Globular proteins
Myoglobin and hemoglobin

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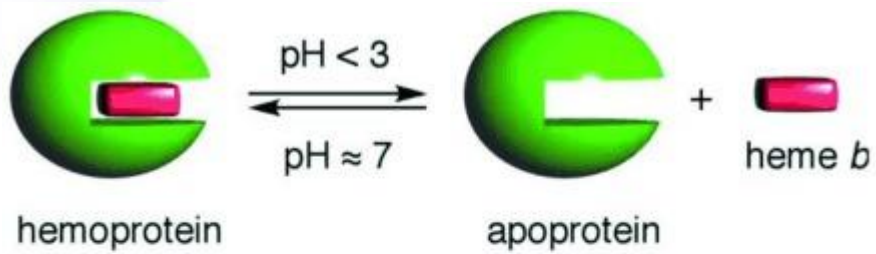
Functions of myoglobin and hemoglobin

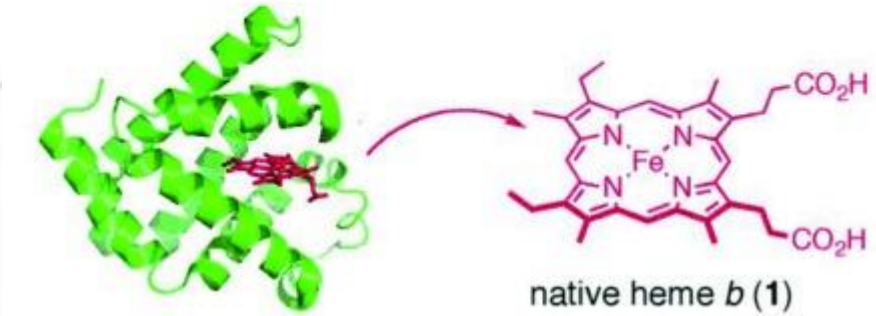
- Myoglobin functions in storing O_2 in muscles. During periods of oxygen deprivation, oxymyoglobin releases its bound oxygen.

- Hemoglobin:
  - transport of O_2 and CO_2
  - blood buffering

Hemoproteins

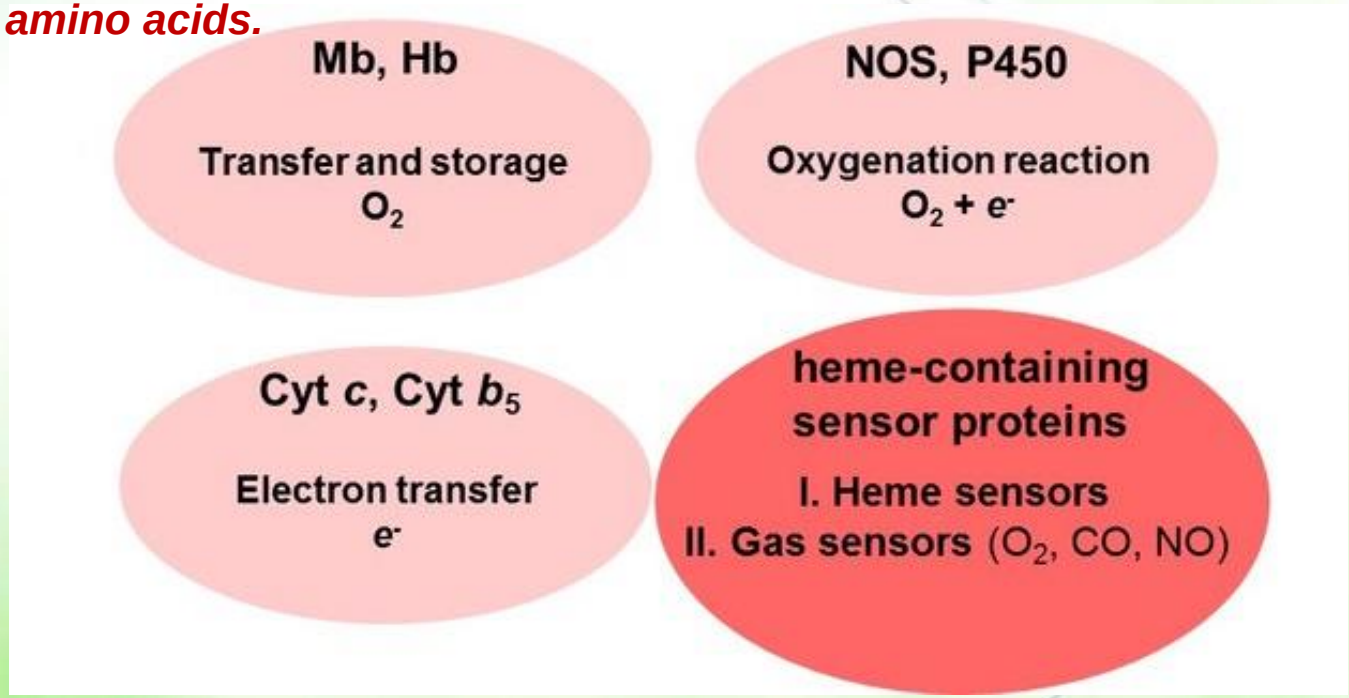
- Many proteins have heme as a prosthetic group called hemoproteins.





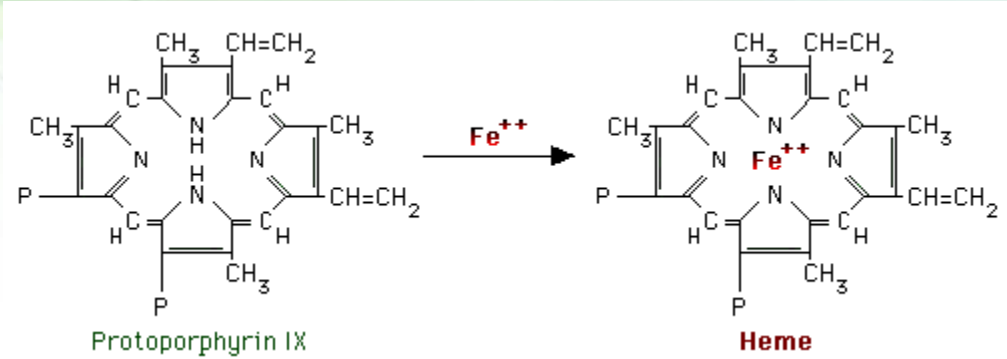
The protein environment dictates the function of the heme.

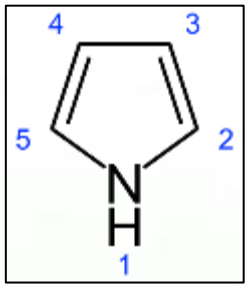
A prosthetic group is a tightly bound, specific non-polypeptide unit required for the biological function of some proteins. The prosthetic group may be organic (such as a vitamin, sugar, or lipid) or inorganic (such as a metal ion), but is not composed of amino acids.

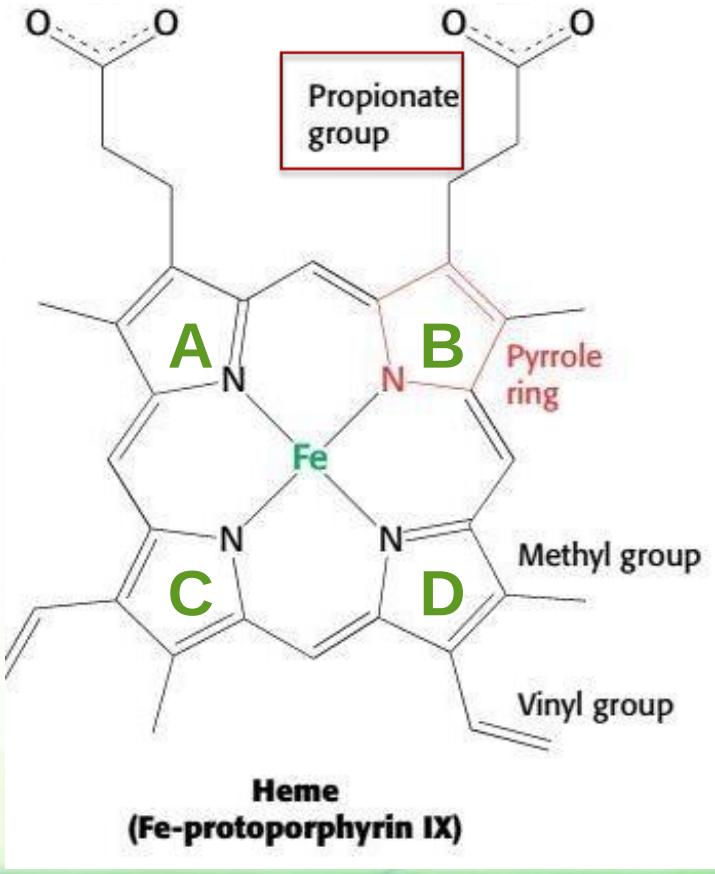


Heme structure

- It is a complex of protoporphyrin IX + Iron (Fe^{2+}).
- The porphyrin is planar and consists of four rings (designated A-D) called pyrrole rings.
- Each pyrrole can bind two substituents.
- Two rings have a propionate group each.
- *Note: the molecule is hydrophobic.*
- Fe has six coordinates of binding.

