

M2D6: Utilize cellular thermal shift assay (CETSA) to test putative small molecule binders

1. Prelab Discussion
2. Treat cells with small molecule ligands
3. Run SDS-PAGE
4. Protein gel transfer for Western Blot

Western Blot Protocol



Sample extraction



Run Gel



Transfer proteins to membrane



Antibodies incubations and washes



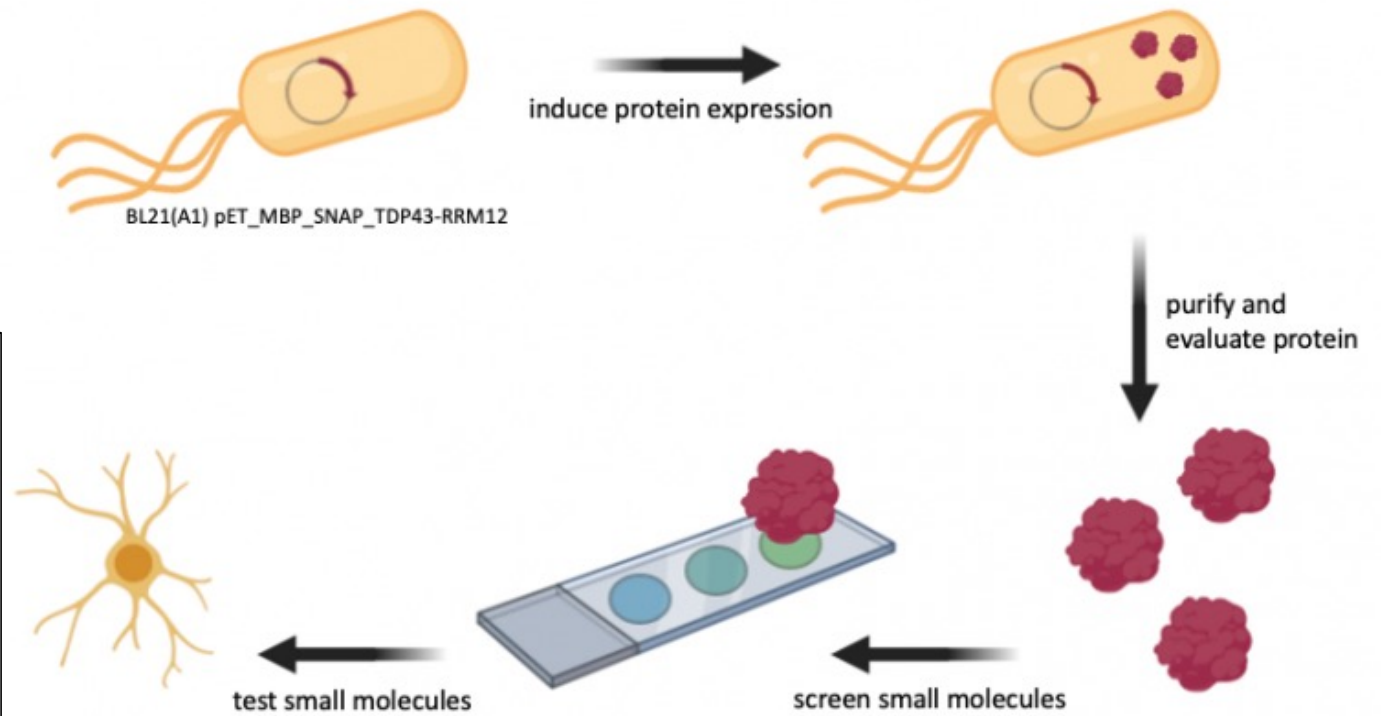
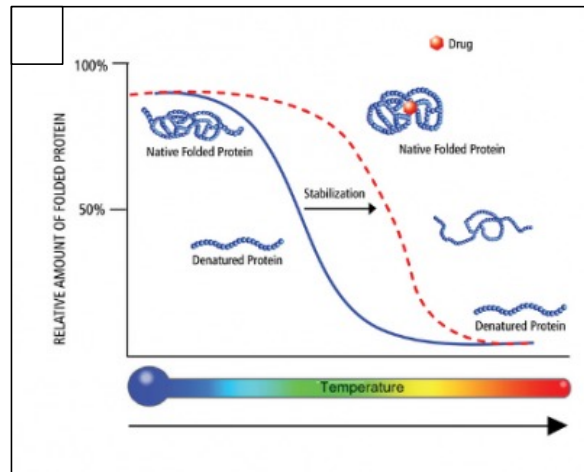
Fail to visualize protein bands



