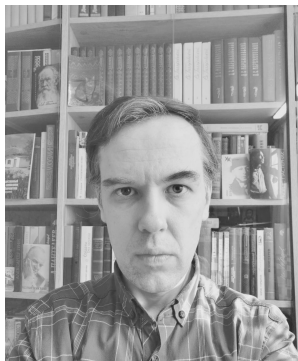


General and Inorganic Chemistry Department,
National Technical University “Kharkiv Polytechnic Institute”

INTRODUCTION TO COORDINATION COMPOUNDS



Responsible Lecturer:
Andrey V. Kramarenko

Lecturer contacts:
andrii.kramarenko@khpі.edu.ua;
avkramar@gmail.com

©Maksym N. Volobuyev
©Andrey V. Kramarenko

Kharkiv 2026



BASIC DEFINITIONS

- The **COORDINATION COMPOUNDS** contain **complex particles**
- The complex particle general formula is as follows:

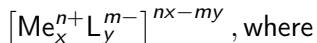


- Me^{n+} is the central ion of metal having unoccupied orbitals;
- L^{m-} is the ligand, the molecule or ion having lone electron pair.
- Hence, the coordination bond is a dative bond



BASIC DEFINITIONS

- The **COORDINATION COMPOUNDS** contain **complex particles**
- The complex particle general formula is as follows:

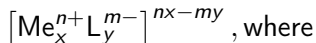


- Me^{n+} is the **CENTRAL ION** of metal having **unoccupied orbitals**;
- L^{m-} is the **LIGAND**, the molecule or ion having **lone electron pair**.
- Hence, the **coordination bond** is a **dative bond**



BASIC DEFINITIONS

- The **COORDINATION COMPOUNDS** contain **complex particles**
- The complex particle general formula is as follows:

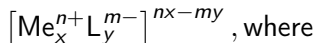


- Me^{n+} is the **CENTRAL ION** of metal having **unoccupied orbitals**;
- L^{m-} is the **LIGAND**, the molecule or ion having **lone electron pair**.
- Hence, the **coordination bond** is a dative bond



BASIC DEFINITIONS

- The **COORDINATION COMPOUNDS** contain **complex particles**
- The complex particle general formula is as follows:

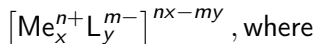


- Me^{n+} is the **CENTRAL ION** of metal having **unoccupied orbitals**;
- L^{m-} is the **LIGAND**, the molecule or ion having **lone electron pair**.
- Hence, the **coordination bond** is a dative bond

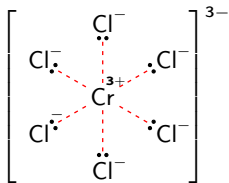


BASIC DEFINITIONS

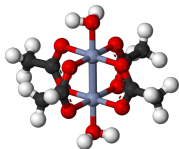
- The **COORDINATION COMPOUNDS** contain **complex particles**
- The complex particle general formula is as follows:



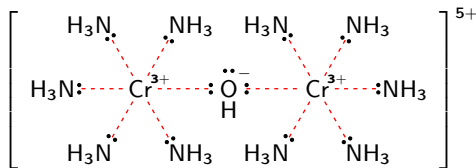
- Me^{n+} is the **CENTRAL ION** of metal having **unoccupied orbitals**;
- L^{m-} is the **LIGAND**, the molecule or ion having **lone electron pair**.
- Hence, the **coordination bond** is a **dative bond**



mononuclear
hexachloridochromate (III)

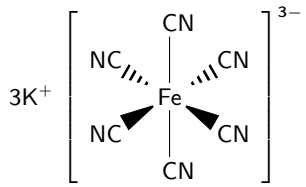
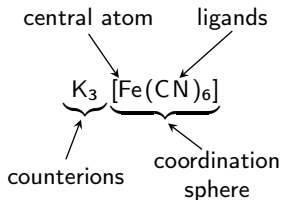


dinuclear cluster
chromium (II)
acetate



dinuclear bridged
 μ -hydroxo-bis(pentaammine)chromium (III)

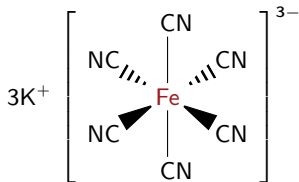
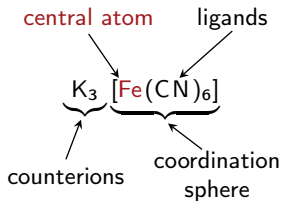