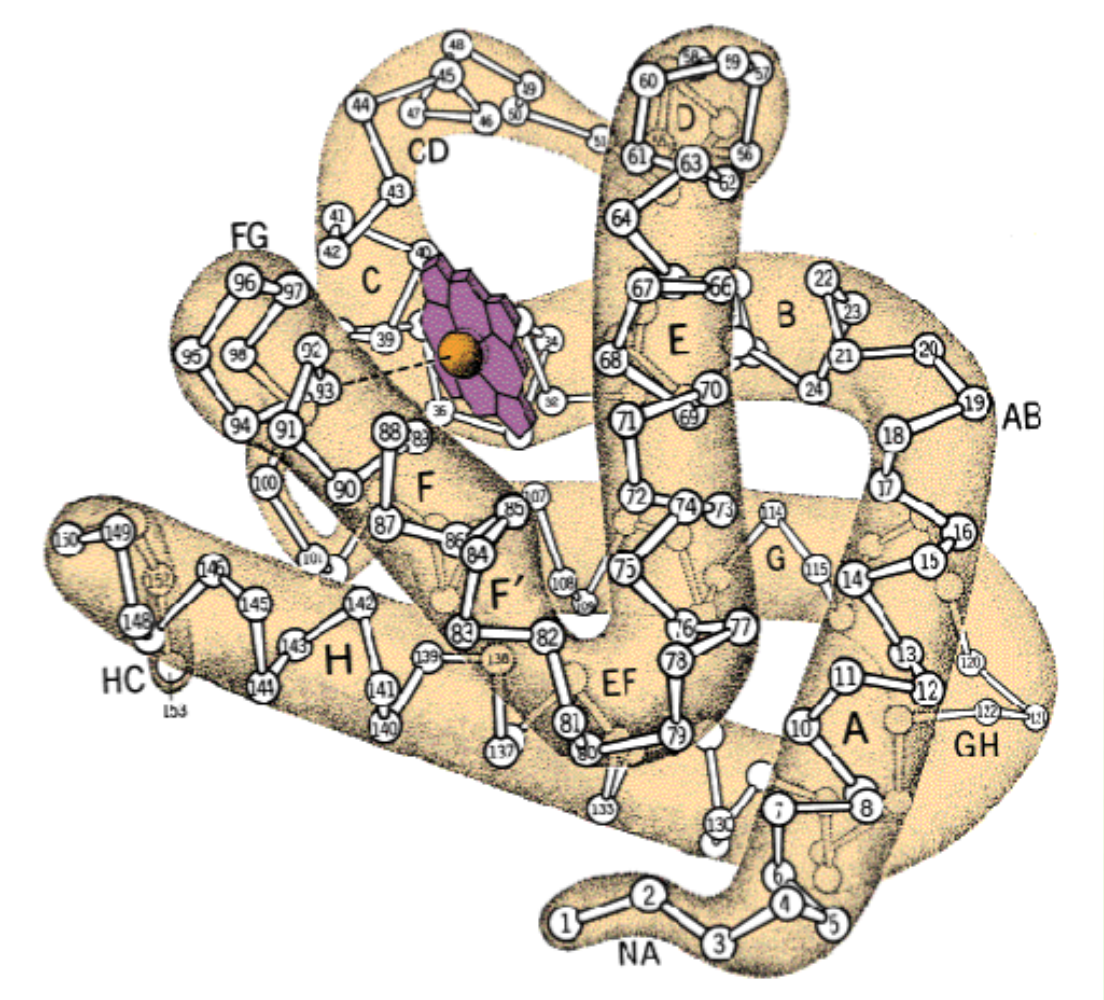






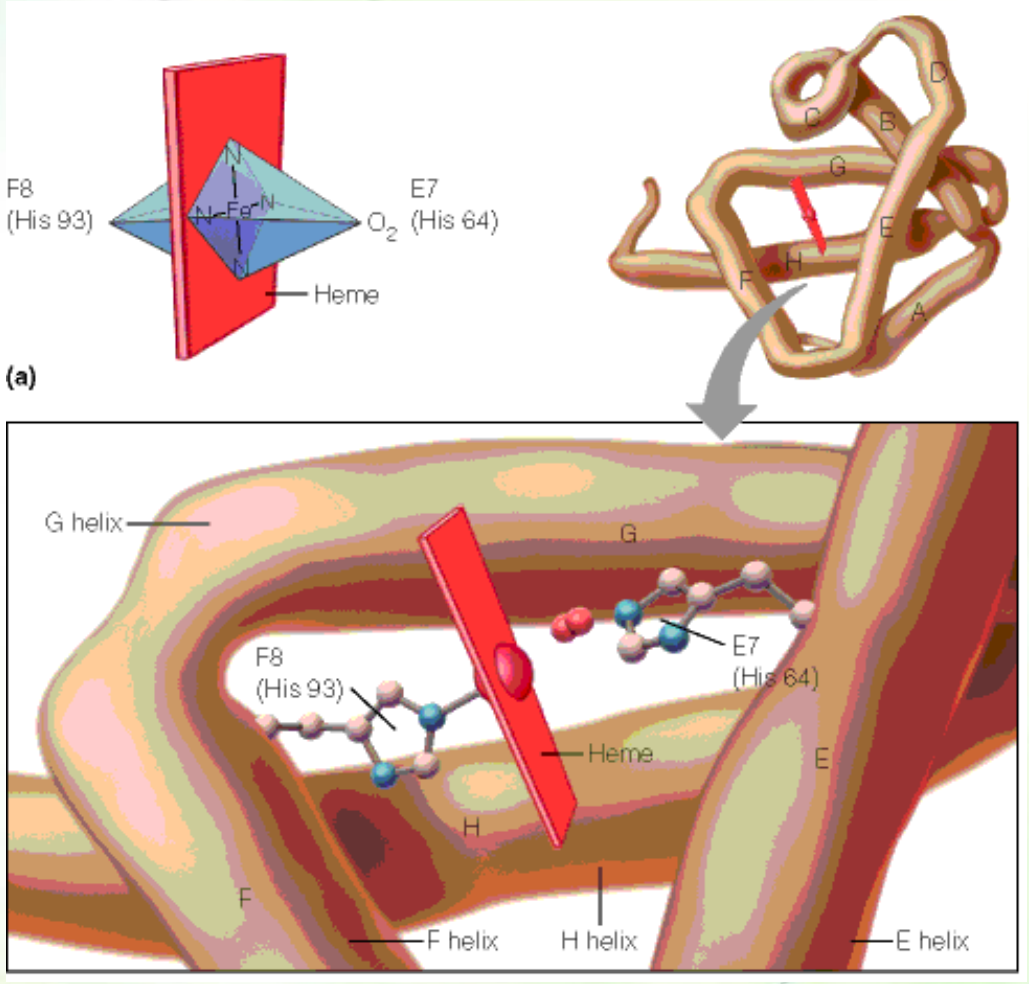
Structure of myoglobin

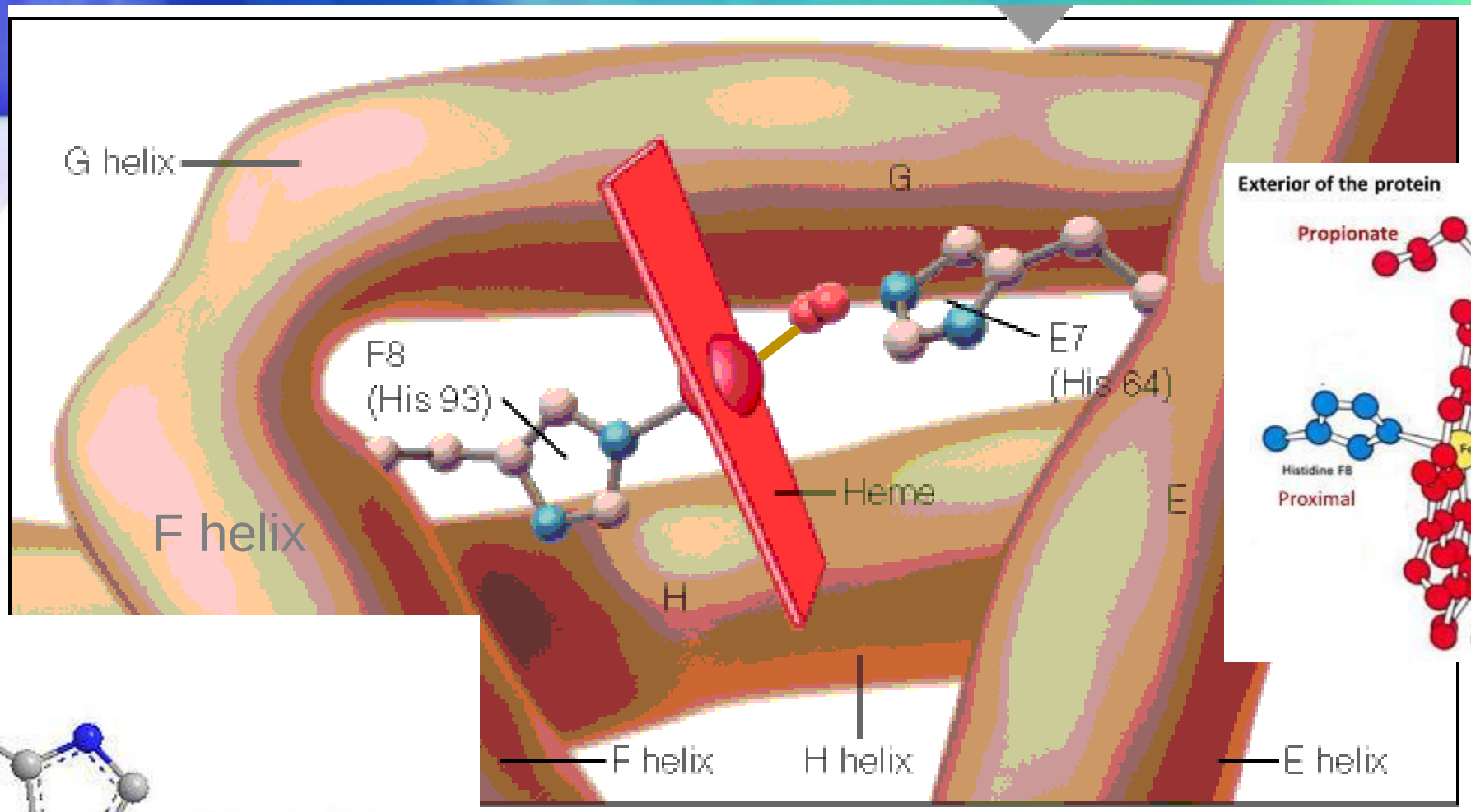
- Myoglobin is a monomeric protein that is mainly found in in muscle tissue.
- It includes a prosthetic group, the heme group
- It can be present in two forms:
  - oxymyoglobin (oxygen-bound)
  - deoxymyoglobin (oxygen-free)
- The tertiary structure of myoglobin 8 α-helices, designated A through H, that are connected by short non-helical regions.

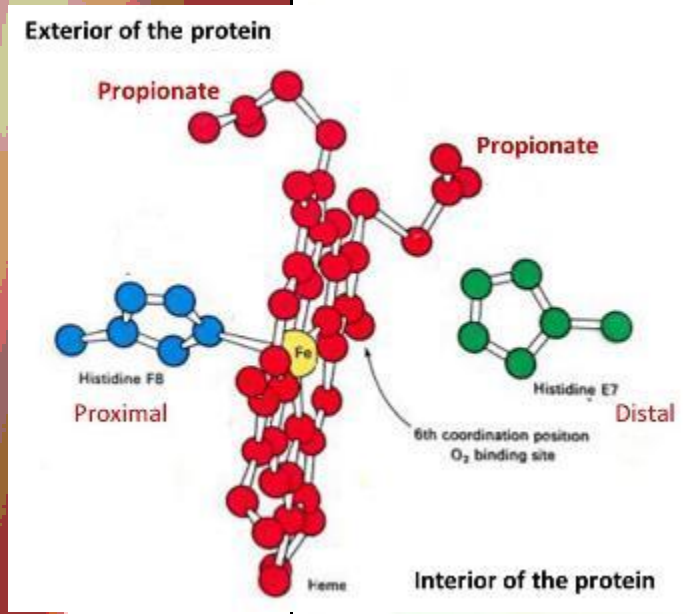


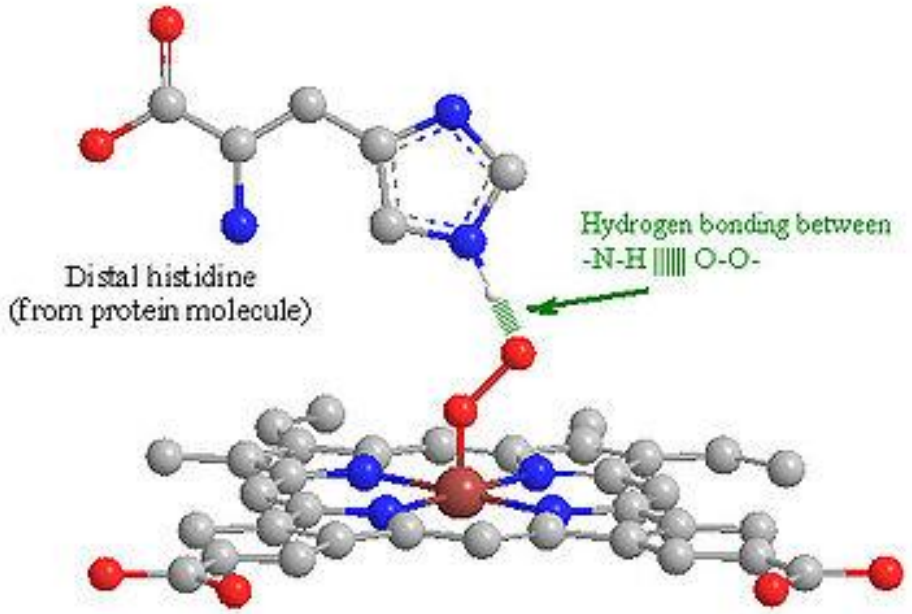
Arrangement of amino acids

- Like other globular protein, amino acid R-groups exposed on the surface of the molecule are generally hydrophilic, while those in the interior are predominantly hydrophobic.
- Except for two histidine residues in helices E and F (known as E7 and F8)
- F8 His is designated as proximal His, whereas E7 His is known as distal His



