

Slides by Carolina Barra Quaglia Presented by Bent Petersen

# Introduction to Protein Structure

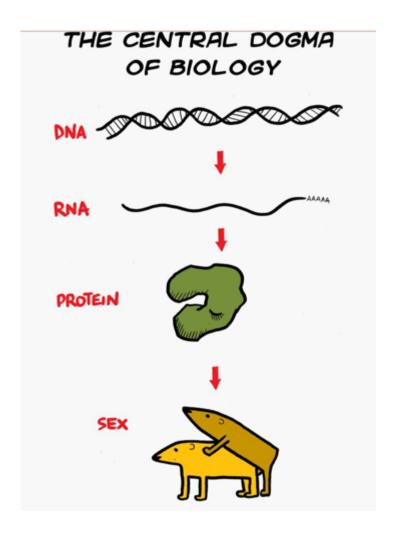


#### **Learning Objectives**

- Identify the four different levels of protein structure
- Discuss experimental assays to obtain protein structures: X-ray crystallography
- Identify protein structure databases
- Interpret relevant parameters for **evaluating the quality** of protein structure determined by X-ray crystallography
- Visualize and manipulate protein structures using PyMol

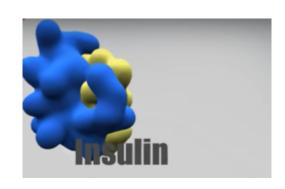


#### Why are proteins so interesting?

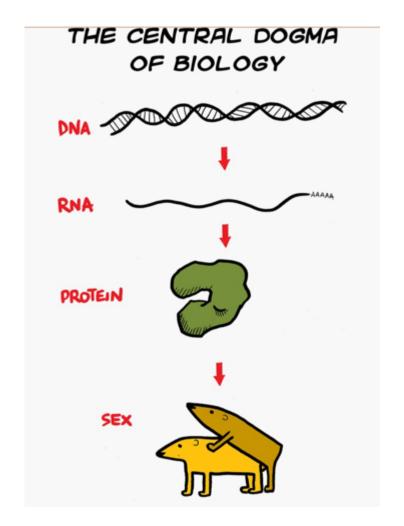


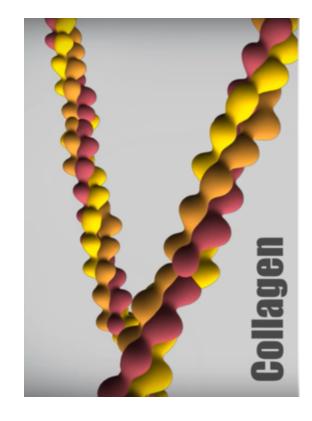


#### Why are proteins so interesting?



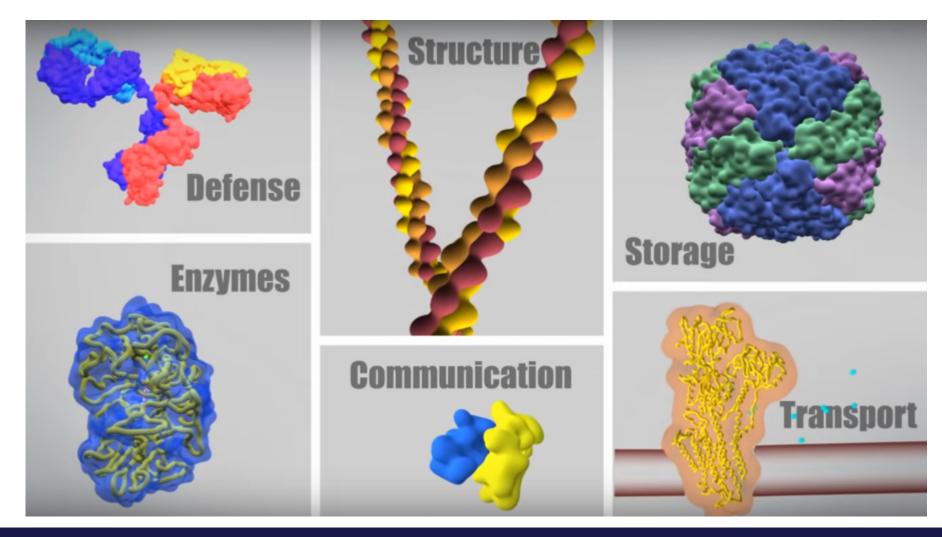






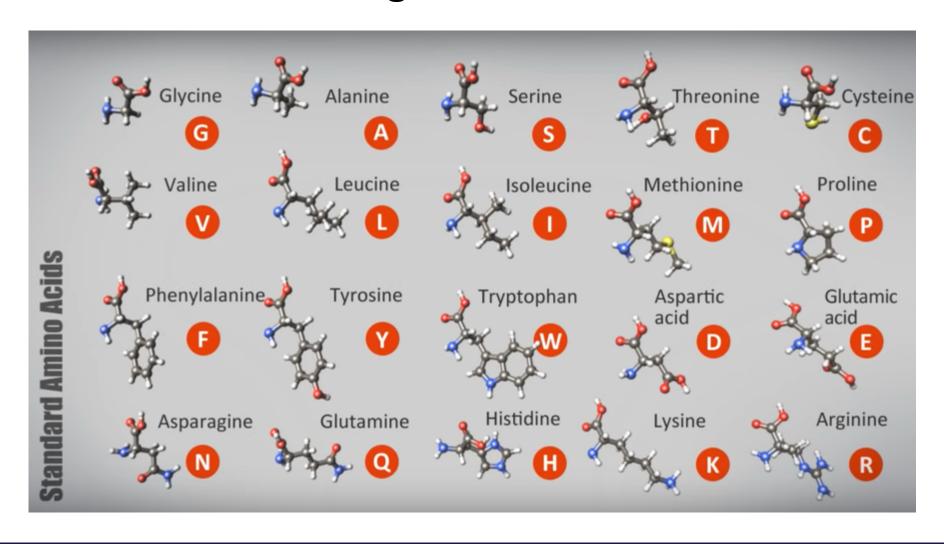


#### Why are proteins so interesting? FUNCTION



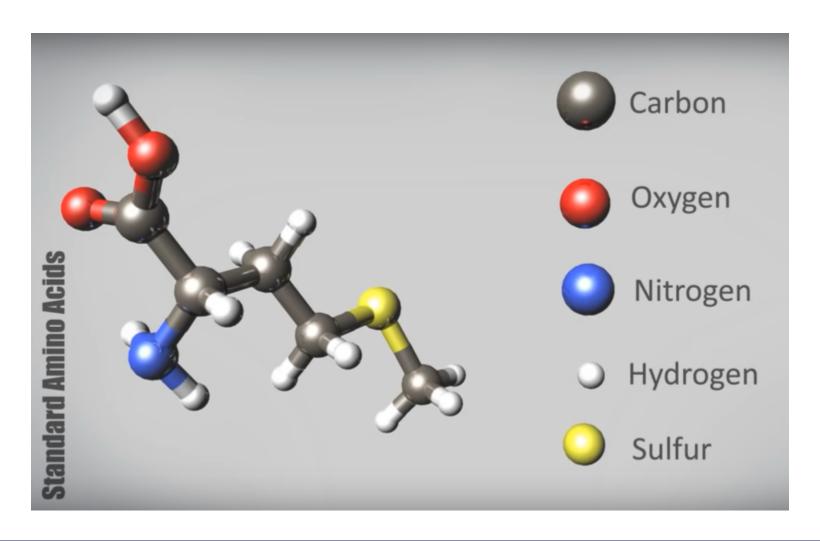


#### Protein's building blocks: the amino acids



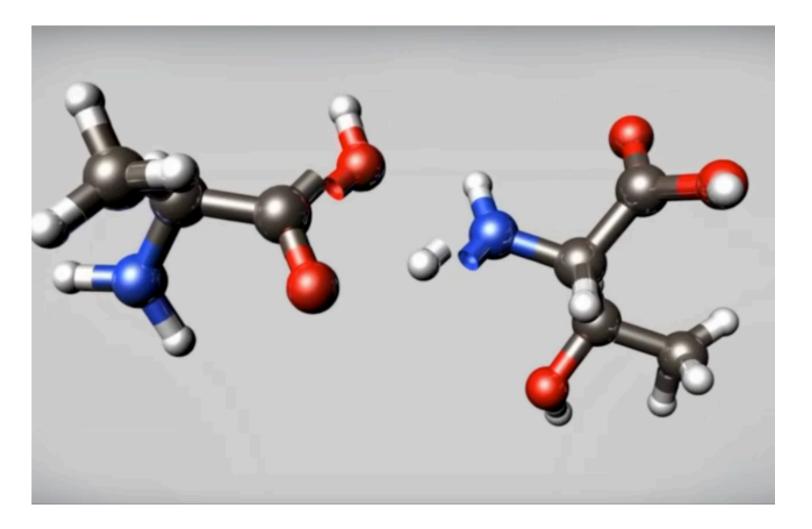


#### What are they made of?



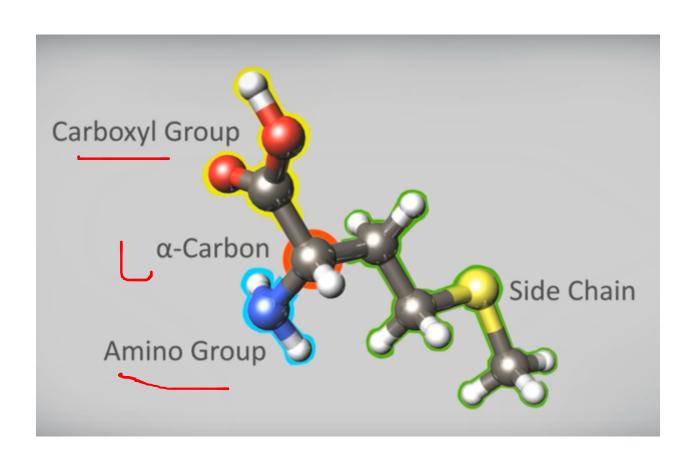