Potassium hexacyanoferrate (III)



The central atom is a **coordination center**

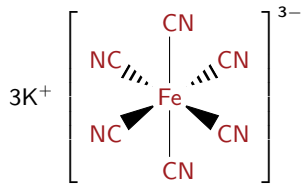
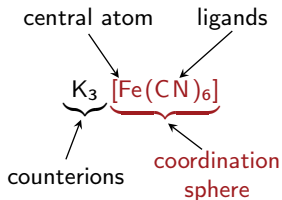


Potassium hexacyanoferrate (III)



COORDINATION COMPOUNDS MAIN COMPONENTS

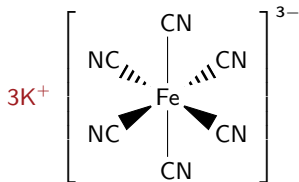
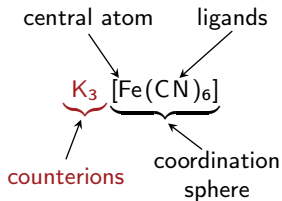
The central atom is a coordination center
 The **coordination sphere** is a metal complex



Potassium hexacyanoferrate (III)



The central atom is a coordination center
 The coordination sphere is a metal complex
 The **outer sphere** contains **counterions**

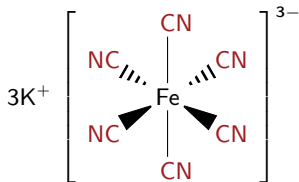
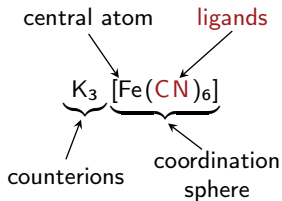


Potassium hexacyanoferrate (III)



COORDINATION COMPOUNDS MAIN COMPONENTS

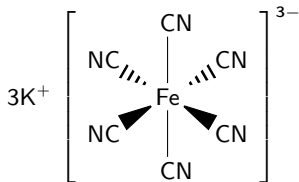
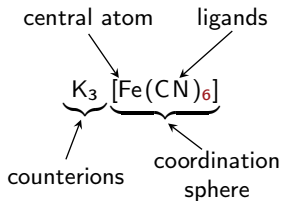
The central atom is a coordination center
The coordination sphere is a metal complex
The outer sphere contains counterions
The **ligands** are ions or molecules



Potassium hexacyanoferrate (III)



The central atom is a coordination center
 The coordination sphere is a metal complex
 The outer sphere contains counterions
 The ligands are ions or molecules
 The **coordination number** (C.N.) is the total number of neighbors of a central atom



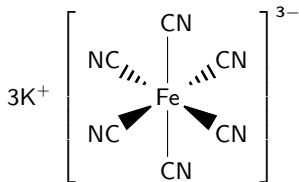
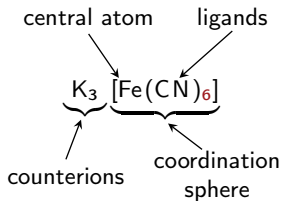
Potassium hexacyanoferrate (III)



COORDINATION COMPOUNDS MAIN COMPONENTS

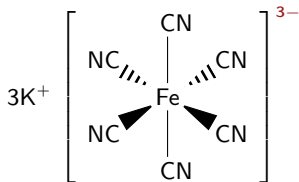
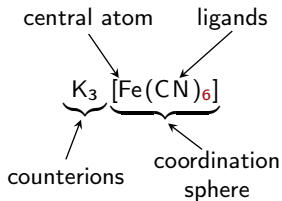
*

The central atom is a coordination center
The coordination sphere is a metal complex
The outer sphere contains counterions
The ligands are ions or molecules
The coordination number (C.N.) is the total number of neighbors of a central atom
C.N. can be from 1 to 12; in the most of the cases is equal to 4 or 6. It very often appears to be **two times more** than total charge of the complex particle



Potassium hexacyanoferrate (III)





Potassium hexacyanoferrate (III)

