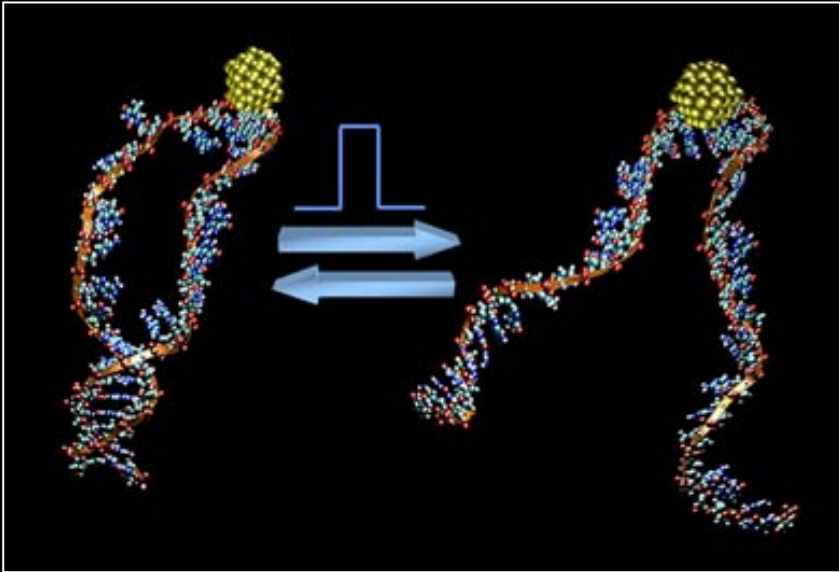
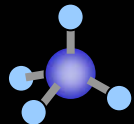


Kimberly Hamad-Schifferli  
Departments of Mechanical  
Engineering and Biological  
Engineering, MIT



Simple  
molecules  
< 1 nm



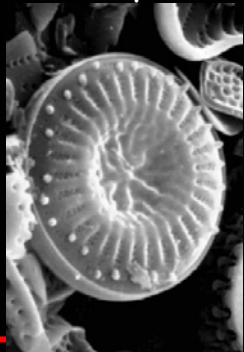
DNA  
proteins  
nm



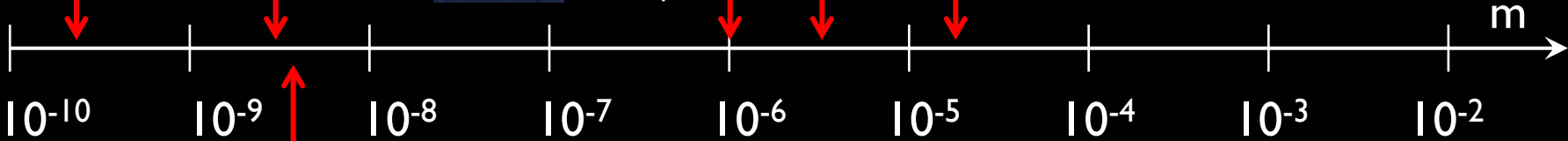
bacteria  
1  $\mu$ m



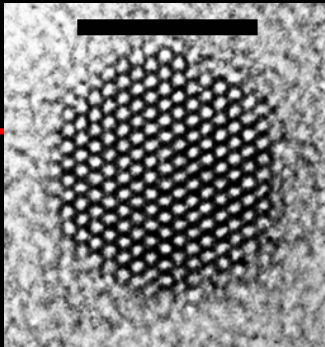
red blood cell  
 $\sim 5 \mu$ m (SEM)



