ChemFactsheet

www.curriculum-press.co.uk

Number 236

Stability Constants (Kstab)

To succeed in this topic you will need to:

- 1. Understand the concepts of 'dynamic equilibrium' and Le Chatelier's Principle (Factsheet 9)
- 2. Understand the concepts of complex formation and ligand substitution in transition metal ions (Factsheet 46)
- 3. Understand the concepts of entropy (Factsheet 194)

After working through this Factsheet you will be able to:

- 1. Understand the concept of stability constants and write out their mathematical expressions from balanced equations
- 2. Describe how stability constants can be used to predict the formation of a complex
- 3. Explain the biochemical importance of ligand substitution in haemoglobin involving O_2 and CO

What are Stability Constants (K_{stab} **)?**

Stability constants (K_{stab}) are a way of measuring the stability of transition metal ion complexes.

They are equilibrium constants and the rules that need to be followed when dealing with them are the same as any other equilibrium constant such as K_c , K_a and K_w and, of course, as with all equilibrium constants, they will vary with temperature.

Definition: K_{stab} is the equilibrium constant for an equilibrium existing between a transition metal ion surrounded by water ligands and the complex formed when the same transition metal ion has undergone a ligand substitution reaction.

Ligands are substituted one at a time and so an equilibrium expression could be written for each step. However, in this Factsheet just the expression for the overall substitution reaction will be used.

Example:

The stability constant for the $[CoCl₄]$ ² complex can be expressed by considering how it is formed from the aqueous cobalt(II) ion.

 $[Co(H₂O)₆]^{2+}(aq) + 4Cl^{-}(aq) \rightleftharpoons [CoCl₄]^{2-}(aq) + 6H₂O(l)$

$$
K_{\text{stab}} = \frac{[[CoCl_4]^2]}{[[Co(H_2O)_6]^2^+].[Cl^4}
$$

[Note: with square brackets representing concentration AND complex ions, the K_{sub} expression has a lot of brackets! If you make square brackets representing concentration slightly larger you will be able to clearly see that you have included all you need to.]

As usual the $[H_2O]$ should be left out of the expression as all the species are dissolved in water and so water is in large excess. The concentration of water can be considered constant.

Units of K_{stab}

The units for K_{stab} can be worked out in the same way as the units for any other equilibrium constant. Each component has units of mol dm-3 and you need to make sure that you take into account the powers to which the concentration terms are raised. Cancel out any units that you can before you express the final unit

$$
K_{\text{stab}} = \frac{[[\text{CoCl}_4]^2]}{[[\text{Co}(H_2\text{O})_6]^2]^2][\text{Cl}^4]} = \frac{\text{mod} \text{dim}^3}{\text{mod} \text{dim}^3 \cdot (\text{mod} \text{ dm}^3)^4} = \frac{1}{\text{mod}^4 \text{dm}^{-12}}
$$

 $=$ mol⁻⁴dm¹²

[Note: some people prefer the units to be written in decreasing index order. i.e. $dm^{12} mol^{-4}$]

The same principles can be applied to the substitution of H_2O ligands by multidentate ligands such as ethane-1,2-diamine (abbreviated to "en"), ethanedioate $(C_2O_4^2)$ and EDTA⁴.

$$
[Cu(H2O)6]2+(aq) + 3en (aq) \rightleftharpoons [Cu(en)3]2+(aq) + 6H2O (l)
$$

$$
Kstab = \frac{[[Cu(en)3]2+]}{[[Cu(H2O)6]2+][en]3}
$$
mol⁻³dm⁹

The Magnitude of K_{stab}

The larger the value of K_{stab} the further the equilibrium lies in favour of the products (i.e. the substituted complex). The more stable the complex, the higher the value of its K_{stab} . This means that you can compare the stability of differently substituted metal ions and predict the complex that will be more likely to form in solution.

[Note: sometimes the values are quoted as log K_{stab} to make the numbers easier to handle]

Questions

- 1. Write the expression for K_{stab} for the complex $[Ni(NH_3)_6]^{2+}$ and give the units.
- 2. Consider the data below. Rank the complex ions in order of increasing stability.

3. An aqueous solution containing 1.0 mol dm⁻³ of $[CuCl₄]$ ² and 0.5 mol dm⁻³ Cl⁻ions was at equilibrium. Calculate the concentration of $[Cu(H₂O)₆]^{2+}$ present if $K_{stab} = 4.17 \times 10^5$ mol⁻⁴ dm¹²

Stability of Multidentate Ligand Complexes

The stability constants for complexes with multidentate ligands show that multidentate complexes are more stable than complexes containing only monodentate ligands.