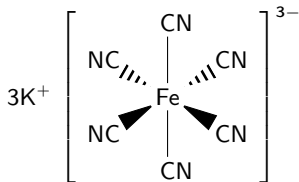
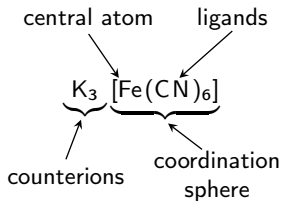
The central atom is a coordination center
 The coordination sphere is a metal complex
 The outer sphere contains counterions
 The ligands are ions or molecules
 The coordination number (C.N.) is the total number of neighbors of a central atom
 C.N. can be from 1 to 12; in the most of the cases is equal to 4 or 6. It very often appears to be two times more than total charge of the complex particle

The **denticity** of the ligand is the number of places that it occupies in the coordination sphere:



COORDINATION COMPOUNDS MAIN COMPONENTS

*

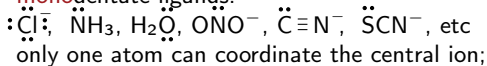


Potassium hexacyanoferrate (III)

The central atom is a coordination center
 The coordination sphere is a metal complex
 The outer sphere contains counterions
 The ligands are ions or molecules
 The coordination number (C.N.) is the total number of neighbors of a central atom
 C.N. can be from 1 to 12; in the most of the cases is equal to 4 or 6. It very often appears to be two times more than total charge of the complex particle

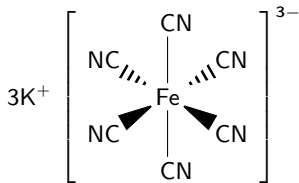
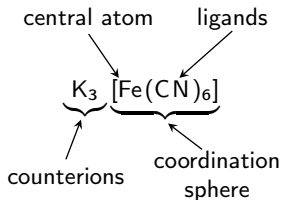
The denticity of the ligand is the number of places that it occupies in the coordination sphere:

monodentate ligands:



COORDINATION COMPOUNDS MAIN COMPONENTS

*

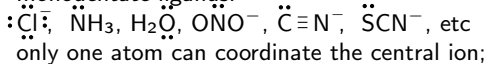


Potassium hexacyanoferrate (III)

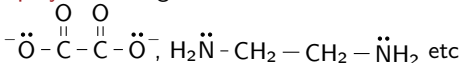
The central atom is a coordination center
 The coordination sphere is a metal complex
 The outer sphere contains counterions
 The ligands are ions or molecules
 The coordination number (C.N.) is the total number of neighbors of a central atom
 C.N. can be from 1 to 12; in the most of the cases is equal to 4 or 6. It very often appears to be two times more than total charge of the complex particle

The denticity of the ligand is the number of places that it occupies in the coordination sphere:

monodentate ligands:



polydentate ligands:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – hexaamminenickel (II) chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – potassium tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
- Oxidation state of the central atom
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
- Oxidation state of the central atom
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $[\text{Co}(\text{NH}_3)_4(\text{H}_2\text{O})_2]\text{Cl}_2$ – **tetraamminedibromocobalt(III)** chloride
 - $[\text{Co}(\text{H}_2\text{O})_4(\text{Cl})_2]\text{Cl}_2$ – **dichlorotetraaquacobalt(III)** chloride
- Oxidation state of the central atom
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaquamonochloridocobalt (III) chloride
- Oxidation state of the central atom
- Structure is connected with the name:



- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- Oxidation state of the central atom
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- Oxidation state of the central atom
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- Oxidation state of the central atom
 - $\text{K}_4[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (II)
 - $\text{K}_3[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (III)
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- **Oxidation state** of the central atom
 - $\text{K}_4[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (II)
 - $\text{K}_3[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (III)
- Structure is connected with the name:



COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- **Oxidation state** of the central atom
 - $\text{K}_4[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (II)
 - $\text{K}_3[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (III)
- Structure is connected with the name:

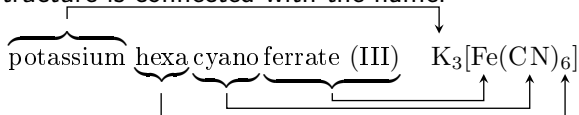


COORDINATION COMPOUNDS NOMENCLATURE

- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaaquamonochloridocobalt (III) chloride
- **Oxidation state** of the central atom
 - $\text{K}_4[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (II)
 - $\text{K}_3[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (III)
- Structure is connected with the name:

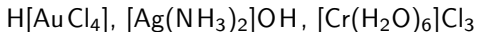


- A **cation** must be called first then anion
 - $[\text{Ni}(\text{NH}_3)_6]\text{Cl}_2$ – **hexaamminenickel (II)** chloride
 - $\text{K}_2[\text{Ni}(\text{CN})_4]$ – **potassium** tetracyanonickelate (II)
- 2 – di, 3 – tri, 4 – tetra, 5 – penta, 6 – hexa, 7 – hepta
- Ligands: should be mentioned in alphabetical order:
 - $\text{K}[\text{Pt}(\text{NH}_3)\text{Br}_5]$ – potassium amminepentabromidoplatinate (IV)
 - $[\text{Co}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$ – pentaquamonochloridocobalt (III) chloride
- **Oxidation state** of the central atom
 - $\text{K}_4[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (II)
 - $\text{K}_3[\text{Fe}(\text{CN})_6]$ – potassium hexacyanoferrate (III)
- Structure is connected with the name:



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:



- By charge: cation, anion, neutral

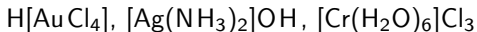


- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:

- aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
- amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
- hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)

- By charge: cation, anion, neutral

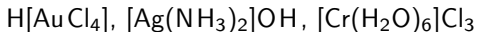


- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:

- aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
- amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
- hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)

- By charge: cation, anion, neutral

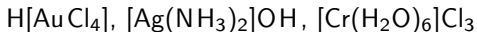


- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:

- aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
- amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
- hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)

- By charge: cation, anion, neutral

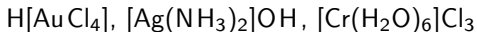


- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:

- aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
- amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
- hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)

- By charge: cation, anion, neutral

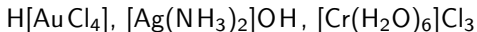


- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



CLASSIFICATION OF COORDINATION COMPOUNDS

- By class: acid, base or salt



- By the nature of ligand:

- aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
- amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
- hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)

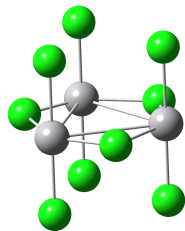
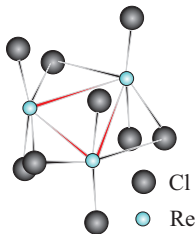
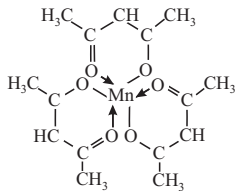
- By charge: cation, anion, neutral



- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



- By class: acid, base or salt
 $\text{H[AuCl}_4\text{]}$, $[\text{Ag}(\text{NH}_3)_2]\text{OH}$, $[\text{Cr}(\text{H}_2\text{O})_6]\text{Cl}_3$
- By the nature of ligand:
 - aquacomplexes ($[\text{Cu}(\text{H}_2\text{O})_4]^{2+}$)
 - amminocomplexes ($[\text{Ag}(\text{NH}_3)_2]^+$)
 - hydroxocomplexes ($[\text{Al}(\text{OH})_6]^{3-}$)
- By charge: cation, anion, neutral
 $[\text{Co}(\text{NH}_3)_6]\text{Cl}_3$, $\text{Li}[\text{AlH}_4]$, $[\text{Co}(\text{NH}_3)_3\text{Cl}_3]$
- By central atoms number: mononuclear, polynuclear (bridged or cluster complexes)



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:

- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:



- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:

$$K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$$

$$K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)} \text{ is recommended by IUPAC}$$

$$K_{st} = 1/K_{ins}$$

- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:

- $$K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$$

- $$K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$$
 is recommended by IUPAC