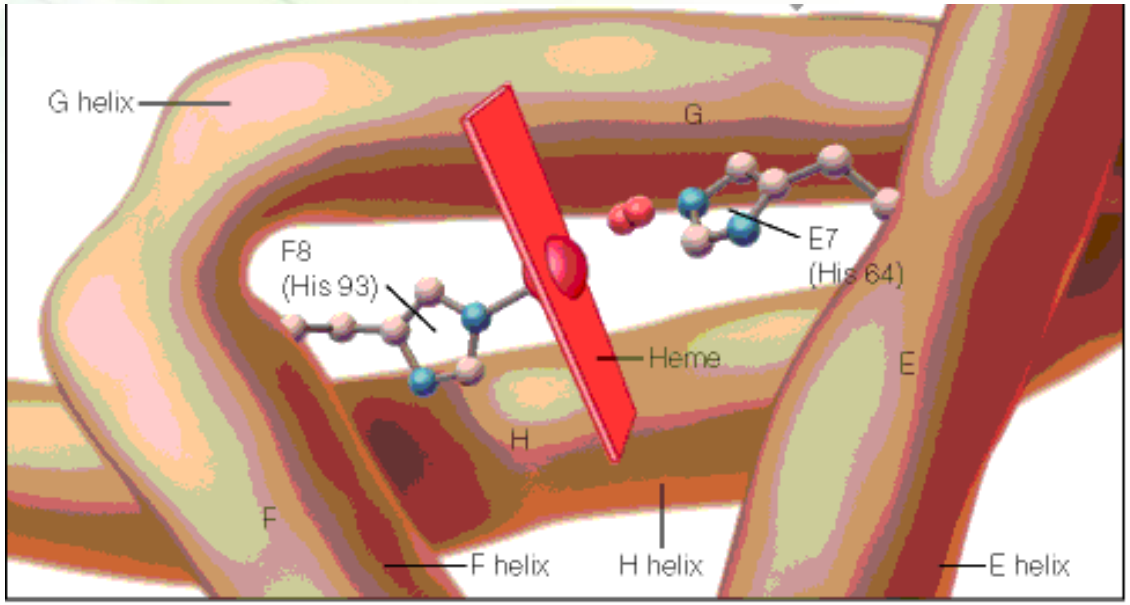

G helix
G
E7
(His 64)
F8
(His 93)
Heme
F helix
H
E
F helix
H helix
E helix


Exterior of the protein
Propionate
Propionate
Histidine F8
Proximal
Fe
Histidine E7
Distal
6th coordination position
O_2 binding site
Heme
Interior of the protein


Distal histidine
(from protein molecule)
Hydrogen bonding between
-N-H ||||| O-O-

Iron

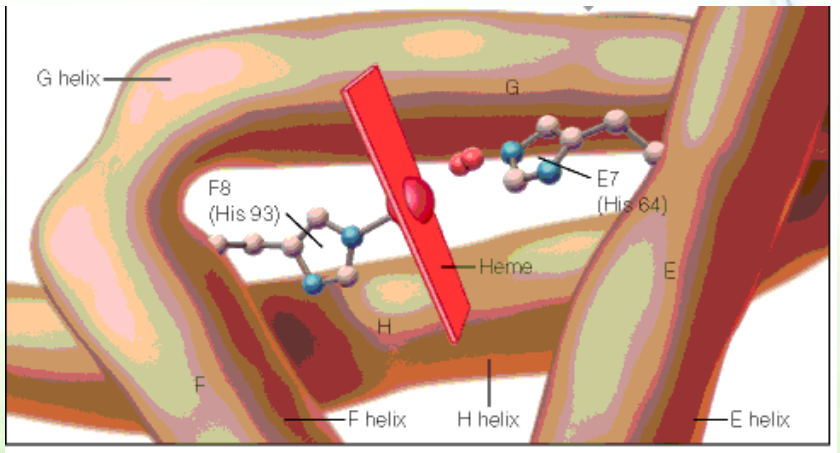
- Iron can bind in the center of the four rings.
- Fe is in the ferrous state (Fe^{2+}) can form 6 bonds:
  - 4 with the nitrogen of the rings,
  - One (known as the fifth coordinate) with the nitrogen of a histidine imidazole (known as proximal His).
  - One with O_2 (the sixth coordinate)



- Oxidation of iron to the Fe^{3+}, ferric, state makes the molecule incapable of normal O_2 binding
- Upon absorption of light, heme gives a deep red color.

