

CAIE Chemistry A-level Topic 28 - An Introduction to the Chemistry of Transition Elements (A level only)

Flashcards

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What is a transition element?







What is a transition element?

A d-block element that can form one or more stable ions with an incomplete d-subshell.







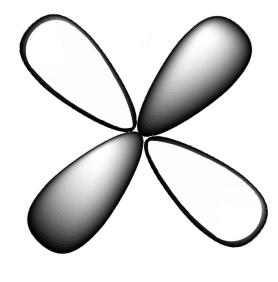
Sketch the shape of a d-orbital







Sketch the shape of a d-orbital









Write the electron configuration for iron and iron(III) ions







Write the electron configuration for iron and iron(III) ions

Fe: 1s²2s²2p⁶3s²3p⁶3d⁶4s²

Fe³⁺: 1s²2s²2p⁶3s²3p⁶3d⁵







Write the electron configurations for chromium and copper







Write the electron configurations for chromium and copper

These are exceptions to the rules for filling up subshells. The 4s subshell holds only one electron in order to give the most stable arrangement.

- Chromium: 1s²2s²2p⁶3s²3p⁶3d⁵4s¹
- Copper: 1s²2s²2p⁶3s²3p⁶3d¹⁰4s¹







Describe and explain how the melting points of transition metals are different to those of typical s-block metals, such as calcium







Describe and explain how the melting points of transition metals are different to those of typical s-block metals, such as calcium

Transition metals have higher melting points than typical s-block metals.

This is because transition metals have the extra 3d electrons which increase the strength of the metallic bonding meaning more energy is required to overcome these bonds when melting the metal.







Describe how the densities of transition metals are different to those of a typical s-block metal, such as calcium







Describe how the densities of transition metals are different to those of a typical s-block metal, such as calcium

Transition metals have greater densities than s-block metals.







Explain why the densities of transition metals are greater than those of s-block metals, such as calcium







Explain why the densities of transition metals are greater than those of s-block metals, such as calcium

- Nuclear charge increases across the period while electron shielding remains the same. This causes the outer shell electrons to be drawn inwards, decreasing the atomic radius.
- As a result, the densities of transition metals are greater than s-block metals because the atoms are smaller so they can pack more closely together and atomic mass increases across the period.







Why are transition elements used as catalysts?







Why are transition elements used as catalysts?

Transition elements are stable in variable oxidation states which means they can easily give/take electrons to/from other molecules.







Why do transition metals have variable oxidation states?







Why do transition metals have variable oxidation states?

Electrons in the 4s and 3d orbitals have very similar energies. This means a relatively similar amount of energy is required to gain/lose a different number of electrons.







How can you predict the likely oxidation states of a transition metal by using its electron configuration?







How can you predict the likely oxidation states of a transition metal by using its electron configuration?

- Most transition metal ions have a 2+ oxidation state due to presence of the 4s electrons.
- To work out the maximum oxidation state: maximum oxidation state = number of 4s electrons + the number of unpaired 3d electrons (copper is an exception to this rule).
- Other oxidation states between +2 and the maximum exist but are much more rare.







What is a ligand? Include the meanings of the terms monodentate, bidentate and polydentate







What is a ligand? Include the meanings of the terms monodentate, bidentate and polydentate

A species that has a lone pair of electrons that forms a dative covalent bond with a central metal atom/ion.

- Monodentate forms 1 coordinate bond
- Bidentate forms 2 coordinate bonds
- Polydentate forms more than 2 coordinate bonds







What is a complex ion?







What is a complex ion?

A molecule containing a central metal ion surrounded by one or more ligands (bonded by coordinate bonds).







What is ligand substitution?







What is ligand substitution?

A reaction in which one ligand in a complex ion is replaced by another.







How does hexaaquacopper(II) form?







How does hexaaquacopper(II) form?

When a copper(II) compound dissolves to release Cu²⁺ ions, water molecules form dative covalent bonds with the metal ion. A complex ion is formed:

$$Cu^{2+}$$
 + $6H_2O$ → $[Cu(H_2O)_6]^{2+}$

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Write an equation for the ligand substitution reaction between ammonia and [Cu(H₂O)]²⁺ ions. Include any colours.