## Primary structure How polypeptide chains are formed?





#### Amino acids are chiral (L - amino acids)

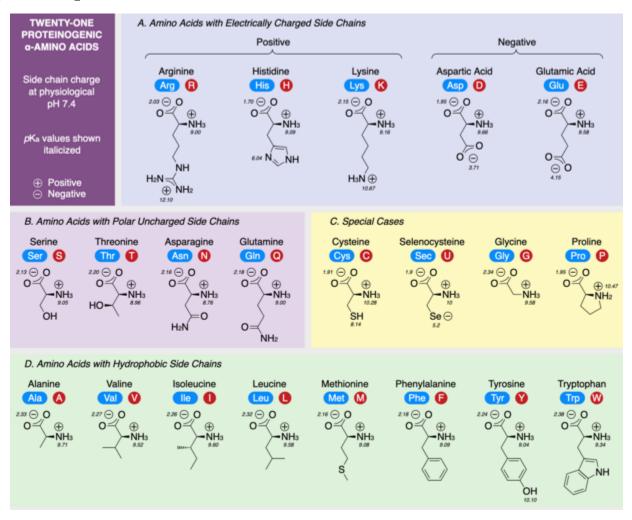


- Carboxyl group
   COO COOH
- Amino group-NH<sub>2</sub> <===> -NH<sub>3</sub>+
- α-Carbon
- Side chain



#### How to group the amino acids?

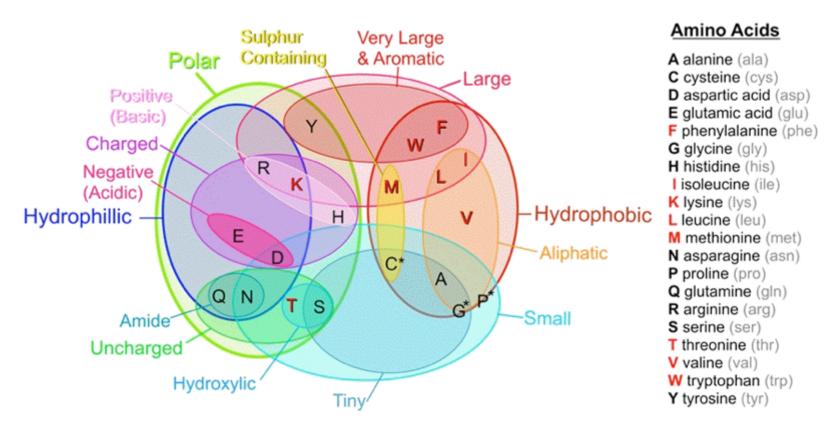
- Charge +/-
  - Acid vs Basic
- Polarity (polar/non-polar)
  - Type, distribution
- Size
  - Length, weight, volume,
  - surface area
- Type (Aromatic/Aliphatic)