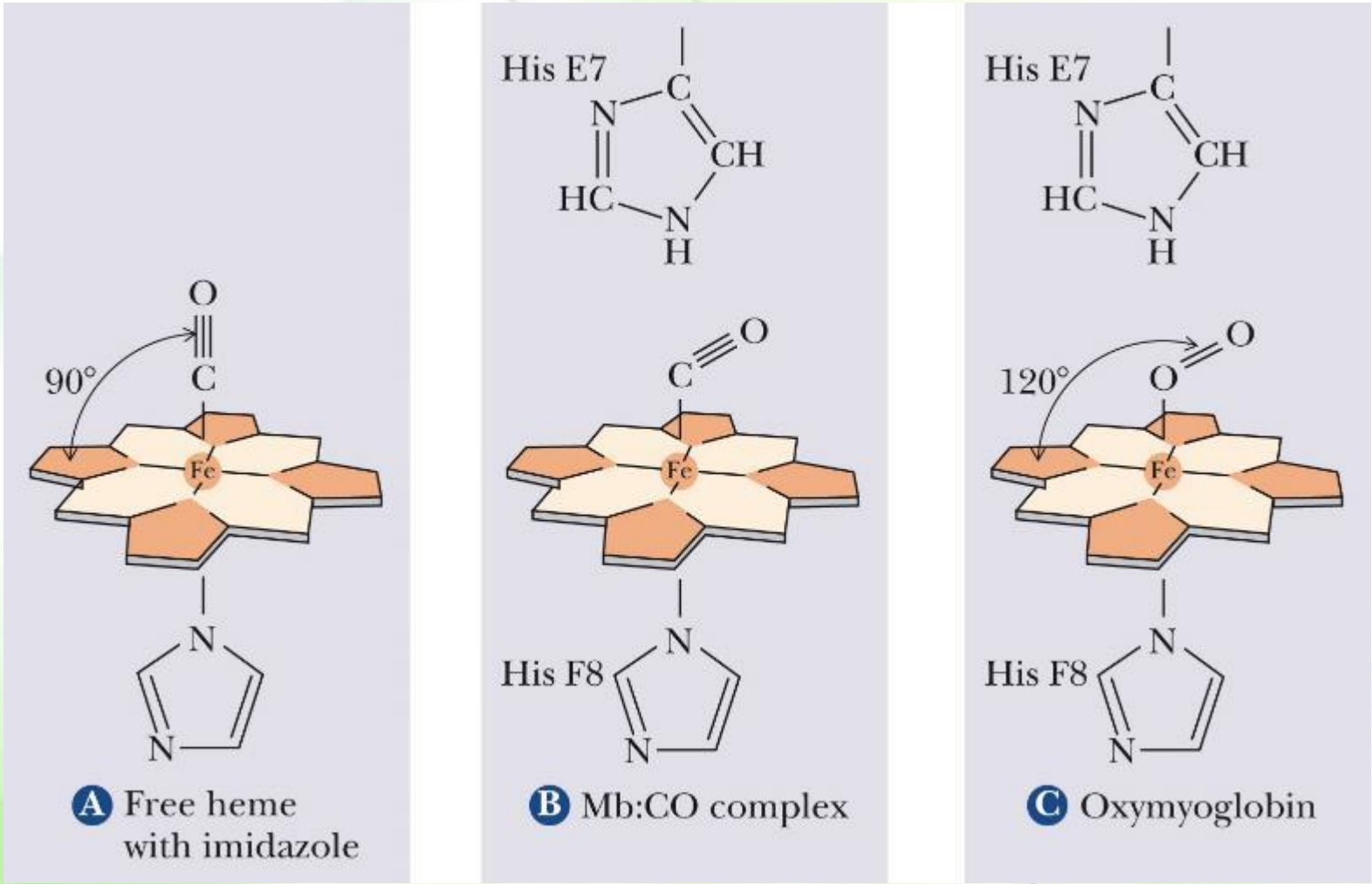
Structure-function relationship

- The planar heme group fits into a hydrophobic pocket of the protein and the myoglobin-heme interaction is stabilized by hydrophobic attractions.
- The heme group stabilizes the tertiary structure of myoglobin.
- The distal histidine acts as a gate that opens and closes as O_2 enters the hydrophobic pocket to bind to the heme.
- The hydrophobic interior of myoglobin (or hemoglobin) prevents the oxidation of iron, and so when O_2 is released, the iron remains in the Fe(II) state and can bind another O_2.



Another significance of distal histidine

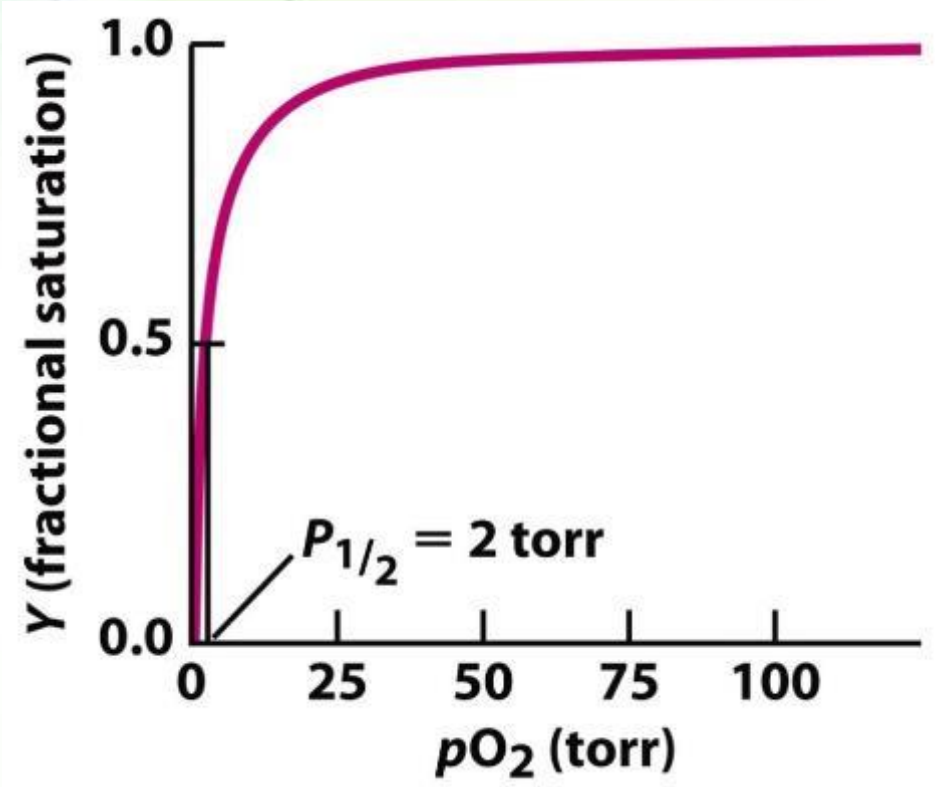
- CO prefers straight bonding, but O_2 prefer bent bonding.
- CO binds to free heme many orders of magnitude compared to O_2.
- CO binding to myoglobin-bound heme only 250 times more than O_2.
- CO occupies 1% of hemoglobin, but 99% if distal His does not exist.



Accidents

Oxygen binding to myoglobin

- Myoglobin binds O_2 with high affinity.
- The P50 (oxygen partial pressure required for 50% of all myoglobin molecules) for myoglobin ~2.8 torrs or mm Hg.
- Given that O_2 pressure in tissues is normally 20 mm Hg, it is almost fully saturated with oxygen at normal conditions.