Cry

Pedromics, Science cartoons by Pedro Velica

Mod2 Overview



Cellular thermal shift assay (CETSA)

ON THE MOD2 OVERVIEW PAGE:

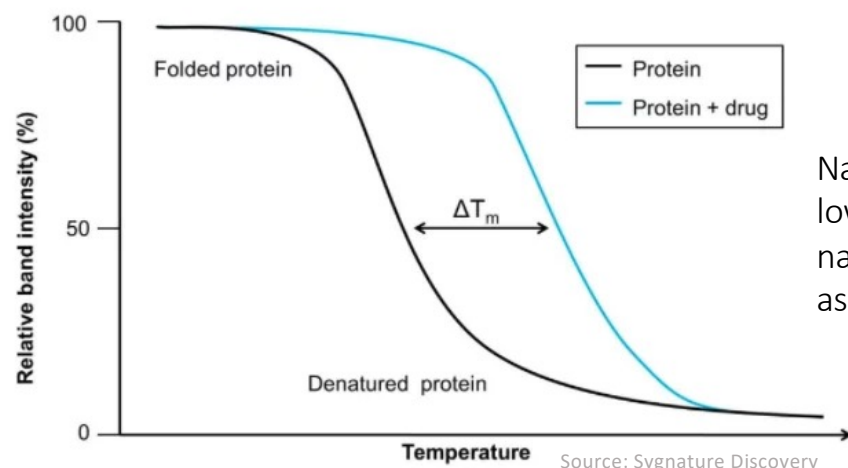
PROTOCOL

The cellular thermal shift assay for evaluating drug target interactions in cells

Rozbeh Jafari¹, Helena Almqvist², Hanna Axelsson², Marina Ignatushchenko¹, Thomas Lundbäck², Pär Nordlund¹ & Daniel Martinez Molina¹

¹Department of Medical Biochemistry and Biophysics, Division of Biophysics, Karolinska Institutet, Stockholm, Sweden. ²Chemical Biology Consortium Sweden, Science for Life Laboratory Stockholm, Division of Translational Medicine and Chemical Biology, Department of Medical Biochemistry and Biophysics, Karolinska Institutet, Solna, Sweden. Correspondence should be addressed to T.L. (thomas.lundback@ki.se), P.N. (par.nordlund@ki.se) or D.M.M. (daniel.martinez.molina@ki.se).

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- The ΔT_m indicates protein stabilization / destabilization compared to control
- Assesses thermal stabilization of protein in presence / absence of drug (small molecule) in the cell

Cellular thermal shift assay (CETSA)