Write an equation for the ligand substitution reaction between ammonia and $[Cu(H_2O)]^{2+}$ ions. Include any colours

$$[Cu(H_2O)_6]^{2+} + 4NH_3 \approx [Cu(NH_3)_4(H_2O)_2]^{2+} + 4H_2O$$

Pale blue solution

Dark blue solution

A pale blue precipitate, $Cu(OH)_2$, initially forms. It dissolves on the addition of excess ammonia.





Describe and explain the shape of $[Cu(NH_3)_4(H_2O)_2]^{2+}$







Describe and explain the shape of $[Cu(NH_3)_4(H_2O)_2]^{2+}$

Distorted octahedral because the Cu-O bonds are longer than the Cu-N bonds.







Write an equation for the ligand substitution reaction between chloride ions and $[Cu(H_2O)]^{2+}$ ions. Include any colours.







Write an equation for the ligand substitution reaction between chloride ions and $[Cu(H_2O)]^{2+}$ ions. Include any colours.

 $[Cu(H_2O)_6]^{2+} + 4CI^- = [CuCI_4]^{2-} + 6H_2O$

Pale blue solution

Yellow solution







Name the shape of $[CuCl_4]^{2-1}$







Name the shape of $[CuCl_4]^{2-1}$

Tetrahedral

Cl⁻ is larger with stronger repulsion than water ligands so only 4 Cl⁻ ligands surround the copper ion.







Write an equation for the reaction between hexaaquacopper(II) and hydroxide ions. Include any colours.







Write an equation for the reaction between hexaaquacopper(II) and hydroxide ions. Include any colours.

$$[Cu(H_2O)_6]^{2+} + 2OH^- \rightarrow [Cu(H_2O)_4(OH)_2] + 2H_2O^-$$

Blue solution

Blue precipitate







How does hexaaquacobalt(II) form?







How does hexaaquacobalt(II) form?

When a cobalt(II) compound dissolves and releases Co²⁺ ions, water molecules form dative covalent bonds with the metal ion. This forms a complex ion:

$$\mathrm{Co}^{2+} + 6\mathrm{H}_{2}\mathrm{O} \rightarrow [\mathrm{Co}(\mathrm{H}_{2}\mathrm{O})_{6}]^{2+}$$

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Write an equation for the ligand substitution reaction between ammonia and $[Co(H_2O)_6]^{2+}$. Include any colours.







Write an equation for the ligand substitution reaction between ammonia and $[Co(H_2O)_6]^{2+}$. Include any colours.

$$[CO(H_2O)_6]^{2+} + 2NH_3 \rightarrow [CO(H_2O)_4(OH)_2]^{2+} + 2NH_4^+$$

Pink solution Blue precipitate

In excess ammonia:

$$[Co(H_2O)_6]^{2+} + 6NH_3 \rightarrow [Co(NH_3)_6]^{2+} + 6H_2O$$

Pink solution

Yellow brown solution







Write an equation for the ligand substitution reaction between chloride ions and $[Co(H_2O)_6]^{2+}$. Include any colours.







Write an equation for the ligand substitution reaction between chloride ions and $[Co(H_2O)_6]^{2+}$. Include any colours.

 $[Co(H_2O)_6]^{2+} + 4CI^- \rightarrow [CoCI_4]^{2-} + 6H_2O$

Pink solution

Blue solution







Write an equation for the reaction between hexaaquacobalt(II) and hydroxide ions. Include any colours.







Write an equation for the reaction between hexaaquacobalt(II) and hydroxide ions. Include any colours.

$$[Co(H_2O)_6]^{2+}_{(aq)} + 2OH_{(aq)}^{-} \rightarrow [Co(H_2O)_4(OH)_2]_{(s)} + 2H_2O_{(l)}$$

Pink solution Blue precipitate







What is meant by coordination number?







What is meant by *coordination number*?

The number of coordinate bonds formed between the ligands and the central metal ion/atom.







What is the shape and bond angle of a complex with a coordination number of 2?

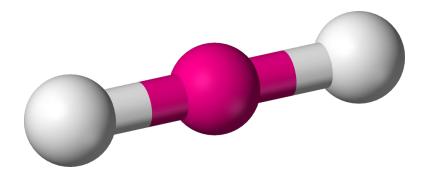






What is the shape and bond angle of a complex with a coordination number of 2?