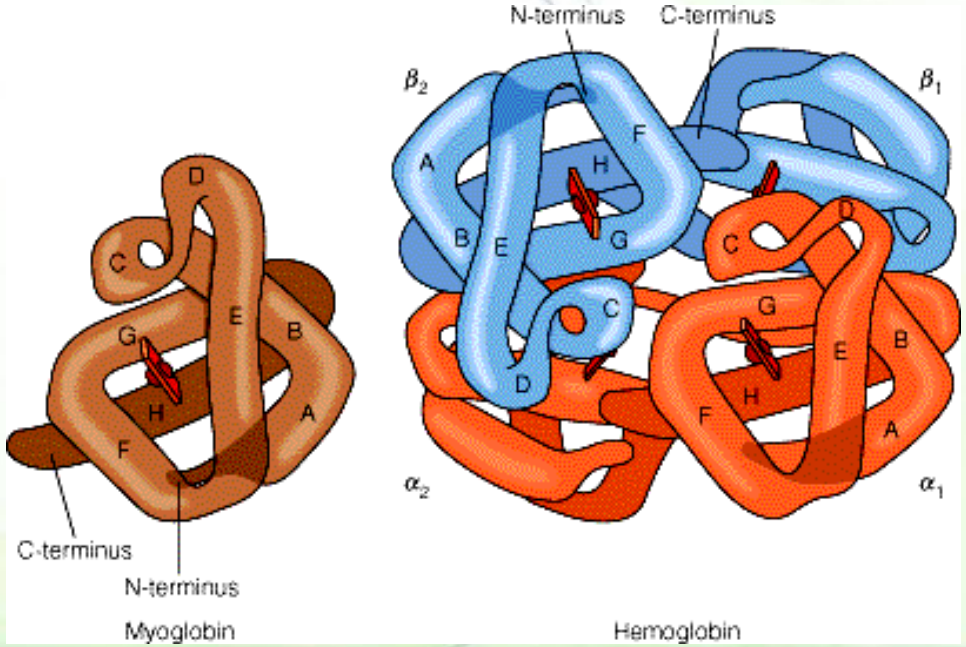
The binding of **O2** to myoglobin follows a hyperbolic saturation curve.

Hemoglobin

Hemoglobin structure

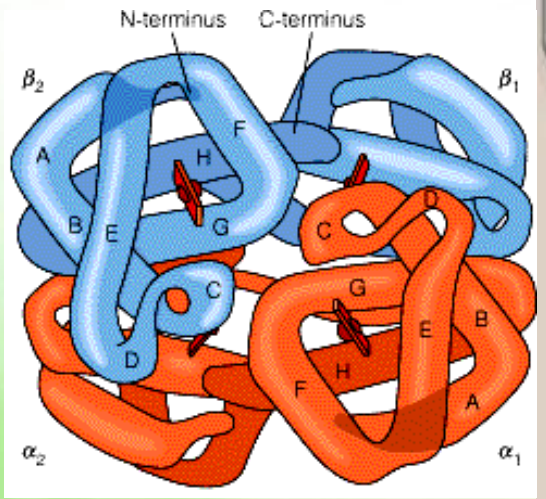
- Hemoglobin is tetrameric hemeprotein (fur protein chains known as globins with each bound to heme.
- In adults, the four globin proteins are of two different types known as α and β, so a hemoglobin protein is an α2β2 globin protein.
- The α and β chains contain multiple α-helices where α contains 7 α-helices and β contains 8 α-helices (similar to myoglobin).

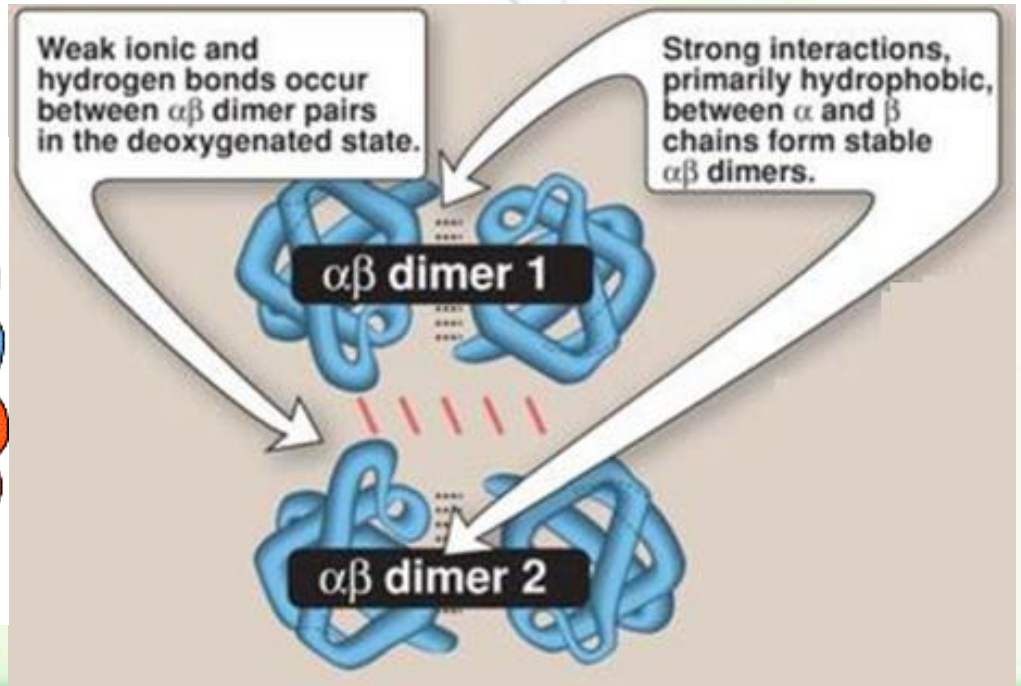
  - The alpha chain is composed of 141 aa residues.
  - The beta chain is composed of 146 aa residues.



How are the subunits bound?

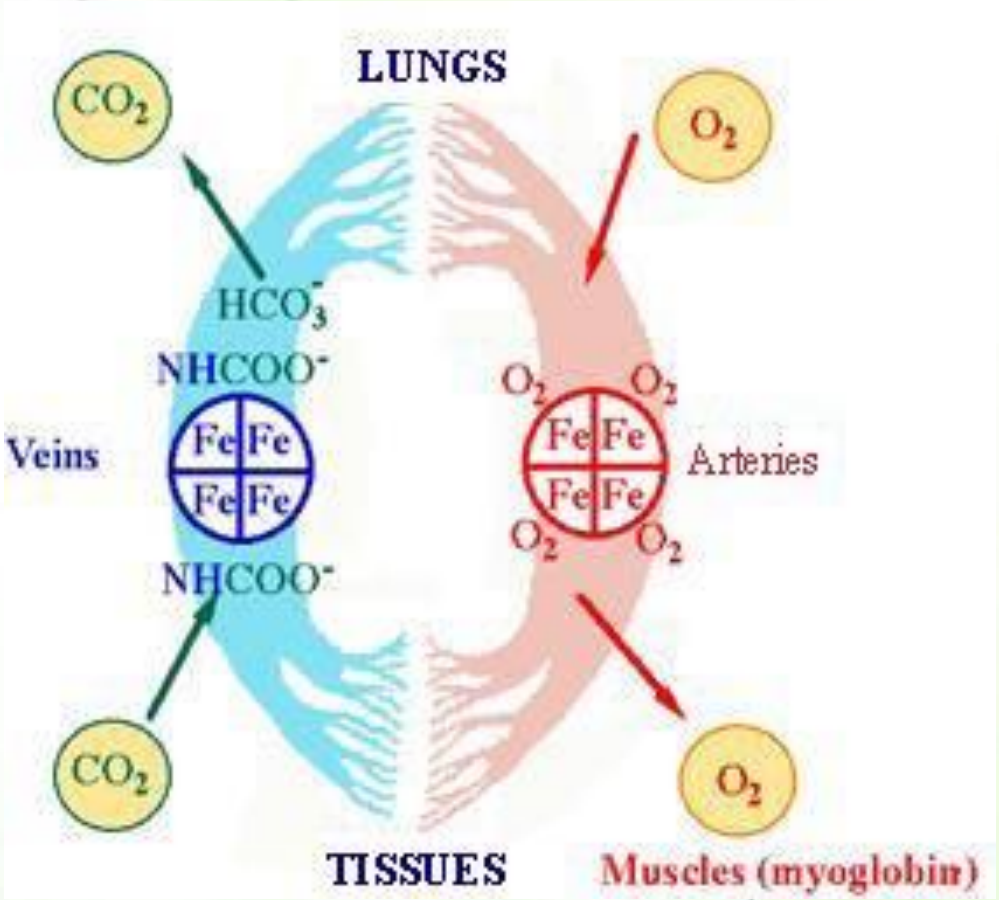
- A dimer of dimers (I made up this term)
  - (α-β)2
- The chains interact with each other via hydrophobic interactions.
  - Therefore, hydrophobic amino acids are not only present in the interior of the protein chains, but also on the surface.
- Electrostatic interactions (salt bridges) and hydrogen bonds also exist between the two different chains.