diatom  
30  $\mu$ m

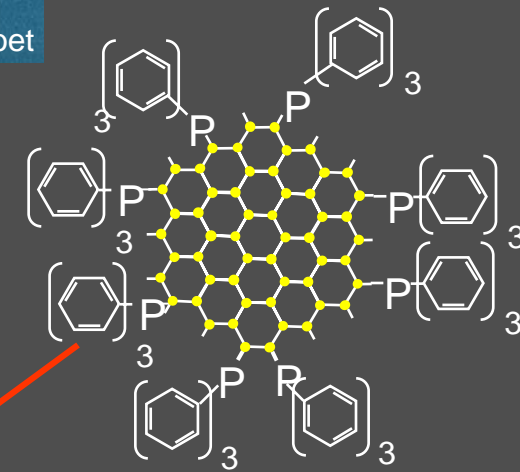
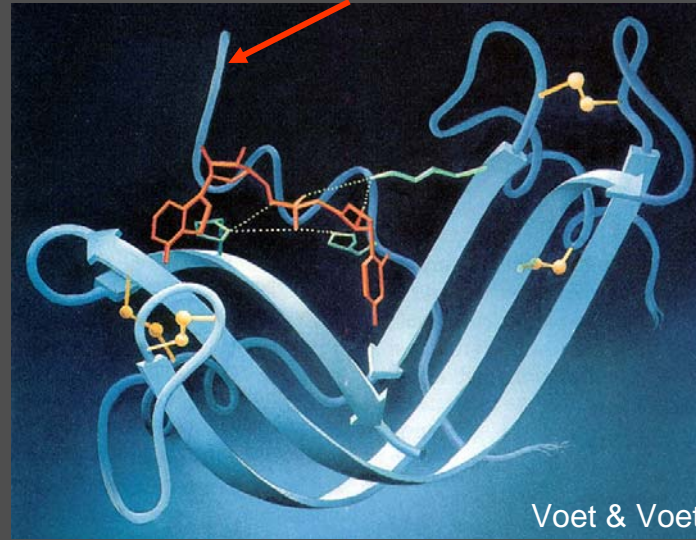
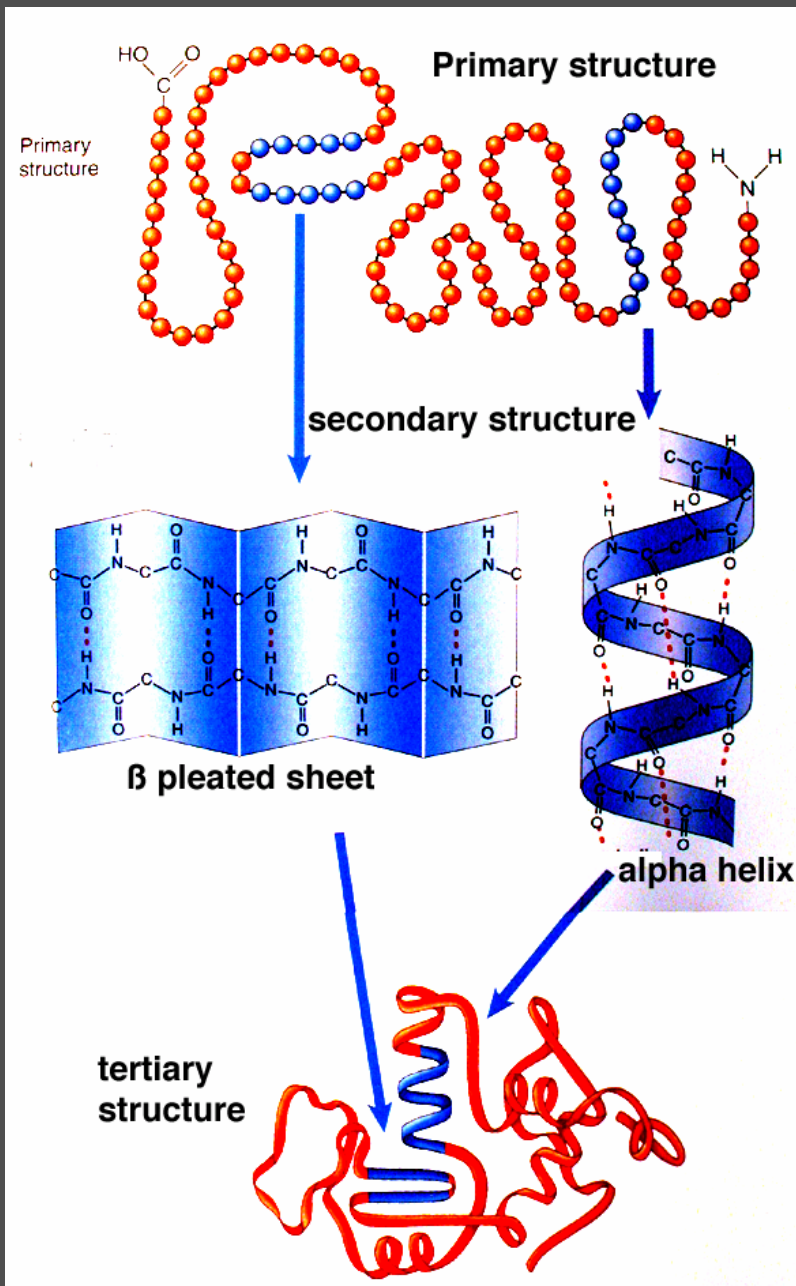