Linear, 180°









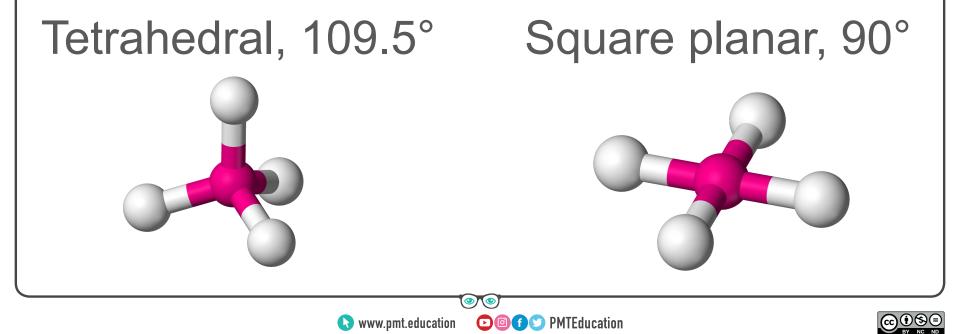
What two shapes and bond angles can complexes with a coordination number of 4 have?







What two shapes and bond angles can complexes with a coordination number of 4 have?





What is the shape and bond angle of a complex ion with a coordination number of 6?

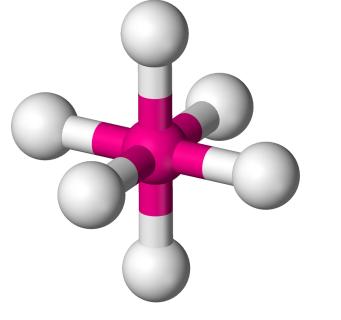






What is the shape bond angle of a complex ion with a coordination number of 6?

Octahedral, 90°









What are the half equations for the reactions which take place when potassium manganate(VII) reacts with an iron(II) salt?







What are the half equations for the reactions which take place when potassium manganate(VII) reacts with an iron(II) salt?

$$MnO_{4 (aq)}^{-} + 8H_{(aq)}^{+} + 5e^{-} \rightarrow Mn^{2+}_{(aq)}^{+} + 4H_{2}O_{(I)}$$

$$Fe^{2+}_{(aq)} \rightarrow Fe^{3+}_{(aq)} + e^{-}$$

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What is the redox equation for the overall reaction which takes place between potassium manganate(VII) and an iron(II) salt?







What is the redox equation for the overall reaction which takes place between potassium manganate(VII) and an iron(II) salt?

Balance the electrons in the half equations and then combine the half equations so that the number of electrons cancel:

$$MnO_{4}^{-} + 8H^{+} + 5Fe^{2+} \rightarrow Mn^{2+} + 5Fe^{3+} + 4H_{2}O^{-}$$







Construct the half equation for the reduction of $Cr_2O_7^{2-}$ ions to Cr^{3+} ions







Construct the half equation for the reduction of $Cr_2O_7^{2-}$ ions to Cr^{3+} ions

$Cr_2O_7^{2-} + 14H^+ + 6e^- \rightarrow 2Cr^{3+} + 7H_2O$







How do you predict which half-cell is being oxidised and which is being reduced?







How do you predict which half-cell is being oxidised and which is being reduced?

- The more negative the E^θ value, the greater the tendency for the system to be oxidised.
- The more positive the E^θ value, the greater the tendency for the system to be reduced.
- Hence the half-cell with the more positive E^θ is oxidised and the half-cell with the more negative E^θ is reduced.

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How do you predict the feasibility of a redox reaction?







How do you predict the feasibility of a redox reaction?

- The reaction is feasible if the oxidising agent has a lower standard cell potential than the reducing agent.
- The greater the difference in E^θ value, the more likely the reaction is to occur.







It has been predicted using standard electrode potentials that a redox reaction is feasible. Why might this reaction not occur spontaneously?







It has been predicted using standard electrode potentials that a redox reaction is feasible. Why might this reaction not occur spontaneously?

- Non-standard conditions
- Ambient energy of the system is lower than the activation energy.