The diagram below shows three complexes of nickel (II) ion. The values of K_{stab} for each complex are also given. You can see by looking at the K_{stab} values that the complex containing the bidentate ligand, en, is more stable than the one containing the monodentate ligand. Similarly, the complex containing the hexadentate ligand, EDTA, is more stable still.

This also means that, since the K_{stab} for the EDTA complex is greater than the other complexes, that EDTA will substitute for the other ligands.

e.g. $[Ni(NH_3)_6]^{2+}$ (aq) + EDTA⁴ (aq) = $[Ni(EDTA)]^{2-}$ (aq) + 6NH₃ (aq)

This is called the chelate effect. A chelate is a complex ion that contains a multidentate ligand. A chelating agent is any multidentate ligand that will substitute for monodentate ligands.

[Note: chelate comes from the Greek word meaning 'claw'.]

Explaining the Chelate Effect

Whichever example you consider, you will always find that a chelate is more stable than an ion with only monodentate ligands. To explain this, you have to consider the substitution equation and the entropy change that occurs in the reaction.

Let's take the equilibrium for the substitution of $NH₃$ ligands by EDTA shown above.

We can predict that the entropy change for the reaction (ΔS) will be positive. Remember that entropy is a measure of disorder – the more disorder, the higher the entropy. Reactions tend to favour an increase in entropy. This is not always the case, as enthalpy has a part to play in reaction predictions too, but in this case, the enthalpy change is fairly small.

In the forward reaction, you can see that 2 species react to form 7 species. You can get more disorder from 7 species than 2 so there will be a significant increase in entropy $(\Delta S$ will be positive). The reverse reaction would involve a significant decrease in entropy $(\Delta S$ will be negative) and so the reaction is unlikely to happen unless there is a large negative change in enthalpy (ΔH needs to be negative). This means that the chelate is very stable.

[Note: There is unlikely to be a significant change in enthalpy in these ligand substitution reactions since the number and type of bonds broken and made in the reaction is similar.]

Questions

4. The K_{stab} values for two cobalt(II) complexes are shown below.

 $[Co(NH₃)₆]²⁺$ 2.5 × 10⁴ mol⁻⁶ dm¹⁸ $[Co(EDTA)]^2$ - 2.0 × 10¹⁶ mol⁻¹ dm³

- (a) Write the equations for the formation of these complexes from the hexaaquacobalt(II) ion, $[Co(H_2O)_6]^{2+}$
- (b) Write expressions for the stability constants for these complexes and give the units.
- (c) Explain why the stability constant of the $[Co(EDTA)]^2$ complex is much greater than that of the $[Co(NH_3)_6]^{2+}$ complex

Explaining Carbon Monoxide Poisoning with Stability Constants

Haemoglobin is a protein in red blood cells that binds and transports oxygen from the lungs to the tissues. The haem group in the haemoglobin molecule is a non-polypeptide co-factor that consists of a central Fe²⁺ ion coordinately bonded to a porphyrin ring. The porphyrin ring can be described as a tetradentate ligand since it forms 4 coordinate bonds to the $Fe²⁺$ ion.

One of the other coordinate bonds to the Fe^{2+} ion is responsible for attaching the haem group to the globin polypeptide chain and the other is where the oxygen molecule is attached. When oxygen is not being carried, a water ligand is usually present instead.

2

There are 4 haem groups per haemoglobin molecule. So each molecule of haemoglobin can bind up to 4 oxygen molecules.

Deoxygenated haemoglobin (deoxyhaemoglobin) can be represented as $Hb(H_2O)₄$

Oxygenated haemoglobin (oxyhaemoglobin) can be represented as $Hb(O_2)_4$

Consequently, when deoxygenated blood reaches the lungs, a ligand substitution reaction occurs.

$$
Hb(H_2O)_4 + 4O_2 \xrightarrow{\text{lungs}} Hb(O_2)_4 + 4H_2O
$$

Of course, in the tissues, the reverse reaction takes place and oxygen is released to be used by the cells for respiration.

[Note: Le Chatelier's Principle can be applied here to describe how under conditions of high oxygen concentration (i.e. the lungs) the forward reaction is favoured and oxygen binds to the haemoglobin and under conditions of low oxygen concentration (i.e. the tissues) the reverse reaction is favoured and oxygen is released.]

The formation of oxyhaemoglobin falls precisely into the type of reactions for which we can describe a stability constant.

$$
K_{\text{stab}} = \frac{[Hb(O_2)_4]}{[Hb(H_2O)_4] \cdot [O_2]^4}
$$

Carbon monoxide molecules also bond to the $Fe²⁺$ ions in haemoglobin to form carboxyhaemoglobin, $Hb(CO)_{4}$.