Nanoparticle  
nm



# Nanoparticle linkages to proteins

- Much more complex than with DNA
- More amino acids to interact with
- Where does nanoparticle link?
- Structure key to function

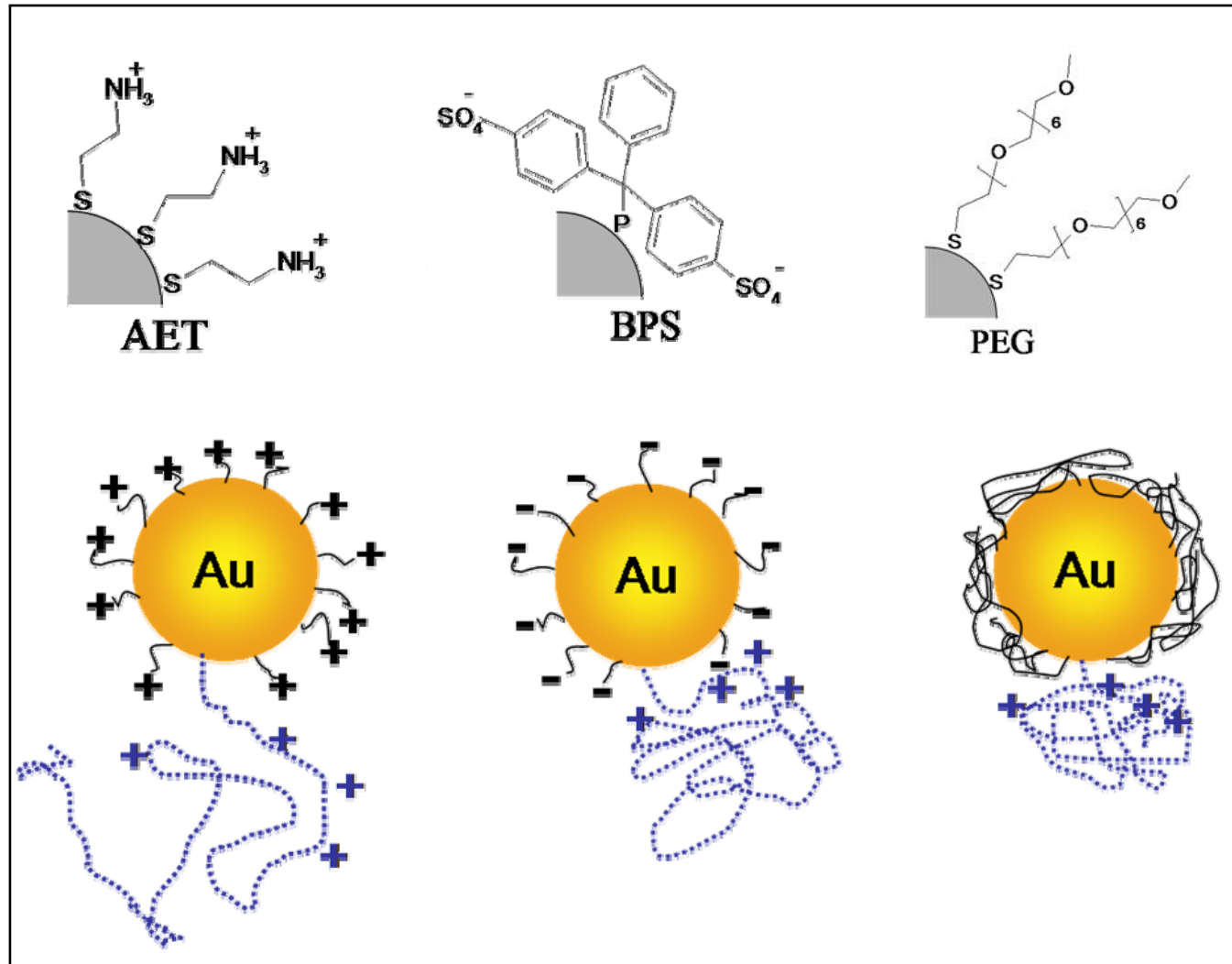
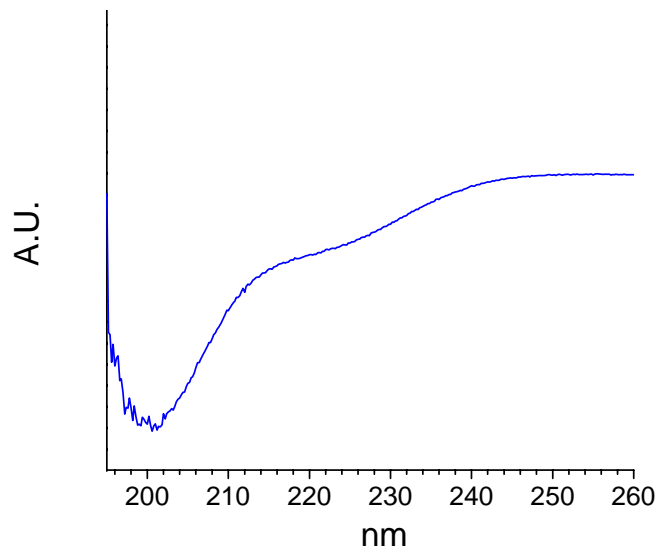
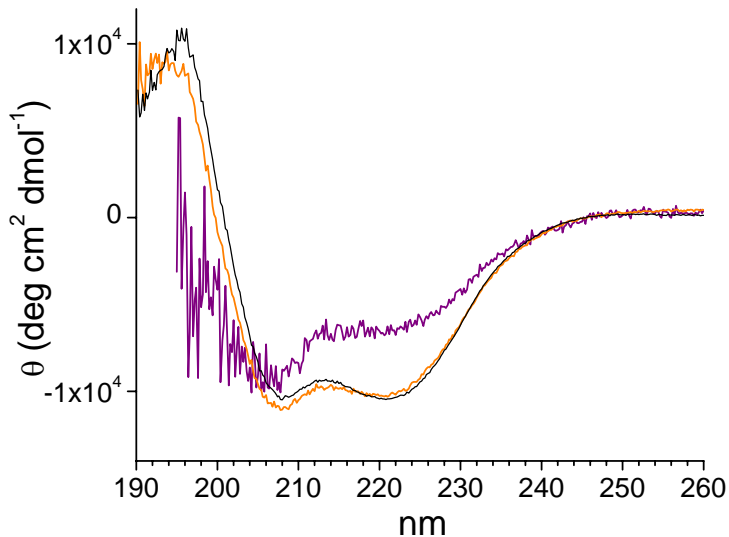