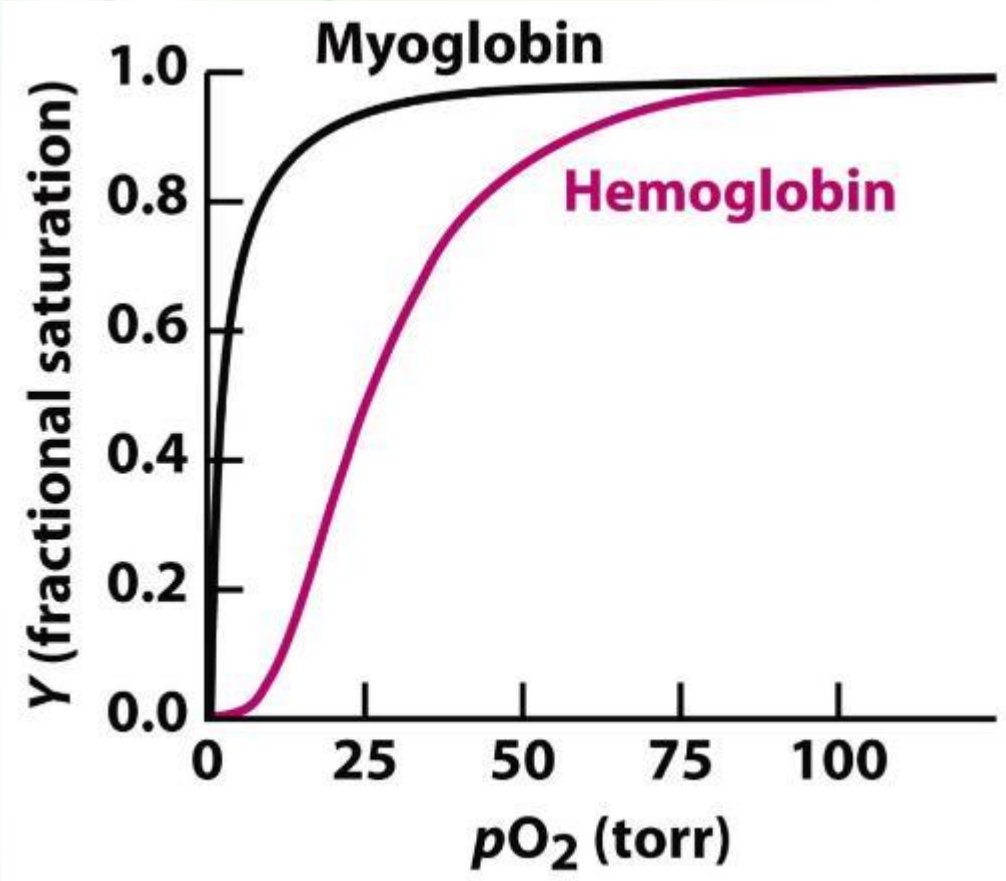
Oxygen binding to hemoglobin

- Hemoglobin must bind oxygen efficiently and become saturated at the high oxygen pressure found in lungs (approximately 100 mm Hg).
- Then, it releases oxygen and become unsaturated in tissues where the oxygen pressure is low (about 30 mm Hg).



The saturation curve

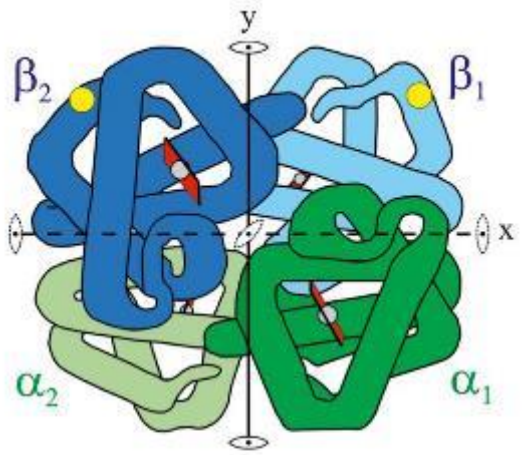
- The saturation curve of hemoglobin binding to O_2 has a sigmoidal shape.
  - A sigmoidal curve indicates that the protein has different structures.
- At 100 mm Hg, hemoglobin is 95-98% saturated (oxyhemoglobin).
- As the oxygen pressure falls, oxygen is released to the cells.
- In contrast to a low p50 for myoglobin, the p50 of hemoglobin is approximately 26 mm.

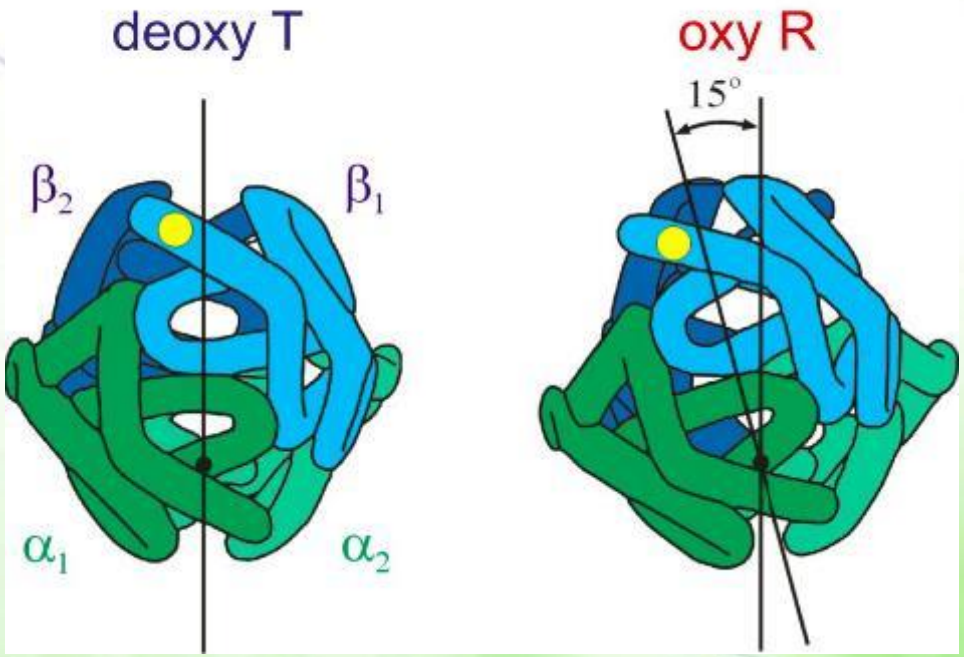


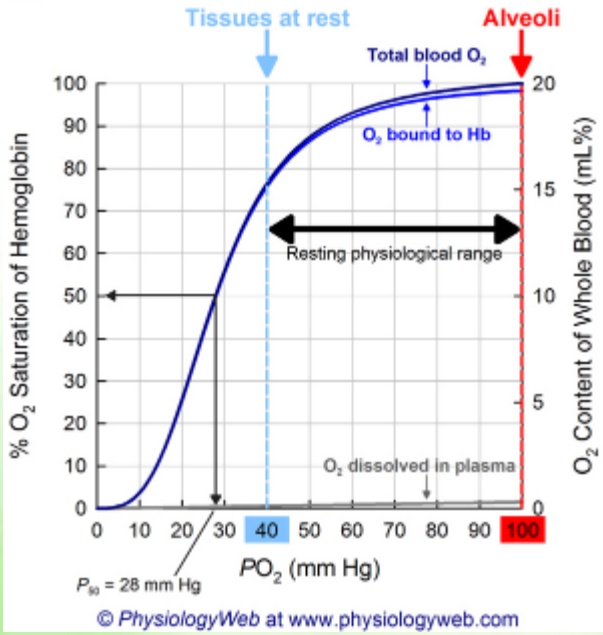
Hemoglobin is allosteric

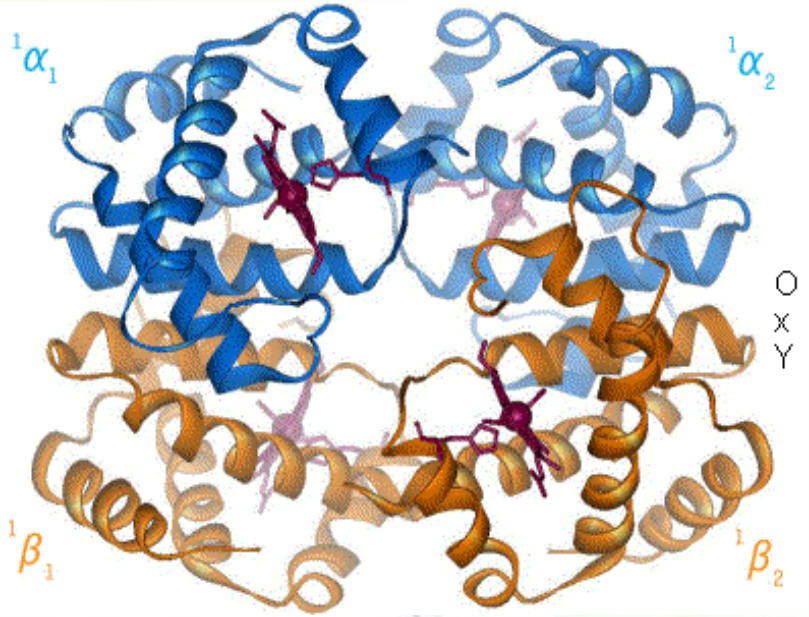
- Hemoglobin is an allosteric protein (from Greek "allos" = "other", and "stereos" = "shape").
  - An allosteric protein: a protein where binding of a molecule (ligand) to one part of the protein affects binding of a similar or a different ligand to another part of the protein.
- Hemoglobin exists in two forms, T-state and R-state
- The T-state is also known as the "taut" or "tense" state and it has a low-binding affinity to oxygen.
- The R-state is known as the "relaxed" state and it has 500 times higher affinity to oxygen than as the T conformation .
- Binding of O_2 causes conformational changes in hemoglobin, converting it from the low affinity T-state to the high affinity R-state .