A ligand substitution equilibrium for the reaction in which carbon monoxide replaces the water ligands can be written.

$$
\text{Hb}(\text{H}_2\text{O})_4 + 4\text{CO} \rightleftharpoons \text{Hb}(\text{CO})_4 + 4\text{H}_2\text{O}
$$

The stability constant for carboxyhaemoglobin is much greater than for oxyhaemoglobin. This means that carboxyhaemoglobin is more stable than oxyhaemoglobin. Therefore carbon monoxide binds in preference to oxygen and forms a stronger coordinate bond to the Fe2+ ion. Consequently if oxyhaemoglobin is exposed to carbon monoxide, the oxygen molecules will be substituted for carbon monoxide.

 $Hb(O_2)_4 + 4CO \rightleftharpoons Hb(CO)_4 + 4O_2$

The equilibrium lies heavily towards the right hand side.

Of course, treatment of carbon monoxide poisoning involves breathing air containing high levels of oxygen. Le Chatelier's Principle can be seen to be applied here as increasing oxygen concentration will encourage the equilibrium to move to the left hand side causing the carbon monoxide to be released.

Questions

- 5. Using ligand substitution reactions, explain how haemoglobin can transport oxygen in the blood.
- 6. Carbon monoxide binds to haemoglobin in preference to oxygen. Explain in terms of stability constants and bond strengths why this is the case.

Answers to Questions

$$
1. \quad K_{\text{stab}} = \frac{[[Ni(NH_3)_6]^{2+}]}{[[Ni(H_2O)_6]^{2+}].[NH_3]^{6}} \quad mol^{-6}dm^{18}
$$

2. lowest stability $\big)_{6}$]²⁺ highest stability $\mathrm{d}_{4}(\mathrm{H}_{2}\mathrm{O}_{2})^{2+}$

[Note: remember as the stability constant increases, the stability of the complex increases]

3. Substituting the data in the question into the expression for $K_{\mu\nu}$ and rearranging gives $1.0/((4.17 \times 10^5) \times 0.5^4) = 3.84 \times 10^{-5}$ mol dm⁻³

[Note: remember to raise the concentration of Cl- ions to the power 4]

4. (a) $[Co(H_2O)_6]^2$ ⁺ (aq) + 6NH₃(aq) $\Rightarrow [Co(NH_3)_6]^2$ ⁺(aq) + 6H₂O(l) $[Co(H₂O)₆]^{2+}$ (aq) + EDTA⁴ (aq) \Rightarrow $[Co(EDTA)]^{2}$ (aq) + 6H₂O(l)

(b)
$$
K_{\text{stab}} = \frac{\text{[[Co(NH_3)_6]^{2+}]} \text{mol} \cdot \text{6} \text{dm}^{18}}{\text{[[Co(H_2O)_6]^{2+}].[NH_3]^{6}}}
$$
mol⁻⁶dm¹⁸

$$
K_{\text{stab}} = \frac{\text{[[Co(EDTA)]^{2-}]} \text{mol}^{-1} \text{dm}^{3}}{\text{[[Co(H_2O)_6]^{2+}].[EDTA^{4-}]}}
$$
mol⁻¹dm³

- (c) The EDTA complex is more stable than the $NH₃$ complex because when the EDTA complex is formed there is a large increase in entropy (7 species formed from 2 species). In the formation of the $NH₃$ complex, there is very little entropy change (7 species formed from 7 species). This effect is called the chelating effect.
- 5. In the lungs, an oxygen molecule binds to the $Fe²⁺$ ion in the haem group of the haemoglobin molecule substituting a water molecule. As the blood circulates through the body, the bond between the oxygen molecule and the $Fe²⁺$ ion is broken and oxygen is released into the tissues.
- 6. K_{stab} for carboxyhaemoglobin is greater than for oxyhaemoglobin. This means that carboxyhaemoglobin is more stable than oxyhaemoglobin. The coordinate bond that is formed between the CO molecule and the Fe^{2+} ion is stronger than the bond between the oxygen molecule and the $Fe²⁺$ ion.

Acknowledgements: This Factsheet was researched and written by Martin Scott. Curriculum Press, Bank House, 105 King Street, Wellington, Shropshire, TF1 1NU. ChemistryFactsheets may be copied free of charge by teaching staff or students, provided that their school is a registered subscriber. No part of these Factsheets may be reproduced, stored in a retrieval system, or transmitted, in any other form or by any other means, without the prior permission of the publisher. ISSN 1351-5136