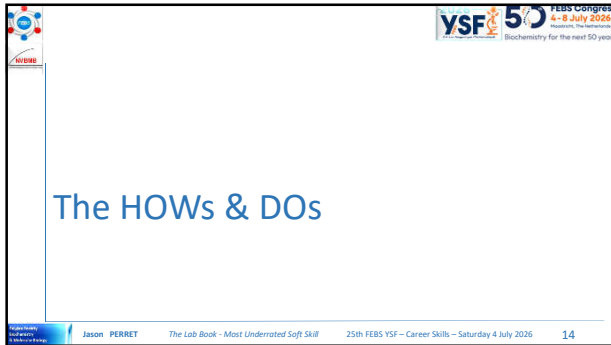
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The HOWs & DOs

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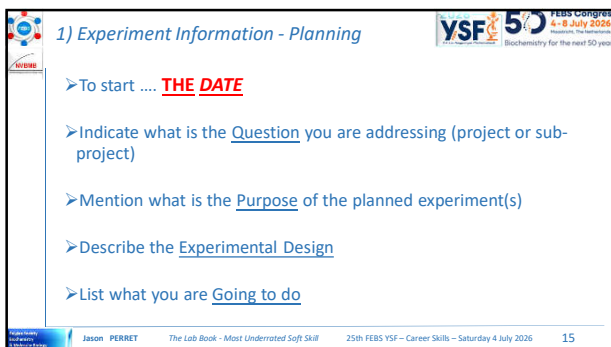
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1) *Experiment Information - Planning*

- To start .... **THE DATE**
- Indicate what is the Question you are addressing (project or sub-project)
- Mention what is the Purpose of the planned experiment(s)
- Describe the Experimental Design
- List what you are Going to do

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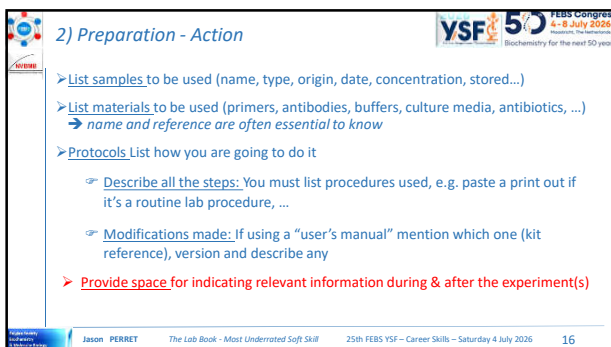
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2) *Preparation - Action*

- List samples to be used (name, type, origin, date, concentration, stored...)
- List materials to be used (primers, antibodies, buffers, culture media, antibiotics, ...)
  - ➔ *name and reference are often essential to know*
- Protocols List how you are going to do it
  - ☞ Describe all the steps: You must list procedures used, e.g. paste a print out if it's a routine lab procedure, ...
  - ☞ Modifications made: If using a "user's manual" mention which one (kit reference), version and describe any
- **Provide space for indicating relevant information during & after the experiment(s)**

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**3) Experiment Outcome**

> Readouts - Results – Comments & INTERPRETATION(s)

- ☞ Copy of
  - ✓ Tables
  - ✓ Picture/Images - if it is a digital file indicate file names(s) and location(s)
  - ✓ Instrument printouts
- ☞ Remarks, Comments, Conclusions for the given experiment(s)
- ☞ **ANNOTATE PROPERLY ALL IMAGES/INSTRUMENT PRINTOUTS**
- ☞ If problems and/or inconsistencies arise point out & indicate possible reasons
- ☞ **Interpretation(s)**
  - ✓ Indicate the follow-up, i.e. what next?
  - ✓ Next steps or controls to do or experiments for problem solving .....

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**BOTTOM LINE – TAKE HOME MESSAGE**

**WHAT DOES YOUR LAB BOOK LOOK LIKE??**

**DOES YOURS ALLOW YOU TO:**

- ☞ Understand what the question(s) is/are addressed.
- ☞ Redo the experiment = Understand what was done, how it was done, and exactly what materials/reagents were used.
- ☞ Interpret the results and answer the question(s).
- ☞ Next step(s)

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**Thanks for your Attention**



Darwin's "Tree of Life" sketch, from his Lab notebook.  
The Text starts with:  
**"I think ...."**