► site specific labeling of protein with NP that preserves protein structure

# NP-cytochrome c interfaces

Site specifically link 1.4nm Au NPs to *Saccharomyces cerevisiae* cytochrome c:

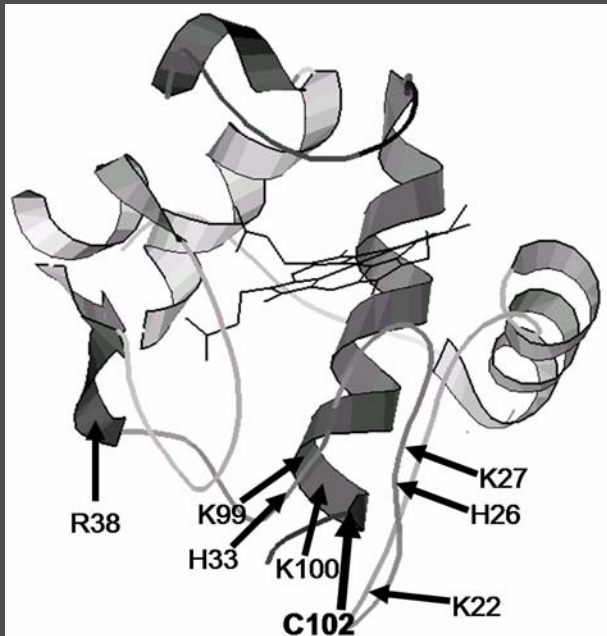
TEFKAGSAKKGATLFFKTR**CLQ**CHTVEKGGPHKVGPNLHGIFGRHSGQAEG  
YSYTDANIKKNVLWDENNMSEYLTNPKKYIPGTKMAFGGLKKEKDRNDLI  
TYLKKACE



