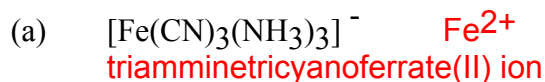
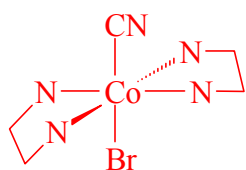
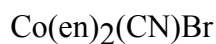


## Review Questions -- Chapter 25

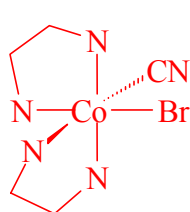
1. For each of the following metal complexes, give the *oxidation state* of the metal and the *complete, systematic name* of the compound or ion.



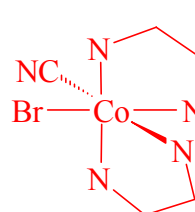
2. Draw *clear, 3-dimensional structures* of all of the isomers, geometric and/or optical, of the following complex. Points will be deducted if the same structure is drawn more than once.



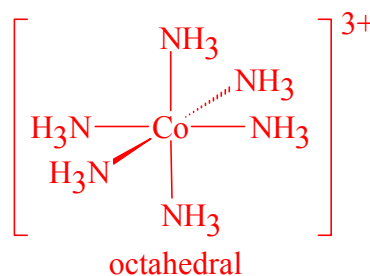
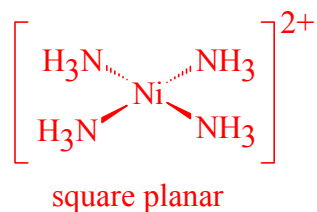
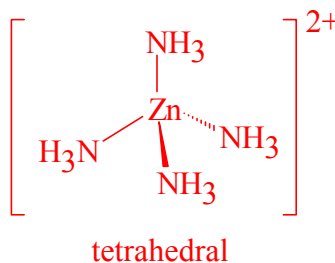
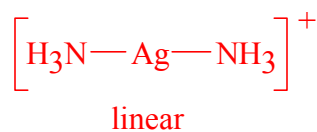
**trans**  
(not optically active)



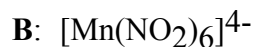
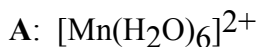
**cis**  
(optically active)



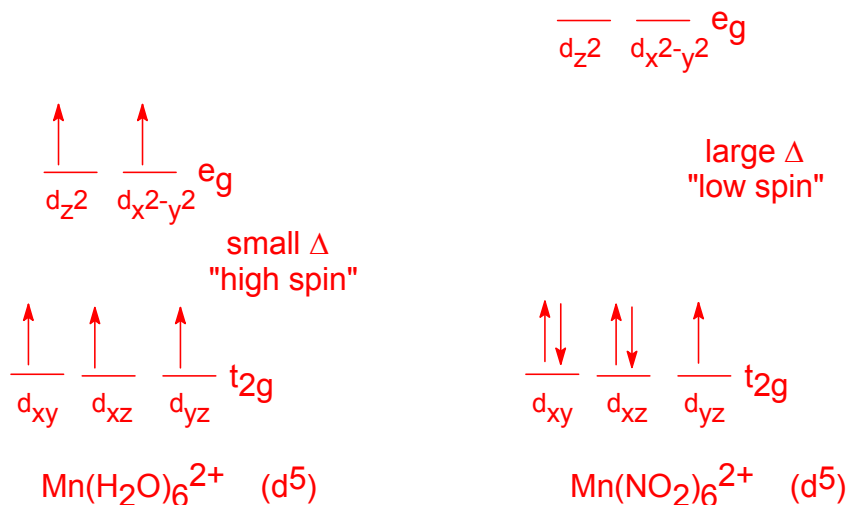
3. When an excess of ammonia ( $\text{NH}_3$ ) is added to aqueous solutions of each of the following metal ions, complexes having various coordination numbers and/or structures are produced. Write the *formula* for each metal- $\text{NH}_3$  complex and clearly draw its *3-dimensional structure*.



4. For each of the following complexes, **A** and **B**, sketch a *properly-labeled d-orbital splitting diagram*. Then answer the questions below in a manner consistent with your diagrams.



The complexes are both  $\text{Mn}^{2+}$ ,  $d^5$  cases but with different ligands. However,  $\text{NO}_2^-$  is a stronger field ligand than is  $\text{H}_2\text{O}$  which leads to a greater d-orbital splitting energy ( $\Delta$ ) in Complex **B**.



(a) How many unpaired electrons does complex **A** have? **5**

(b) Which complex, **A** or **B**, will absorb light of longer wavelength?

Since  $\text{Mn}(\text{H}_2\text{O})_6^{2+}$  has a smaller d-orbital splitting energy ( $\Delta$ ), it should absorb light of lower energy (i.e., longer wavelength) than  $[\text{Mn}(\text{NO}_2)_6]^{4-}$ .