https://upload.wikimedia.org/wikipedia/commons/4/4f/ProteinogenicAminoAcids.svg



#### How to group the amino acids?



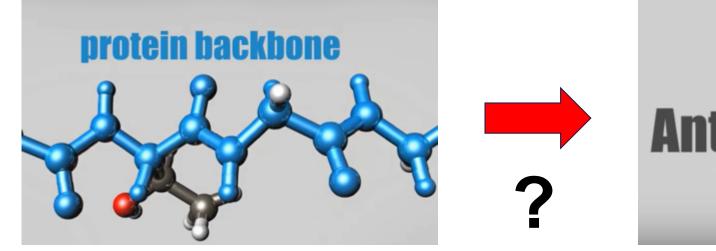
<sup>\*</sup> Unlike some other sources, J. Mol. Recognit., 17, 17-32 (2004) lists G and P as 'Neutral' (not hydrophobic), and does not include them in the group of Aliphatics C is sometimes listed as polar, uncharged and weakly acidic.

Essential amino acids are shown in red. These cannot be synthesized by the human body, and must be obtained from food.

http://betarhythm.blogspot.com



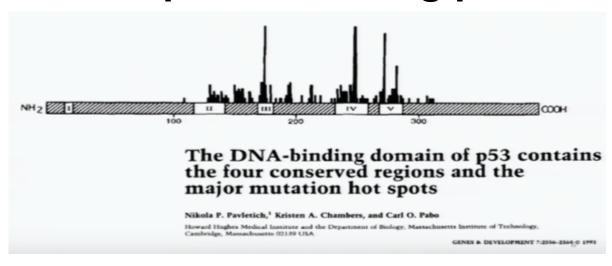
#### The protein folding problem





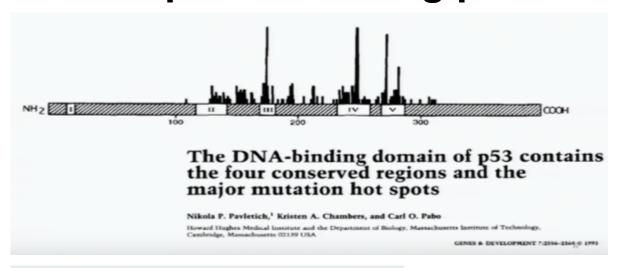


#### The protein folding problem





#### The protein folding problem





#### Crystal Structure of a p53 Tumor Suppressor-DNA Complex: Understanding Tumorigenic Mutations

Yunje Cho, Svetlana Gorina, Philip D. Jeffrey, Nikola P. Pavletich

SCIENCE • VOL. 265 • 15 JULY 1994



#### Amino acids are held together by different forces

- Hydrophobicity (Entropy)
- Salt bridges
- H bonds
- Di-sulfide bridges
- Aromatic interactions

Mostly proteins are held together by polar interactions and entropy

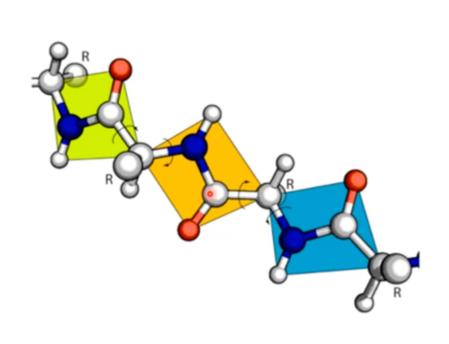


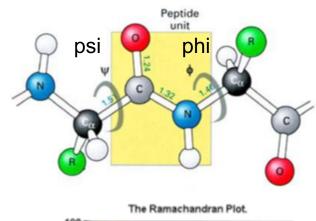
## Proteins reach the conformation that minimizes their Free Energy