- $$K_{st} = 1/K_{ins}$$

- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:
 - $K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$
 - $K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$ is recommended by IUPAC
 - $K_{st} = 1/K_{ins}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:
 - $K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$
 - $K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$ is recommended by IUPAC
 - $K_{st} = 1/K_{ins}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:
 - $K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$
 - $K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$ is recommended by IUPAC
 - $K_{st} = 1/K_{ins}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:
 - $K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$
 - $K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$ is recommended by IUPAC
 - $K_{st} = 1/K_{ins}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[Fe(CN)_6] \rightarrow 4K^+ + [Fe(CN)_6]^{4-}$
 - “secondary”: $[Fe(CN)_6]^{4-} \rightleftharpoons Fe^{2+} + 6CN^-$
- Dissociation of the complex is an equilibrium:
 - $K_{ins} = \frac{c(Fe^{2+})c^6(CN^-)}{c([Fe(CN)_6]^{4-})}$
 - $K_{st} = \frac{c([Fe(CN)_6]^{4-})}{c(Fe^{2+})c^6(CN^-)}$ is recommended by IUPAC
 - $K_{st} = 1/K_{ins}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[\text{Fe}(\text{CN})_6] \rightarrow 4\text{K}^+ + [\text{Fe}(\text{CN})_6]^{4-}$
 - “secondary”: $[\text{Fe}(\text{CN})_6]^{4-} \rightleftharpoons \text{Fe}^{2+} + 6\text{CN}^-$
- Dissociation of the complex is an equilibrium:
 - $K_{\text{ins}} = \frac{c(\text{Fe}^{2+})c^6(\text{CN}^-)}{c([\text{Fe}(\text{CN})_6]^{4-})}$
 - $K_{\text{st}} = \frac{c([\text{Fe}(\text{CN})_6]^{4-})}{c(\text{Fe}^{2+})c^6(\text{CN}^-)}$ is recommended by IUPAC
 - $K_{\text{st}} = 1/K_{\text{ins}}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons [\text{Ag}(\text{NH}_3)]^+ + \text{NH}_3, \beta_1 = \frac{c([\text{Ag}(\text{NH}_3)]^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)_2]^+)}$
 - $[\text{Ag}(\text{NH}_3)]^+ \rightleftharpoons \text{Ag}^+ + \text{NH}_3, \beta_2 = \frac{c(\text{Ag}^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)]^+)}$
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons \text{Ag}^+ + 2\text{NH}_3, K_{\text{ins}} = \beta_1 \cdot \beta_2$



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[\text{Fe}(\text{CN})_6] \rightarrow 4\text{K}^+ + [\text{Fe}(\text{CN})_6]^{4-}$
 - “secondary”: $[\text{Fe}(\text{CN})_6]^{4-} \rightleftharpoons \text{Fe}^{2+} + 6\text{CN}^-$
- Dissociation of the complex is an equilibrium:
 - $K_{\text{ins}} = \frac{c(\text{Fe}^{2+})c^6(\text{CN}^-)}{c([\text{Fe}(\text{CN})_6]^{4-})}$
 - $K_{\text{st}} = \frac{c([\text{Fe}(\text{CN})_6]^{4-})}{c(\text{Fe}^{2+})c^6(\text{CN}^-)}$ is recommended by IUPAC
 - $K_{\text{st}} = 1/K_{\text{ins}}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons [\text{Ag}(\text{NH}_3)]^+ + \text{NH}_3, \beta_1 = \frac{c([\text{Ag}(\text{NH}_3)]^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)_2]^+)}$
 - $[\text{Ag}(\text{NH}_3)]^+ \rightleftharpoons \text{Ag}^+ + \text{NH}_3, \beta_2 = \frac{c(\text{Ag}^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)]^+)}$
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons \text{Ag}^+ + 2\text{NH}_3, K_{\text{ins}} = \beta_1 \cdot \beta_2$



DISSOCIATION OF COMPLEX COMPOUNDS

- Dissociation occurs in two stages:
 - “primary”: $K_4[\text{Fe}(\text{CN})_6] \rightarrow 4\text{K}^+ + [\text{Fe}(\text{CN})_6]^{4-}$
 - “secondary”: $[\text{Fe}(\text{CN})_6]^{4-} \rightleftharpoons \text{Fe}^{2+} + 6\text{CN}^-$
- Dissociation of the complex is an equilibrium:
 - $K_{\text{ins}} = \frac{c(\text{Fe}^{2+})c^6(\text{CN}^-)}{c([\text{Fe}(\text{CN})_6]^{4-})}$
 - $K_{\text{st}} = \frac{c([\text{Fe}(\text{CN})_6]^{4-})}{c(\text{Fe}^{2+})c^6(\text{CN}^-)}$ is recommended by IUPAC
 - $K_{\text{st}} = 1/K_{\text{ins}}$
- The more the K_{st} the more stable complex is
- Step-by-step dissociation
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons [\text{Ag}(\text{NH}_3)]^+ + \text{NH}_3, \beta_1 = \frac{c([\text{Ag}(\text{NH}_3)]^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)_2]^+)}$
 - $[\text{Ag}(\text{NH}_3)]^+ \rightleftharpoons \text{Ag}^+ + \text{NH}_3, \beta_2 = \frac{c(\text{Ag}^+)c(\text{NH}_3)}{c([\text{Ag}(\text{NH}_3)]^+)}$
 - $[\text{Ag}(\text{NH}_3)_2]^+ \rightleftharpoons \text{Ag}^+ + 2\text{NH}_3, K_{\text{ins}} = \beta_1 \cdot \beta_2$