Structural change of hemoglobin



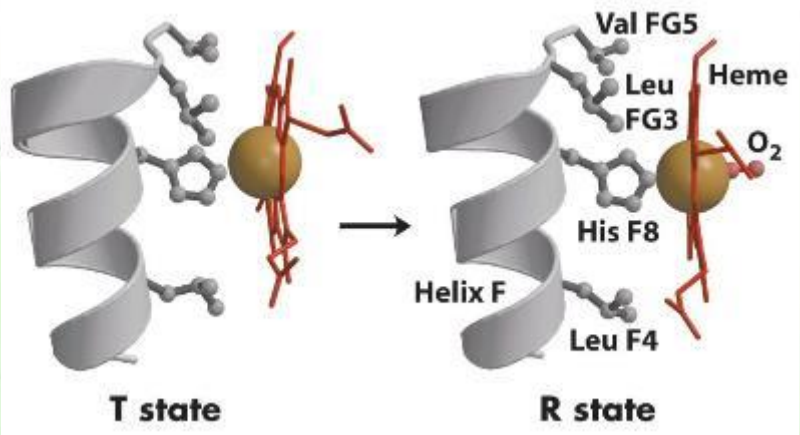


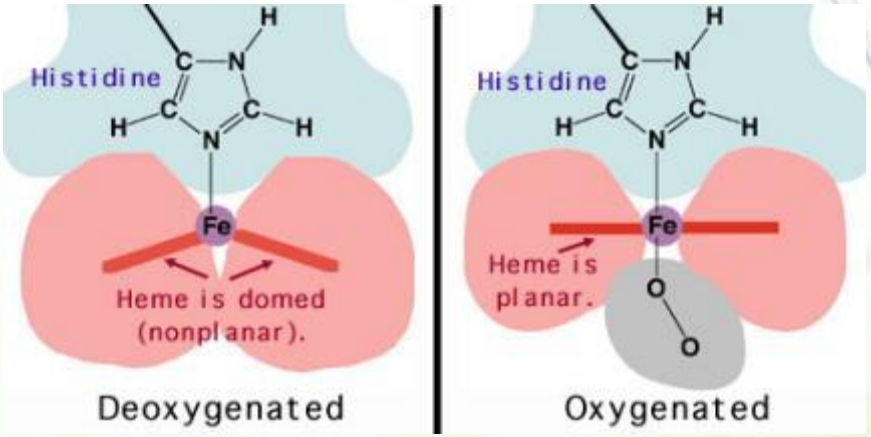




How does the structure change? (1)

- When heme is free of oxygen, it has a domed structure and iron is outside the plane of the heme group.
- When oxygen binds to an iron atom, heme adopts a planar structure and the iron moves into the plane of the heme pulling proximal histidine (F8) along with it.

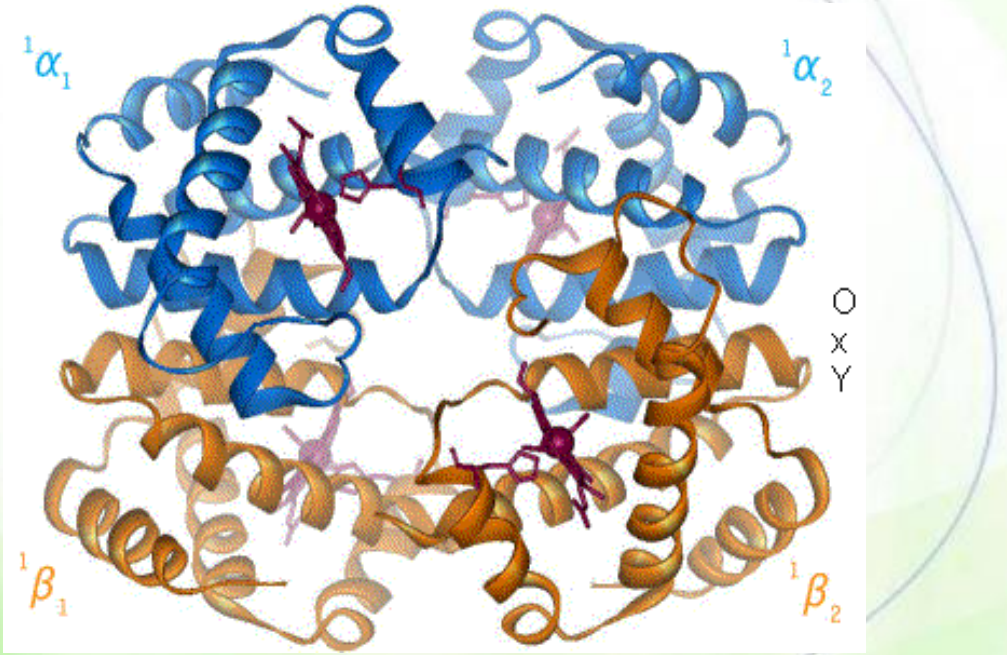




How does the structure change? (2)

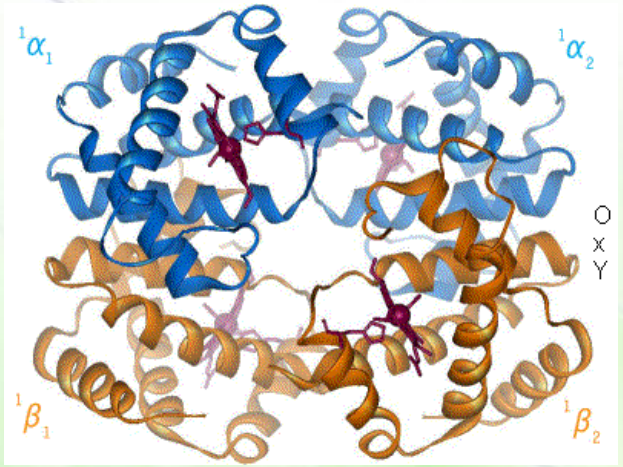
- This movement triggers
  - changes in tertiary structure of individual hemoglobin subunits
  - breakage of the electrostatic bonds at the other oxygen-free hemoglobin chains.

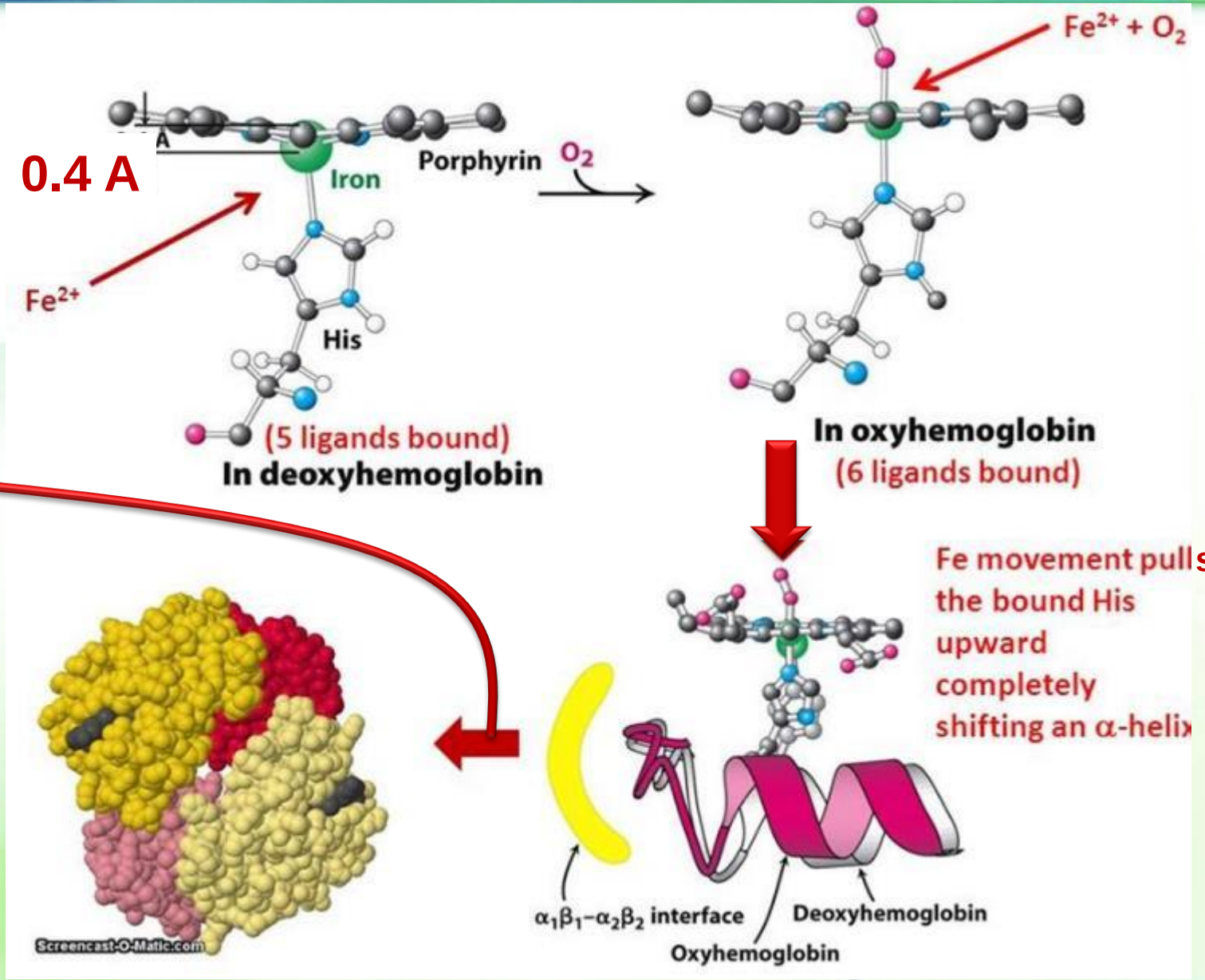
In myoglobin, movement of the helix does not affect the function of the protein.