What are degenerate orbitals?







What are degenerate orbitals?

Orbitals with the same energy level.







How are degenerate d-orbitals split into two energy levels in octahedral complexes?







How are degenerate d-orbitals split into two energy levels in octahedral complexes?

- The ligand electrons repel the d-orbital electrons of the transition metal. This raises the energy of the d orbitals.
- These orbitals are split into two groups, one with a slightly higher energy than the other.
- The group with the higher energy has two d-orbitals while the other group has three d-orbitals.







How are degenerate d-orbitals split into two energy levels in tetrahedral complexes?







How are degenerate d-orbitals split into two energy levels in tetrahedral complexes?

- Ligand electrons repel d-orbital electrons of the transition metal. This raises the energy level of the d-orbitals.
- These orbitals are split into two groups, one with a slightly higher energy than the other.
- The group with a higher energy contains three d-orbitals while the other group contains two d-orbitals.
- This is the opposite way round to octahedral complexes.







How does the quantity of energy absorbed due to the split d-orbitals correspond with the colour of the transition metal ion complex?

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How does the quantity of energy absorbed due to the split d-orbitals correspond with the colour of the transition metal ion complex?

- When white light is passed through the complex, some energy is absorbed and used to raise an electron from the lower d-orbital energy level to the higher d-orbital energy level.
- The equivalent wavelength/frequency for this quantity of energy is the wavelength/frequency of light that is absorbed.
- The transition metal will be seen as the complementary colour to the colour that was absorbed (e.g. blue will be seen when red is absorbed).







What factors affect the colour of transition metal ion complexes?







What factors affect the colour of transition metal ion complexes?

- The nature of the ligand.
- The oxidation state of the metal.
- The coordination number of the transition metal ion.







How does the nature of the ligand affect the transition metal ion complex colour?







How does the nature of the ligand affect the transition metal ion complex colour?

- If the ligands have a strong electric field, there will be a larger gap between d-orbitals.
- If the ligands have a weak electric field, there will be a smaller gap between d-orbitals.

The larger the gap, the more energy is absorbed which means the corresponding wavelength of light gets smaller and hence, if the light is absorbed from the red end of the spectrum, the colour will move away from the red end of the spectrum.







How does the oxidation state of the metal affect the colour of the transition metal ion complex?







How does the oxidation state of the metal affect the colour of the transition metal ion complex?

- As the oxidation state of the metal increases, the amount of d-orbital splitting increases.
- This therefore affects the amount of energy absorbed and hence the corresponding wavelength of light.







How does the coordination number of the ion affect the colour of the transition metal ion complex?







How does the coordination of the ion affect the colour of the transition metal ion complex?

- For an octahedral molecule, splitting is greater than in a tetrahedral molecule.
- Greater splitting leads to more energy being absorbed, which in turn means the wavelength of light is smaller.

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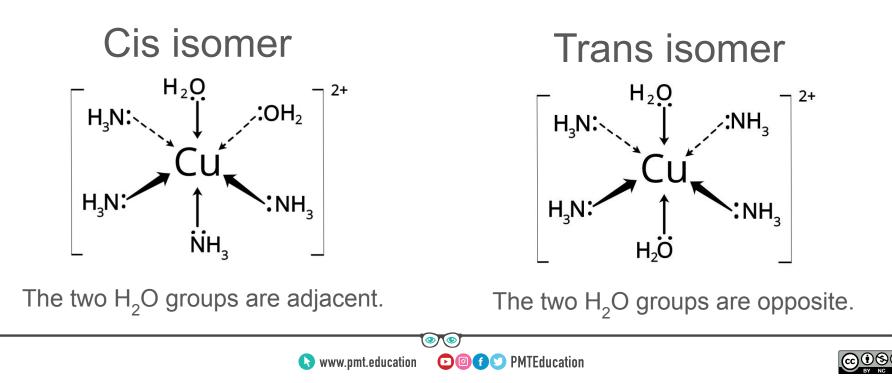
Use diagrams of [Cu(NH₃)₄(H₂O)]²⁺ to show how complex ions can show cis/trans isomerism