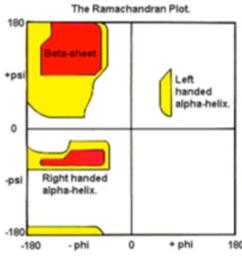




#### Can polypeptide chains adopt any conformation?

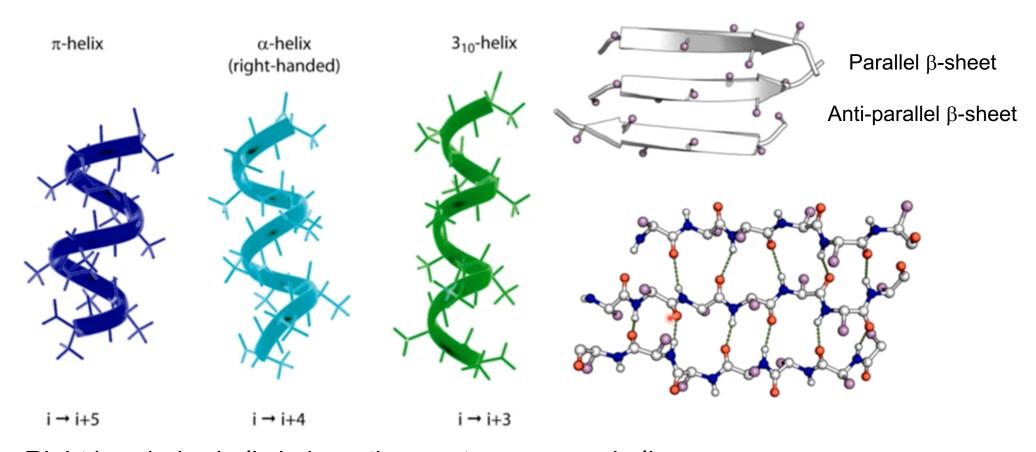








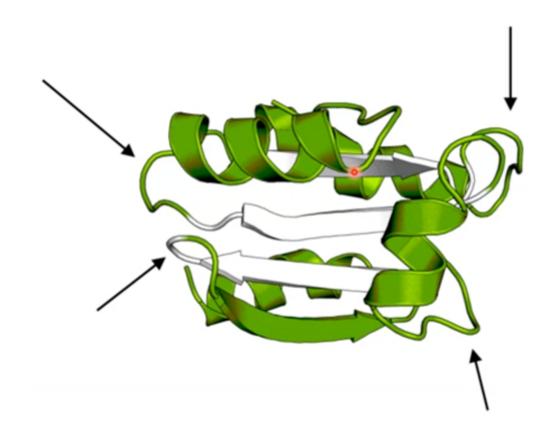
#### Secondary structure: $\alpha$ -helices and $\beta$ -sheets



Right handed  $\alpha\text{-helix}$  i+4 are the most common  $\alpha\text{-helices}$ 



## Secondary structure: Turn, loops and bends and disordered regions

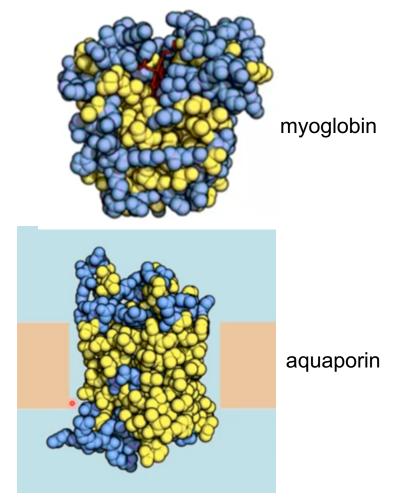


Disordered regions are regions with **non-stable** conformations



#### Hydrophobicity and hydrophilicity

Hydrophobic side chains such as in Val, Iso, Leu, Met, Phe, Tyr go into the core of the protein or into the membrane