PROTOCOL

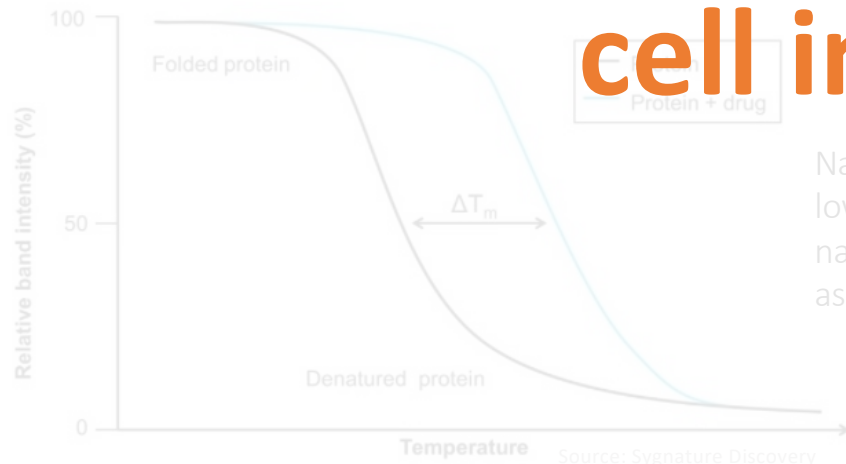
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¹Department of Medical Biochemistry and Biophysics, Division of Biophysics, Karolinska Institutet, Stockholm, Sweden. ²Chemical Biology Consortium Sweden, Science for Life Laboratory Stockholm, Division of Translational Medicine and Chemical Biology, Department of Medical Biochemistry and Biophysics, Karolinska Institutet, Solna, Sweden. Correspondence should be addressed to T.L. torbjorn.lundqvist@ki.se or D.M. daniel.martinez.molina@ki.se

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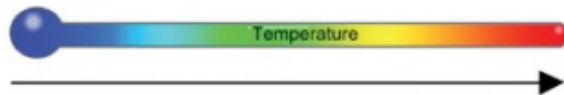
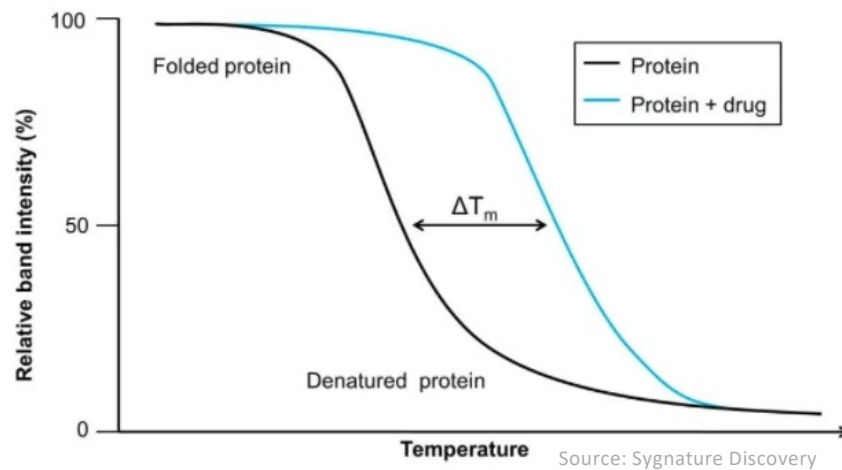
Why is testing for ligand binding in a cell important?



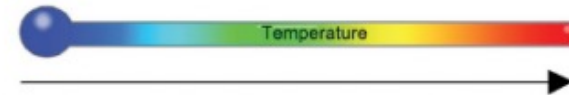
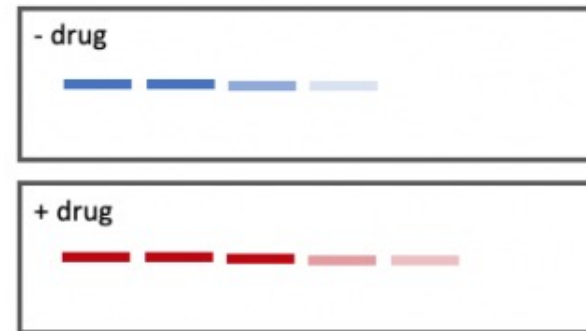
Native proteins denature at lower temperature than native folded proteins associated with drug

- The ΔT_m indicates protein stabilization / destabilization compared to control
- Assesses thermal stabilization of protein in presence / absence of ligand in the cell