Use diagrams of $[Cu(NH_3)_4(H_2O)]^{2+}$ to show how complex ions can show cis/trans isomerism





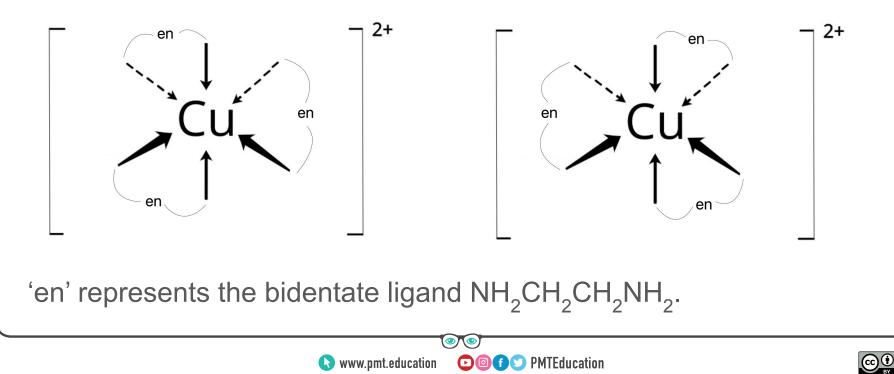
Use diagrams of [Cu(en)₃] to show how complex ions can show optical isomerism







Use diagrams of $[Cu(en)_3]$ to show how complex ions can show optical isomerism





Draw diagrams of cisplatin and transplatin. What is the shape of these stereoisomers?

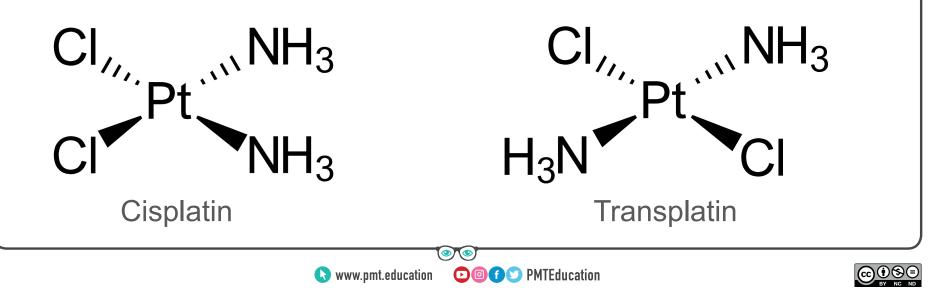






Draw diagrams of cisplatin and transplatin. What is the shape of these stereoisomers?

The shape is square planar.





Cisplatin is used as an anti-cancer drug. What does it do?







Cisplatin is used as an anti-cancer drug. What does it do?

Cisplatin binds to DNA, preventing cells replicating. This leads to cell death (apoptosis), preventing uncontrolled cell division.







What is K_{stab} ?







What is K_{stab}?

The equilibrium constant for the formation of a complex ion from its constituent molecules/ions, in a solvent.







Write an expression for K_{stab} for the ligand exchange reaction below: $[Cr(H_2O)_6]^{3+} + 6NH_3 \rightleftharpoons [Cr(NH_3)_6]^{3+} + 6H_2O$







Write an expression for K_{stab} for the ligand exchange reaction below: $[Cr(H_2O)_6]^{3+} + 6NH_3 \rightleftharpoons [Cr(NH_3)_6]^{3+} + 6H_2O$

- Square brackets now represent concentration (ignore any current square brackets).
- Power of each species = balancing number in chemical equation.

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• Water isn't included in the equation because the reaction is happening in solution so it is in excess and so does not affect the expression.

$$K_{stab} = \frac{[Cr(NH_3)_6^{3+}]}{[Cr(H_2O)_6^{3+}][NH_3]^6}$$

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What does it mean if the K_{stab} value is large?







What does it mean if the $\mathrm{K}_{\mathrm{stab}}$ value is large?

The larger the stability constant, the greater the stability of the complex ion.







Describe ligand exchange in terms of competing equilibria







Describe ligand exchange in terms of competing equilibria

When there are two competing equilibria in a ligand exchange reaction, the reaction that forms the most stable complex will be prioritised. This will be the equilibrium with the greatest K_{stab} value.