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - Fe^{2+} gives a precipitate with K_2S that dissolves in excess amount of precipitant
 - Fe^{3+} gives a precipitate with K_2S that dissolves in excess amount of precipitant
 - Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:



- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:



- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation



- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation



COORDINATION COMPOUNDS: SOME REACTIONS

- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AgBr}, \text{AgI} + 2\text{NH}_3 \rightarrow$



- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AgBr}, \text{AgI} + 2\text{NH}_3 \rightarrow$
 - $\text{AgHal} + 2\text{Na}_2\text{S}_2\text{O}_3 = \text{Na}_3[\text{Ag}(\text{S}_2\text{O}_3)_2] + \text{NaHal}$



- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AgBr}, \text{AgI} + 2\text{NH}_3 \rightarrow$
 - $\text{AgHal} + 2\text{Na}_2\text{S}_2\text{O}_3 = \text{Na}_3[\text{Ag}(\text{S}_2\text{O}_3)_2] + \text{NaHal}$



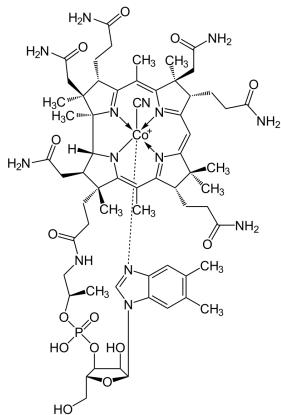
- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AgBr}, \text{AgI} + 2\text{NH}_3 \rightarrow$
 - $\text{AgHal} + 2\text{Na}_2\text{S}_2\text{O}_3 = \text{Na}_3[\text{Ag}(\text{S}_2\text{O}_3)_2] + \text{NaHal}$



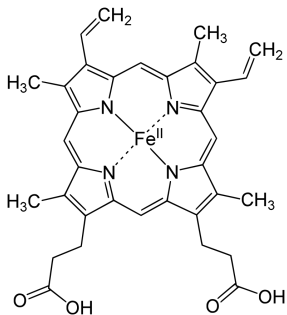
- Formation of the complexes
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AuCN} + \text{KCN} = \text{K}[\text{Au}(\text{CN})_2]$
- Reaction to Fe^{2+} and Fe^{3+} cations:
 - $\text{FeSO}_4 + \text{K}_3[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Turnbull's blue}} \downarrow + 2\text{K}_2\text{SO}_4$
 - $\text{FeCl}_3 + \text{K}_4[\text{Fe}(\text{CN})_6] = \underbrace{\text{KFe}[\text{Fe}(\text{CN})_6]}_{\text{Prussian blue}} \downarrow + 3\text{KCl}$
- Bi^{3+} gives a precipitate with KI that dissolves in excess amount of precipitant
 - $\text{BiCl}_3 + 3\text{KI} = \text{BiI}_3 \downarrow + 3\text{KCl}$
 - $\text{BiI}_3 + \text{KI} = \text{K}[\text{BiI}_4]$
- Silver halogenides separation
 - $\text{AgCl} + 2\text{NH}_3 = [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 - $\text{AgBr}, \text{AgI} + 2\text{NH}_3 \nrightarrow$
 - $\text{AgHal} + 2\text{Na}_2\text{S}_2\text{O}_3 = \text{Na}_3[\text{Ag}(\text{S}_2\text{O}_3)_2] + \text{NaHal}$



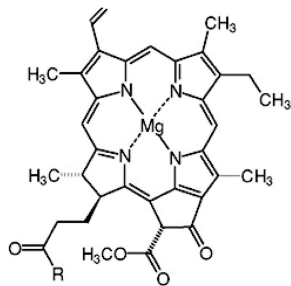
FAMOUS COMPLEXES



Cyanocobalamin (B12)



Hemoglobine heme



Chlorophyll



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_5(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S → 2Na⁺ + S²⁻
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	–	–
c_{rctd}	x	–	–
c_{frmd}	–	x	4x
c_{eq}	0.05 – x	x	4x