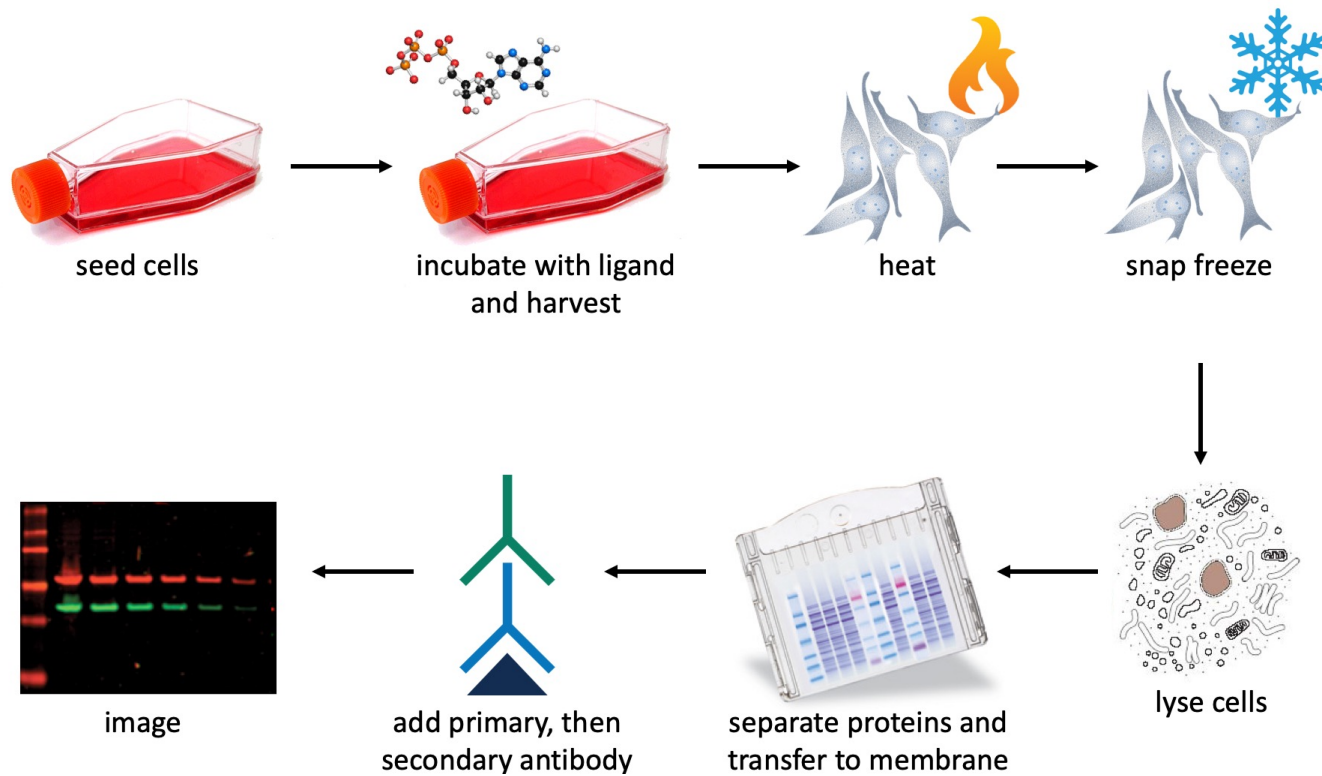
CETSA : Protein stability is measured by Western blot