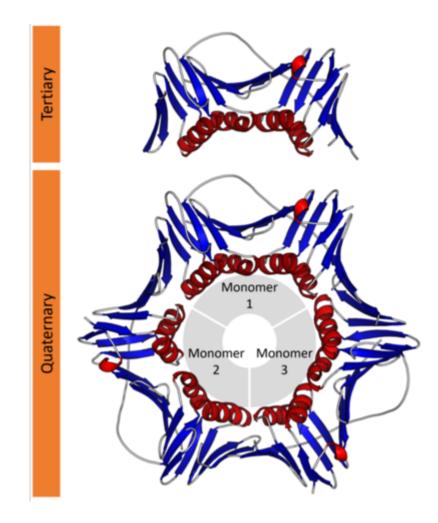




#### **Tertiary and Quaternary protein structure**

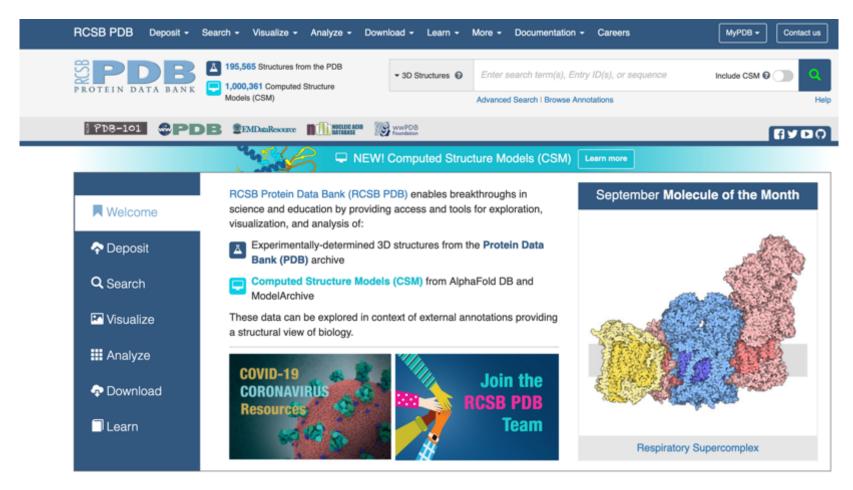
**Tertiary structures** are arrangements of secondary structure elements within one protein chain