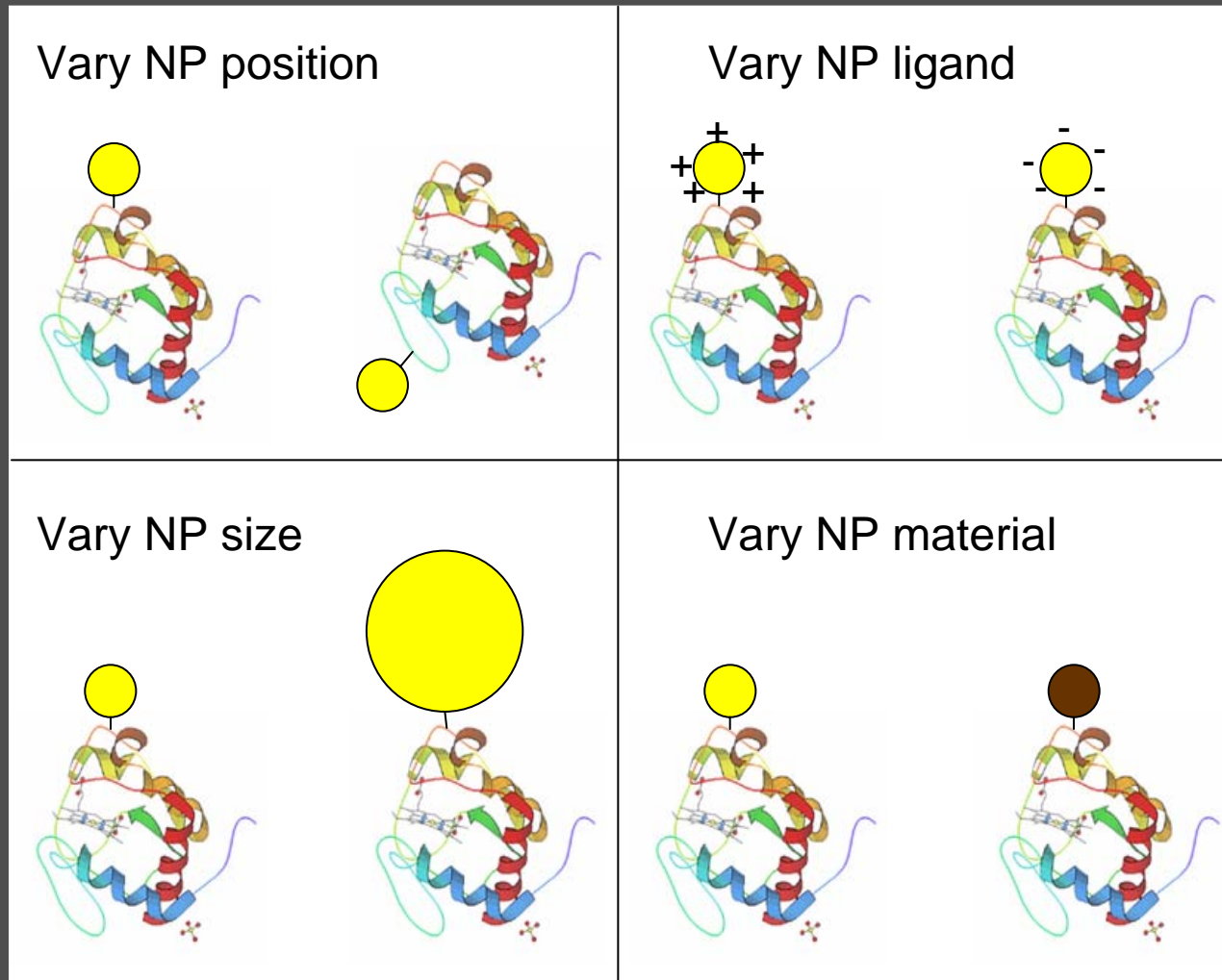
► Charged ligands result in greatest denaturation