Identify presence of native folded proteins

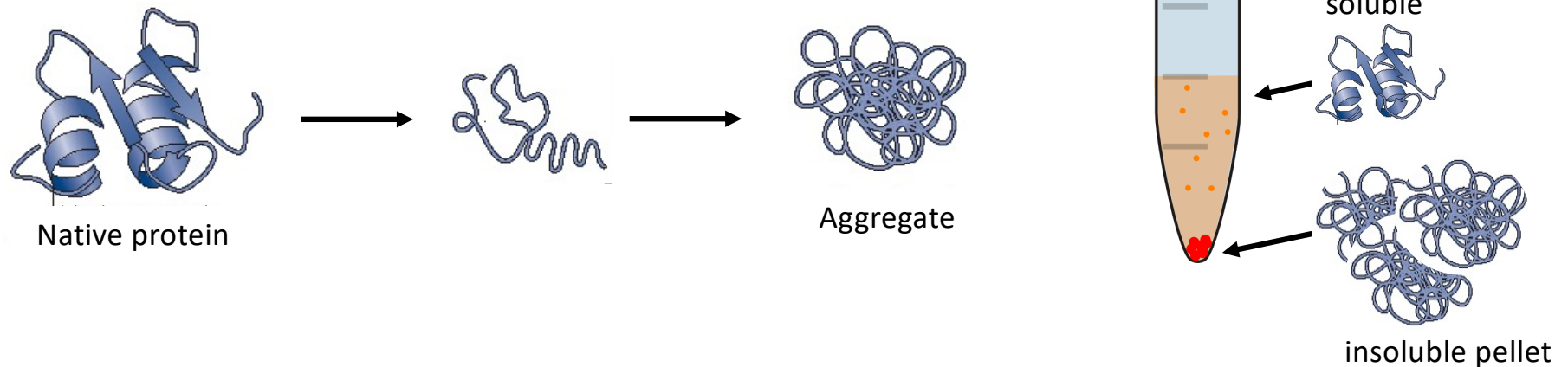


Major experimental steps of CETSA



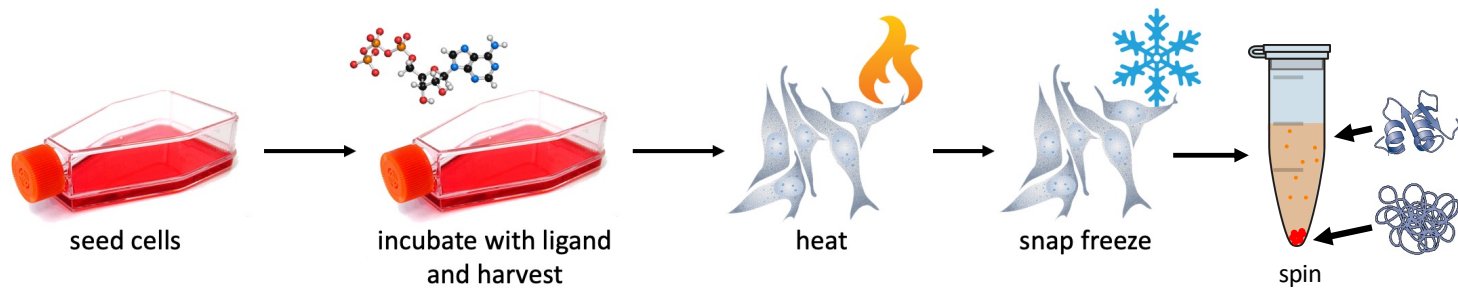
Heat causes protein denaturation

- As proteins denature, 'melted' primary structures aggregate
- Aggregates precipitate out of solution and can be removed via centrifugation