**Do you???**

From: <https://betterscienceteaching.com/2013/04/18/famous-science-notebooks/>

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**CHECK OUT SOME EXAMPLES**

**AND NOW SEE WHAT YOU « THINK »  
WHAT DOES YOUR LAB BOOK LOOK LIKE??**

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**EXAMPLE 1 - WB page 1**

**Very Good! A date**

**Good! Experiment Purpose**

**Good! Clear Table with conc. & volumes a reagents used**

**Good! Experiment details**

**Acronyms used not consistent between table and gel loading chart**

**Lacking electrophoresis conditions**

**Good! Gel Loading details**

**Good! Cross gel control used & described**

**Good! Detail on how blots were processed**

**Good! Experiment Title for continuity**

**Washing & Blocking volumes missing**

**No info on ab brand and cat no**

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**EXAMPLE 1 - WB page 2**

**Date missing on this page**

**Good! Process detailed**

**Bad! Wash/block volumes missing**

**Good! Process nicely detailed**

**Good! Images annotated for samples/conditions**

**GENERALLY NO COMMENTS/REMARKS RELATED GEL PHOTOS**

**Good! Ab details + brand**

**Bad! Cat no missing**

**Lab Book should always have a scan of the WHOLE GEL image. Don't be broken down alongside for analysis.**

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**Example 3 - Western Blot page 2**

**DATE missing.**

**GOOD** but could have been in the previous table to have an overview of reaction mixture and tick off as added

**Transferred how? Method?**

**Is this a typical Lab protocol???**

**GOOD!** Ab dilutions and volume stock used in the 2 ml binding solution...

**BAD!** Brand, cat#, species specificity, host species, ...

**Who is/what is "Mac" ?????? No way of reproducing experiment with this information**

**What marker?, Brand, cat#, eventually a copy of the molecular weights and distribution in the same type of gel.**

**What type of gel, %, gradient, denaturing, commercial, ...**

**Does not mean a thing, useless information, does not explain how and what was done!**

**What is "overnight" ?? 12, 14, 16, 18 or 20 hours???**

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**Example 3 - Western Blot pages 3/4**

**GOOD!** A Date

**Ab info: Species, brand, cat# ... There are millions of commercial HSP abs**

**Detection reagent info: Brand, cat#, protocol (amounts, incubation times, ...??)**

**GOOD!** Blot image annotated and MW as well!

**Is this the 1<sup>st</sup> or 10<sup>th</sup> exposure? Expected MW of the specific band?**

**IN GENERAL: NO COMING UP WITH MARKERS RELATED TO GEL PHOTOS**

**DATE missing.**

**GOOD!** Blot image annotated and MW as well!

**No explanations on what should be seen, what is seen, interpretation, meaning ... what next!**

**No explanation concerning IMP3 ... Was this to be expected? Unexpected results? Then some explanation/hypothesis to why. What workaround??**

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**Example 4 - Inhibition of CDK8 in CDK8As cell line page 1**

**Title a bit short to describe experiment purpose.**

**Good. Experimental conditions, well layout, concentrations, volumes**

**Good. Next steps**

**BAD!** Date??

**What cell line (passage number, confluency, culture conditions, ...)**

**GOOD!** Reagent addition, timing

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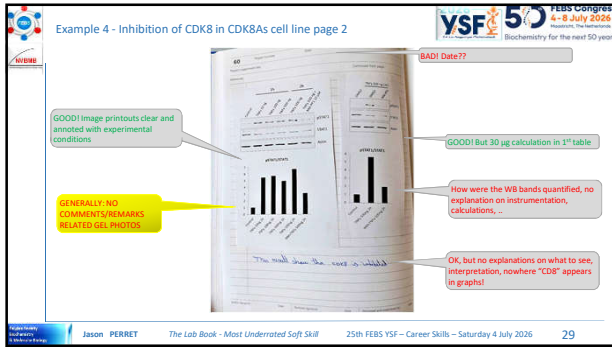
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Example 4 - Inhibition of CDK8 in CDK8As cell line page 2



Annotations on the slide:

- GOOD! Image printouts clear and annotated with experimental conditions
- BAD! Date??
- GOOD! But 30 µg calculation in 1<sup>st</sup> table
- How were the WB bands quantified, no explanation on instrumentation, calculations, ...
- OK, but no explanations on what to see, interpretation, nowhere "CD8" appears in graphed
- GENERALLY NO COMMENTS/REMARKS RELATED GEL PHOTOS

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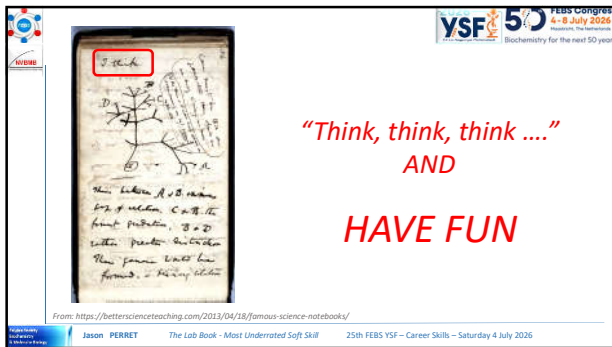
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“Think, think, think ....”  
AND  
HAVE FUN

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