**Quaternary structures** are assemblies of multi-chain complexes





#### Where do we find protein structure data?



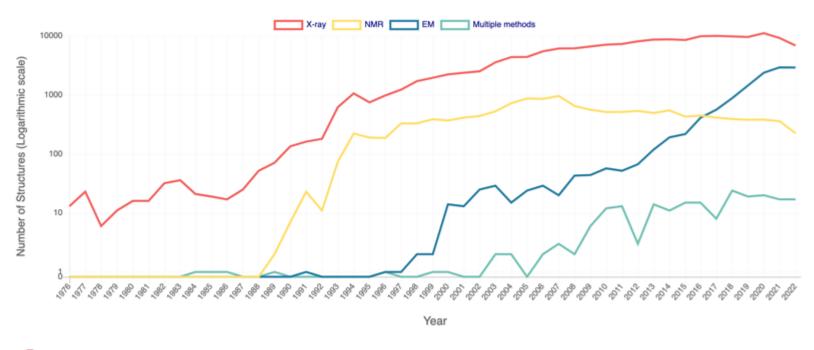
https://www.rcsb.org/



## How do we experimentally determine the protein structure?



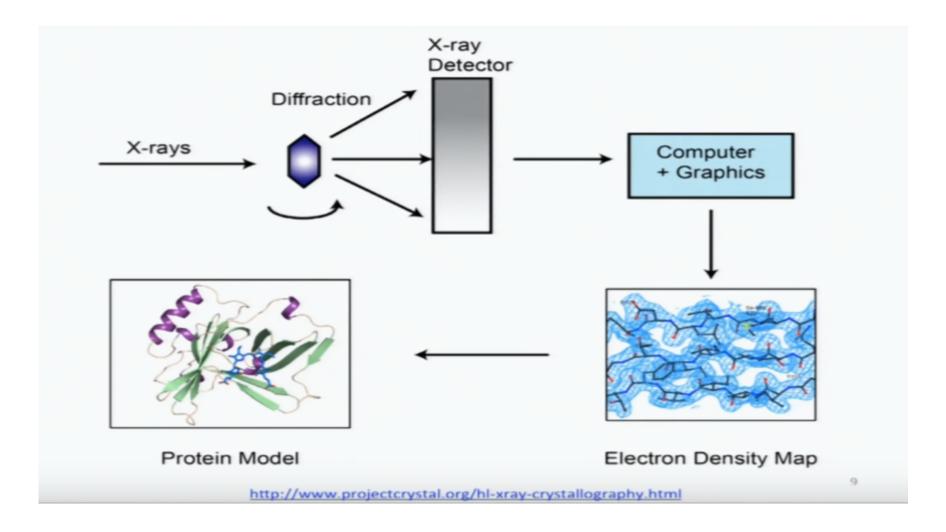




- X-ray: X-RAY DIFFRACTION, FIBER DIFFRACTION, or POWDER DIFFRACTION
- NMR: SOLUTION NMR or SOLID-STATE NMR
- EM: ELECTRON MICROSCOPY or ELECTRON CRYSTALLOGRAPHY or ELECTRON TOMOGRAPH.
- MULTIPLE METHODS: Multiple experimental methods. For example, if a structure is solved by X-RAY DIFFRACTION AND NEUTRON DIFFRACTION, it will be counted only in