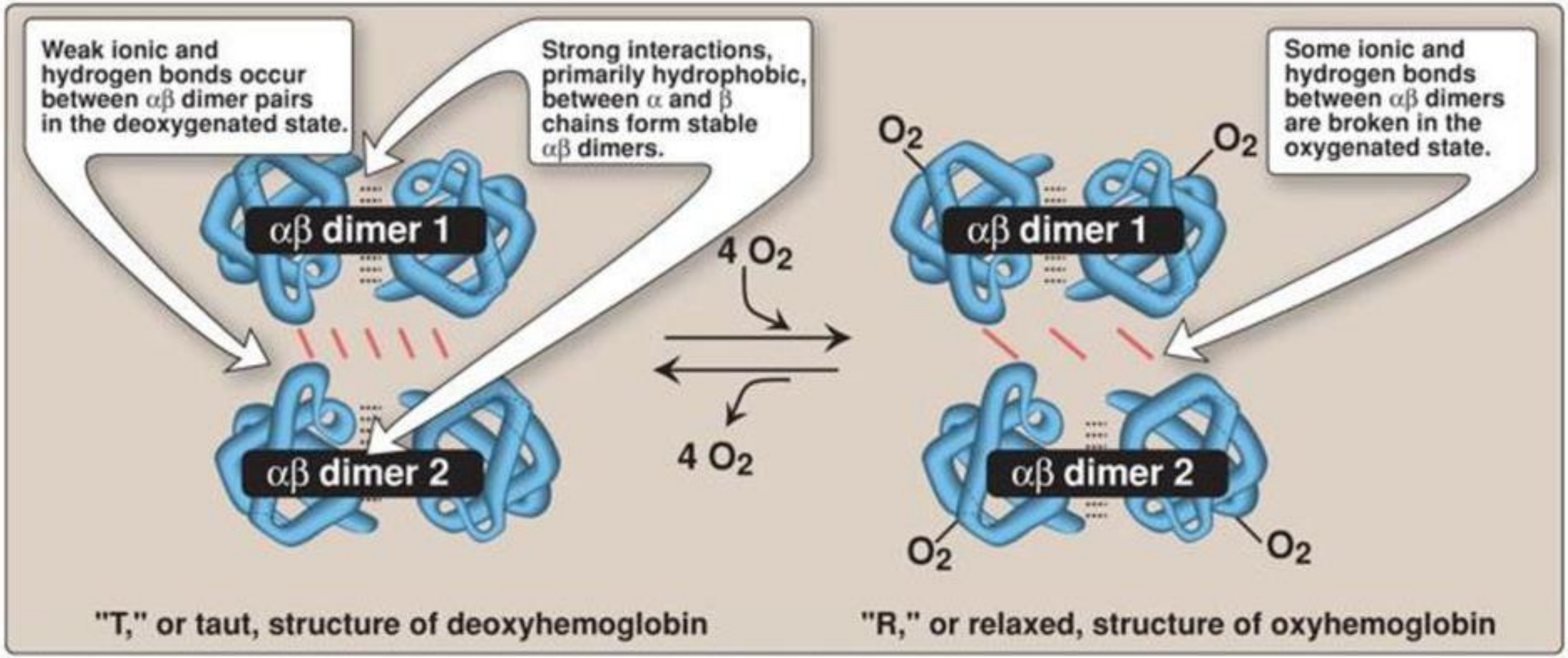
Structural amplification change

- **Changes in tertiary structure of individual hemoglobin subunits**
- **Breakage of the electrostatic bonds at the other oxygen-free hemoglobin chains.**



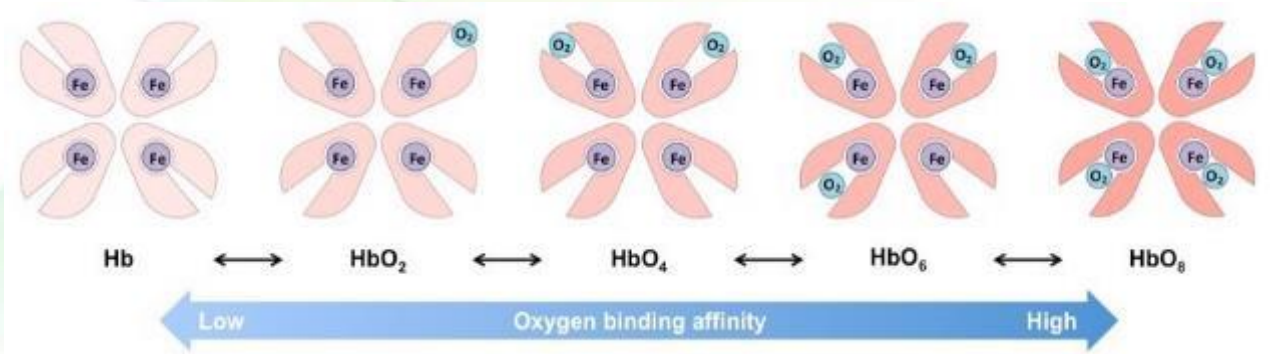


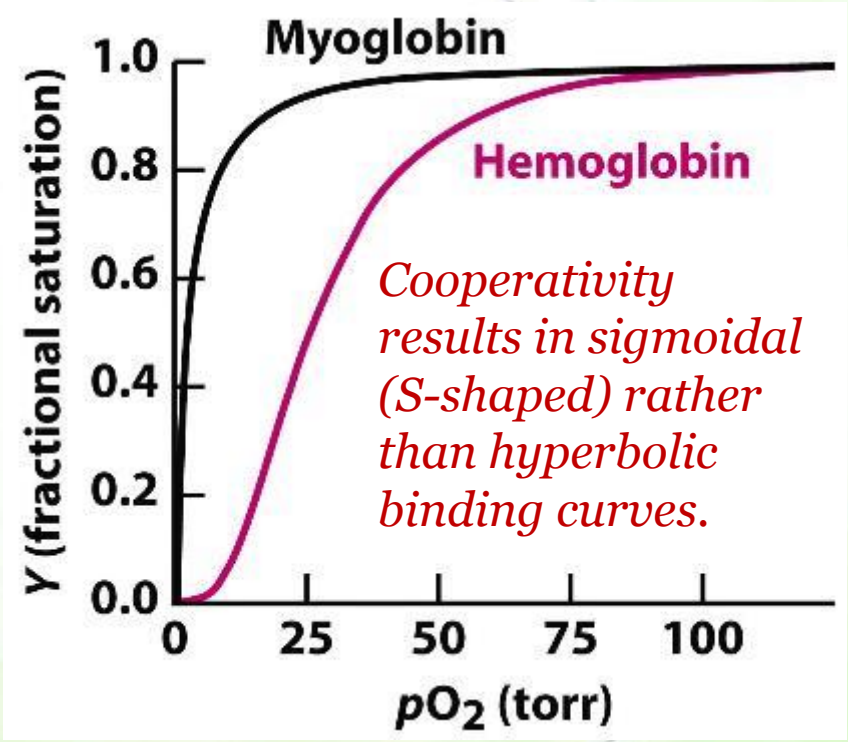
Electrostatic interactions are broken



Binding is cooperative

- Conformational changes lead to cooperativity among binding sites.
- Binding of the first O_2 breaks some salt bridges with the other chains increasing the affinity of the binding of a second molecule.
- Binding of the second O_2 molecule breaks more salt bridges increasing the affinity towards binding of a third O_2 even more, and so on.
- Binding is cooperative.
- Oxygen is a homotropic effector (the allosteric modulator is the substrate itself).





Some terminologies

- Homotropic allosteric regulator/effector: effector and ligand regulated by the effector are the same molecule (e.g., O_2 binding affects subsequent O_2 binding).
- Heterotropic allosteric regulator: effector and ligand are different molecules (e.g., H^+ or BPG binding affects O_2 binding).
- Positive allosteric interaction: effector binding increases affinity for ligand.
- Negative allosteric interaction: effector binding decreases affinity for ligand.