# Amino acids in vicinity

- Many charged amino acids closest to linking site

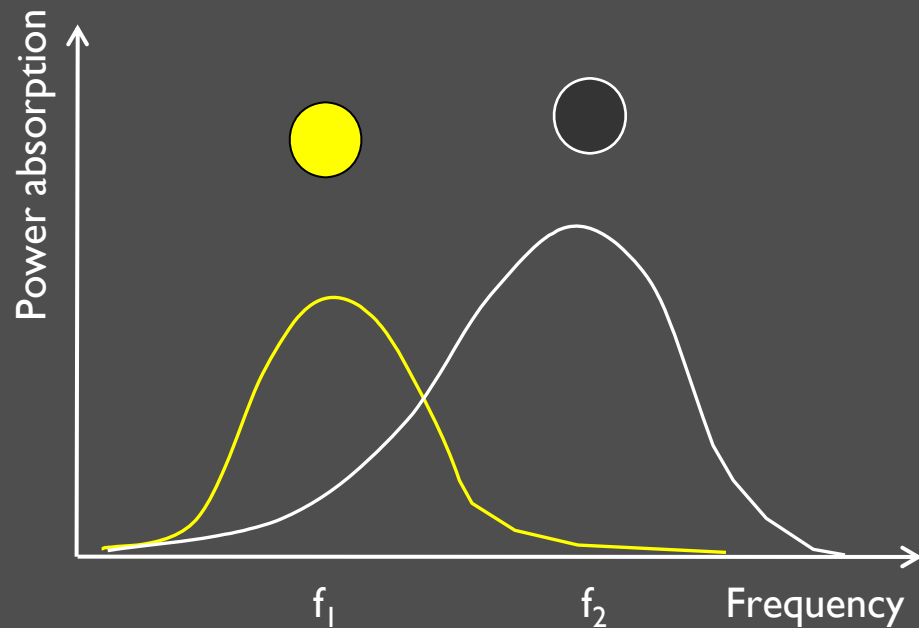
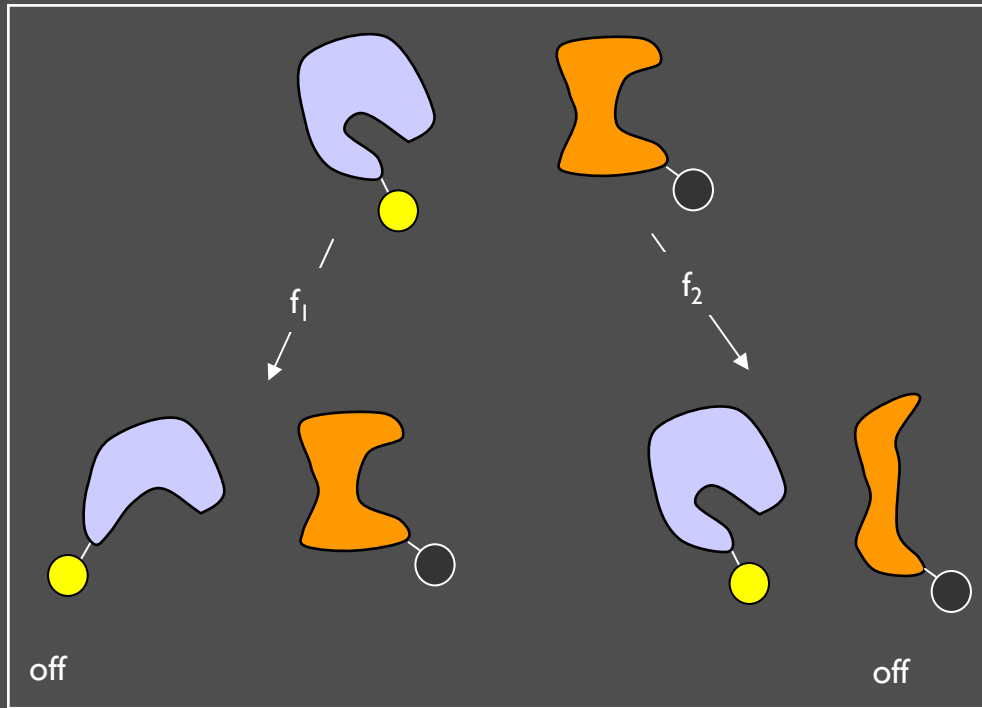