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_5$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_5(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from $\text{Na}_2\text{S} \rightarrow 2\text{Na}^+ + \text{S}^{2-}$
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	–	–
c_{rctd}	x	–	–
c_{frmd}	–	x	4x
c_{eq}	0.05 – x	x	4x

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_5$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_5(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S → 2Na⁺ + S²⁻
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	–	–
c_{rctd}	x	–	–
c_{frmd}	–	x	4x
c_{eq}	0.05 – x	x	4x

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_5$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_5(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S $\rightarrow 2\text{Na}^+ + \text{S}^{2-}$
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	–	–
c_{rctd}	x	–	–
c_{frmd}	–	x	4x
c_{eq}	0.05 – x	x	4x

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_5$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_5(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S $\rightarrow 2\text{Na}^+ + \text{S}^{2-}$
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	–	–
c_{rctd}	x	–	–
c_{frmd}	–	x	4x
c_{eq}	0.05 – x	x	4x

$$K_{\text{st}}([\text{Cd}(\text{CN})_4]^{2-}) = 1.3 \cdot 10^{17}$$

$$\frac{c([\text{Cd}(\text{CN})_4]^{2-})}{c(\text{Cd}^{2+})c^4(\text{CN}^-)} = 1.3 \cdot 10^{17}$$

$$\frac{0.05-x}{x \cdot (4x)^4} = 1.3 \cdot 10^{17}$$

If $x \ll 0,05$ then

$$256x^5 = 3.9 \cdot 10^{-19}$$

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_5$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_s(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S $\rightarrow 2\text{Na}^+ + \text{S}^{2-}$
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

$$K_{\text{st}}([\text{Cd}(\text{CN})_4]^{2-}) = 1.3 \cdot 10^{17}$$

$$\frac{c([\text{Cd}(\text{CN})_4]^{2-})}{c(\text{Cd}^{2+})c^4(\text{CN}^-)} = 1.3 \cdot 10^{17}$$

$$\frac{0.05 - x}{x \cdot (4x)^4} = 1.3 \cdot 10^{17}$$

If $x \ll 0,05$ then

$$256x^5 = 3.9 \cdot 10^{-19}$$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	—	—
c_{rctd}	x	—	—
c_{frmd}	—	x	$4x$
c_{eq}	$0.05 - x$	x	$4x$

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_s$
- The conclusion is that precipitate will appear



TASK EXAMPLE

- Does the CdS precipitate will form, if some volume of 0.1 solution of Na₂S will be added to the equal volume of 0.1 K₂[Cd(CN)₄] solution?
- Condition of sedimentation: $c(\text{Cd}^{2+})c(\text{S}^{2-}) > K_s(\text{CdS}) = 8 \cdot 10^{-27}$
- It's easy to determine $c(\text{S}^{2-})$ from Na₂S $\rightarrow 2\text{Na}^+ + \text{S}^{2-}$
 $c(\text{S}^{2-}) = 1/2 c_{\text{st}}(\text{Na}_2\text{S}) = 0.05$ (because of mixing dilution)
- $c(\text{Cd}^{2+})$ can be found using scheme $[\text{Cd}(\text{CN})_4]^{2-} \rightleftharpoons \text{Cd}^{2+} + 4\text{CN}^-$

$$K_{\text{st}}([\text{Cd}(\text{CN})_4]^{2-}) = 1.3 \cdot 10^{17}$$

$$\frac{c([\text{Cd}(\text{CN})_4]^{2-})}{c(\text{Cd}^{2+})c^4(\text{CN}^-)} = 1.3 \cdot 10^{17}$$

$$\frac{0.05 - x}{x \cdot (4x)^4} = 1.3 \cdot 10^{17}$$

If $x \ll 0,05$ then

$$256x^5 = 3.9 \cdot 10^{-19}$$

	$[\text{Cd}(\text{CN})_4]^{2-}$	Cd^{2+}	CN^-
c_{st}	0.05	—	—
c_{rctd}	x	—	—
c_{frmd}	—	x	4x
c_{eq}	0.05 - x	x	4x

- $x = c(\text{Cd}^{2+}) = 6.9 \cdot 10^{-5}$
- $c(\text{Cd}^{2+})c(\text{S}^{2-}) = 6.9 \cdot 10^{-5} \cdot 0.05 = 3.5 \cdot 10^{-6} > K_s$
- The conclusion is that precipitate will appear