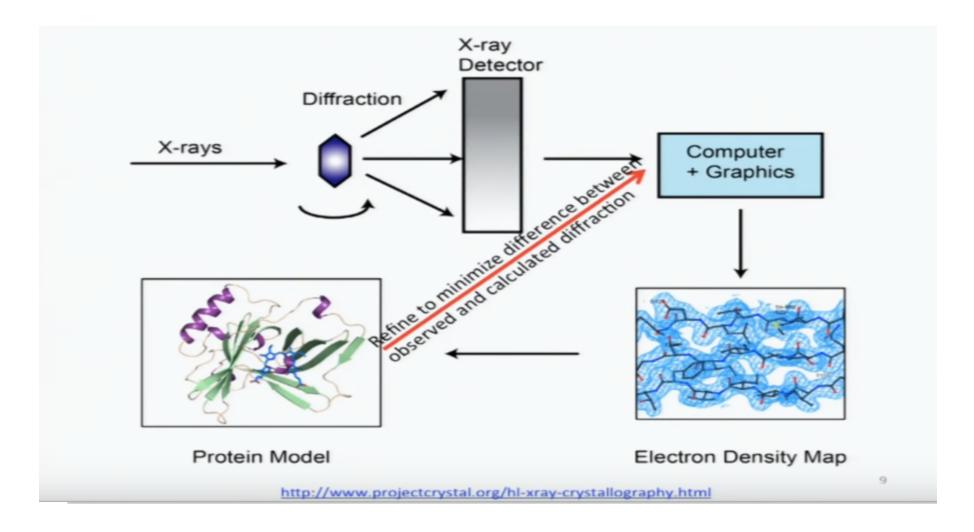


#### Overview of the X-ray crystallographic method



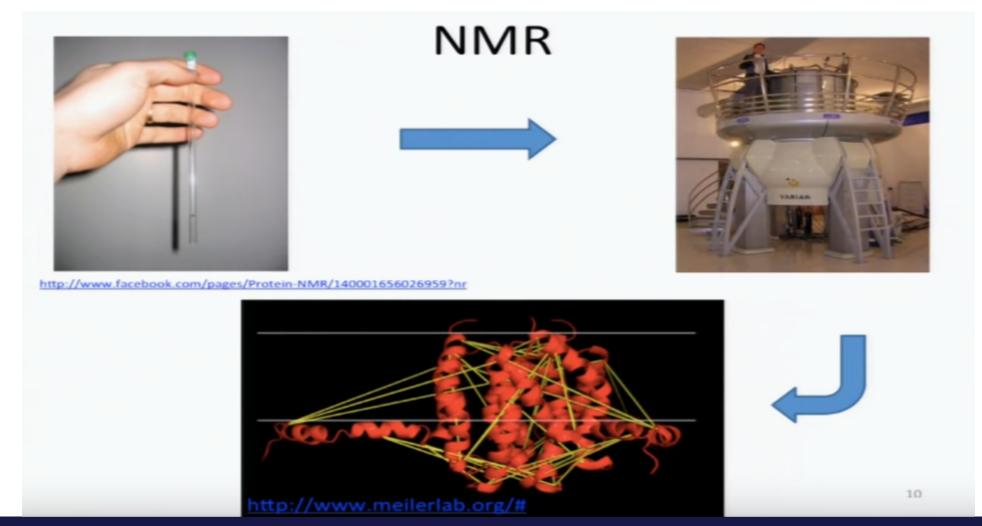


#### Overview of the X-ray crystallographic method



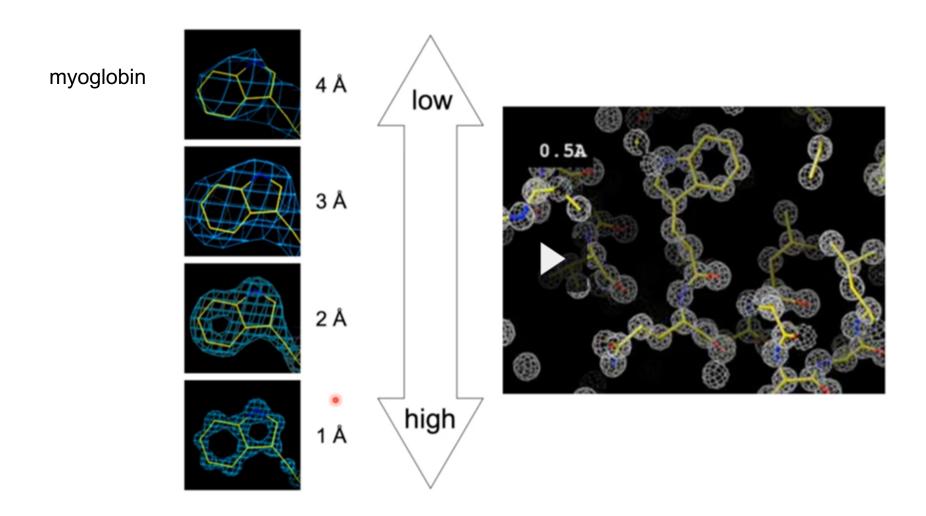


## Nuclear magnetic resonance spectroscopy of proteins





#### The importance of protein structure resolution





#### **Quality of the protein structure: Goodness of model fitness**

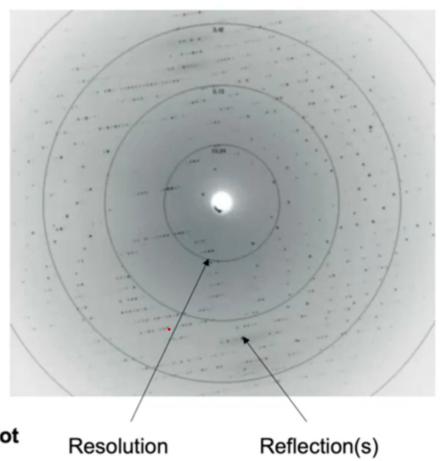
Individual reflections

$$I_{hkl} \propto \left| F_{obs}(hkl) \right|^2$$

R-factor:

$$R = \frac{\sum \left\| F_{obs} \right| - \left| F_{calc} \right\|}{\sum \left| F_{obs} \right|}$$

- - Like R-factor
  - Unbiased measure.
  - Calculated on 5-10% of data **not** included in refinement.



rings

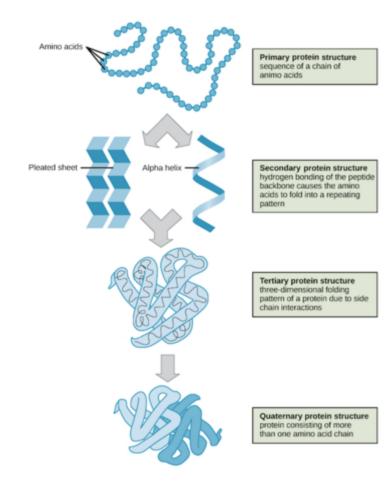


#### Where do we find the information in PDB?





### Summary: The four levels of protein structure





### Video link to protein structure



https://www.youtube.com/watch?v=qBRFIMcxZNM