- Systematic study:



► electrostatic interactions with amino acids in local vicinity

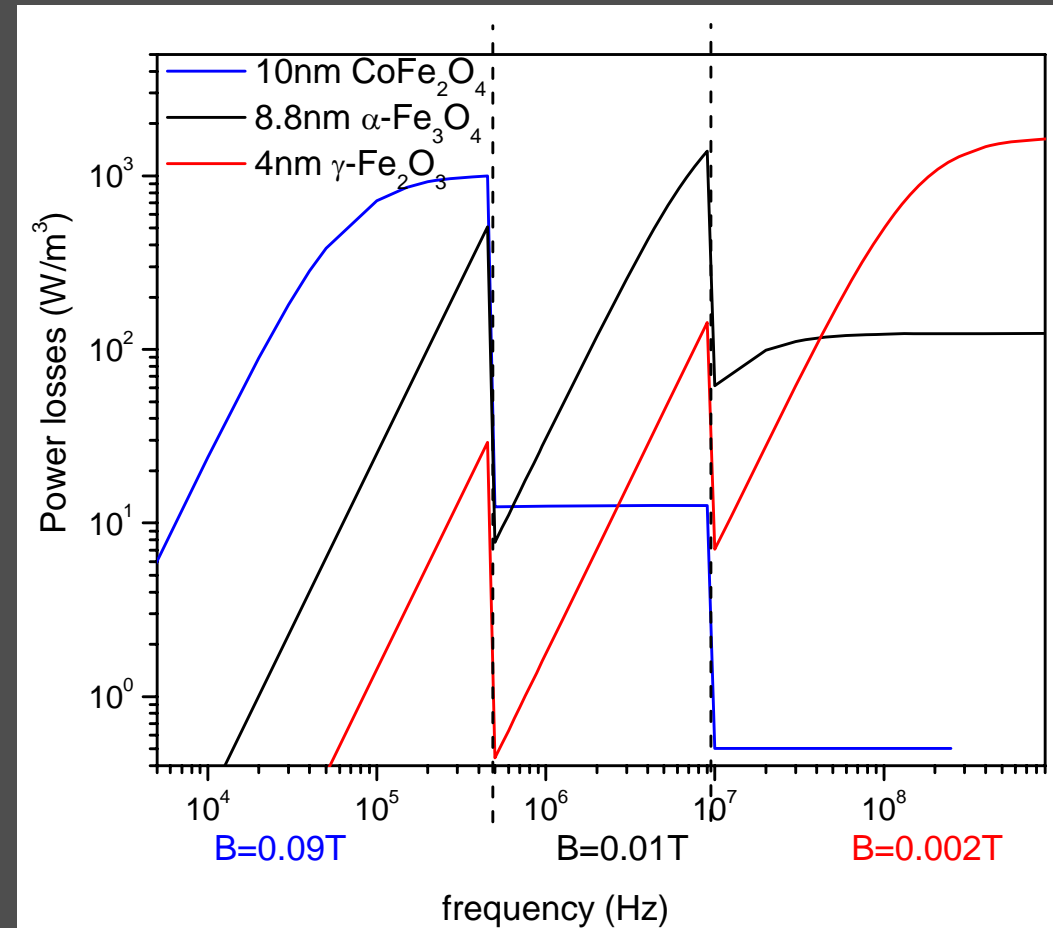
# Orthogonal heating of NPs



A. Wijaya et al., 2006

$$P = f(\text{material properties, } R, H, \omega)$$

3-Variable Tuning = Multiple Control



► independent heating possible