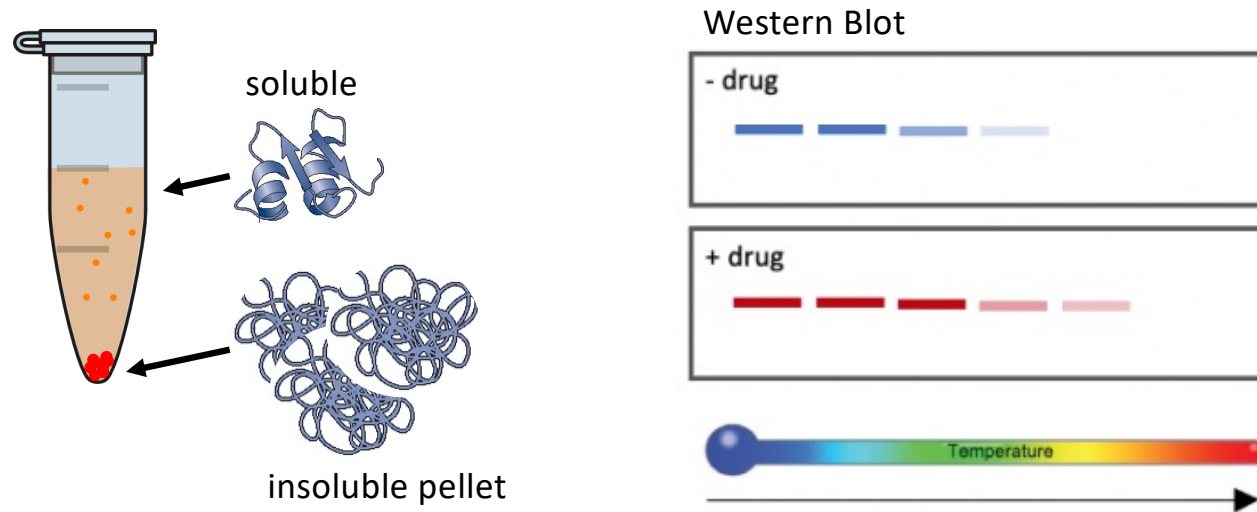


Purpose of CETSA steps

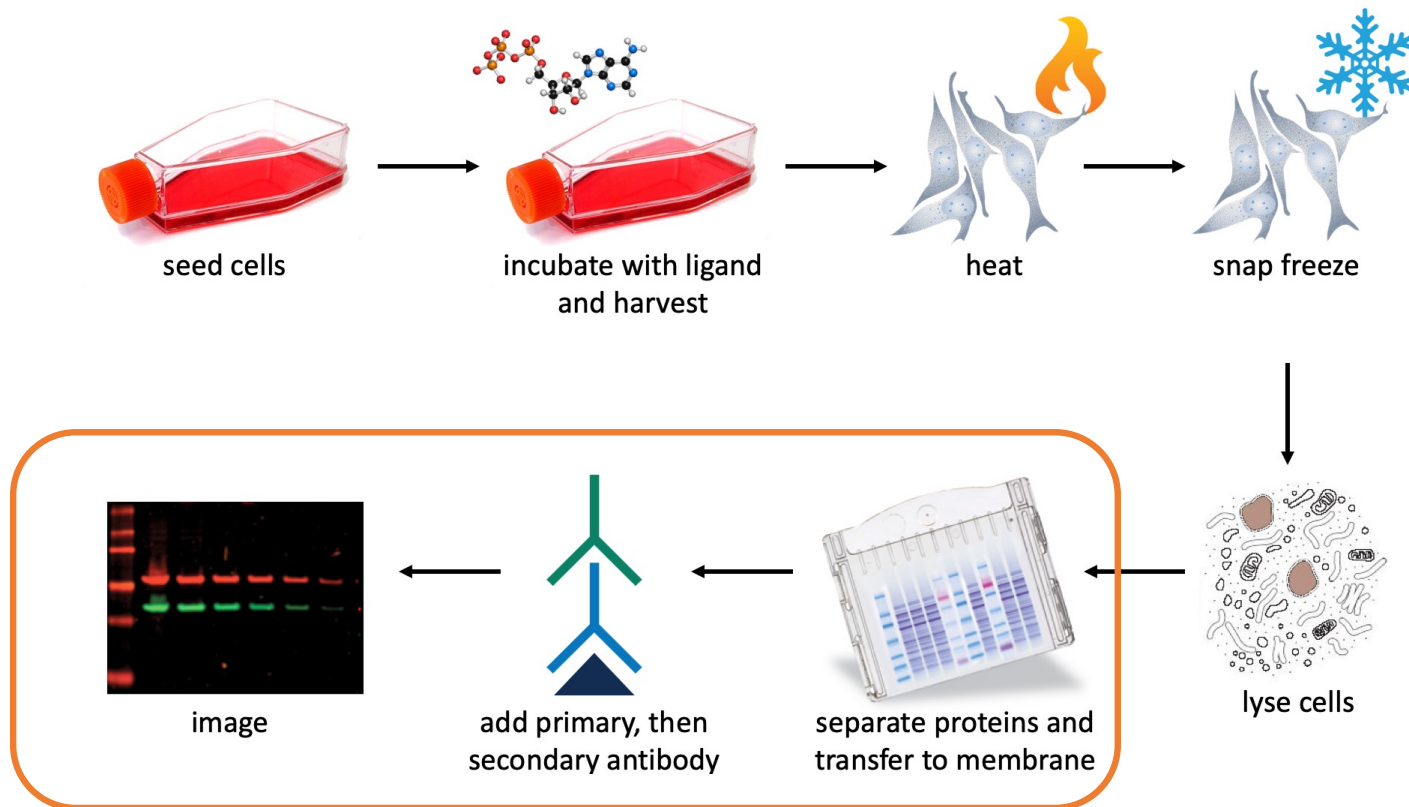
- Heat:
- Snap freeze:
- Spin:



Only soluble proteins are measured by Western blot

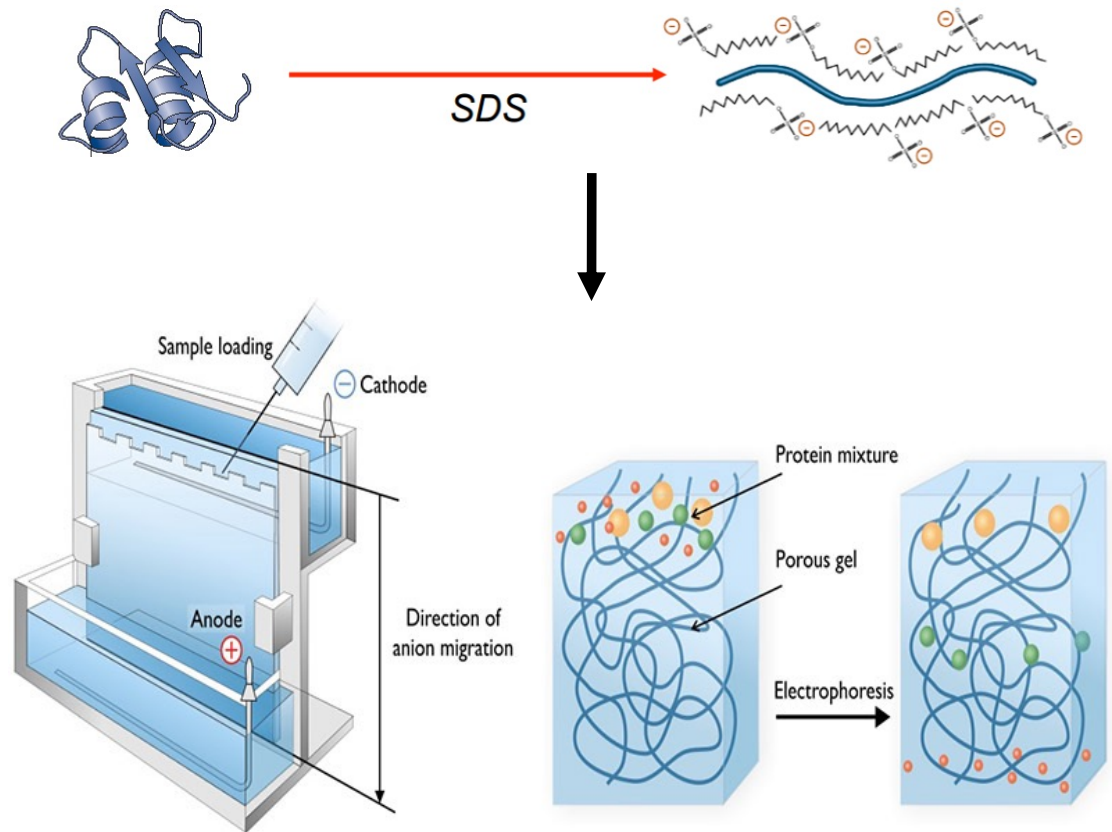


Imaging for CETSA



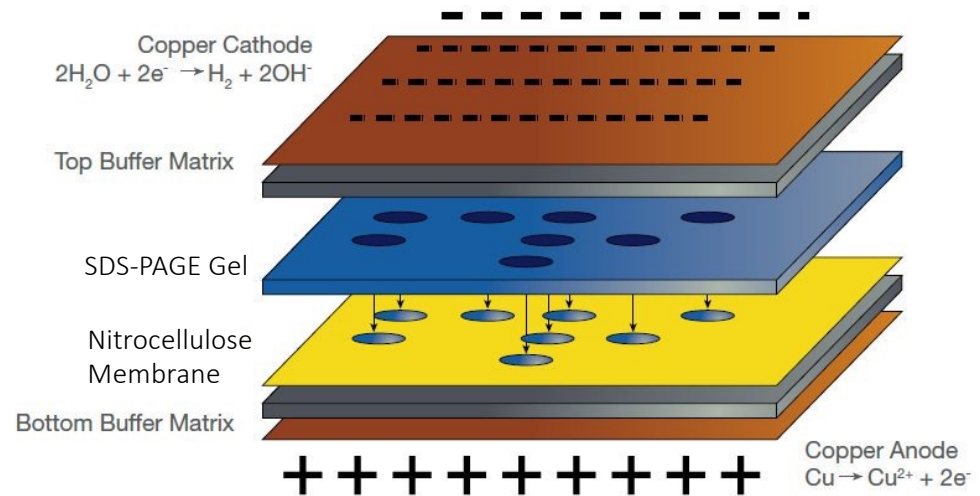
Separate proteins using SDS-PAGE (Review of M2D2 prelab)

- SDS-PAGE separates proteins according to
 - Charge and secondary structure are alleviated by:
- SDS imparts uniform charge-to-mass ratio



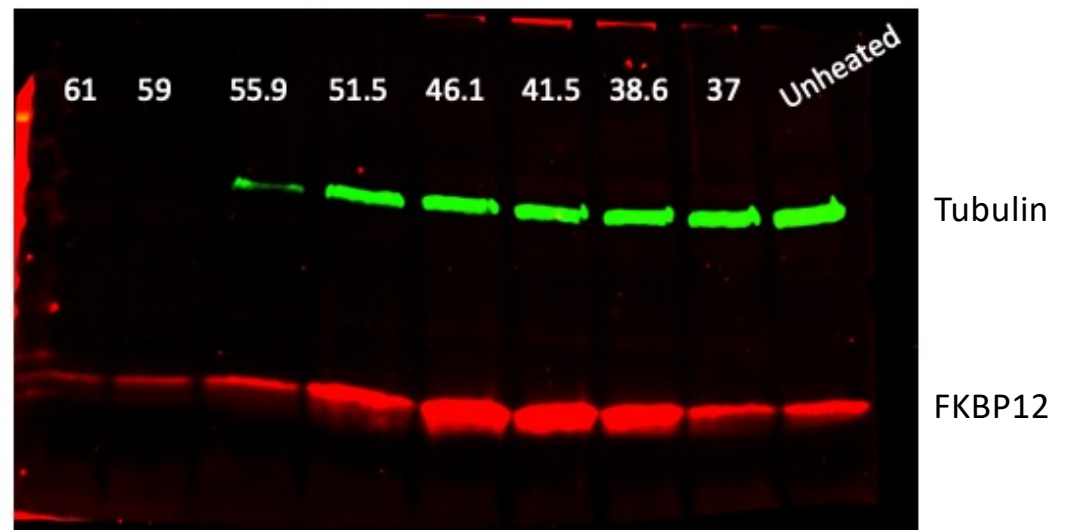
Transfer of proteins from SDS-PAGE gel to Nitrocellulose membrane

- Allows for protein identification with detection antibodies
- Move protein bands from polyacrylamide gel to a nitrocellulose membrane for further assessment
- Net negatively charged proteins are migrated using a current from the gel to a membrane



Proteins are visualized through Western Blot

- Membrane with proteins is 'blotted' using antibodies to probe for specific for protein of interest



For today

- Work through wiki to choose small molecule ligands and perform CETSA experiment

For M2D7 (one week from today, 4/23)...

- Write an outline of the research article discussion
 - Use citations
 - Propose 2 follow up experiments
- See wiki for additional guidance