

Consider The Reaction $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$ What Is The Form Of The Equilibrium Constant K_c For This Reaction?

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Understanding the form of the equilibrium constant, K_c , for chemical reactions is fundamental in chemistry, especially when analyzing reaction dynamics and predicting the direction of reactions. The given reaction: $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$, involves different phases of matter—solids, liquids, and gases—and each phase contributes differently to the expression of the equilibrium constant. This article aims to comprehensively explore the nature of K_c for this specific reaction, elucidate how it is formulated, and discuss the underlying principles governing its behavior.

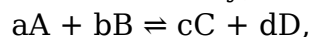
Overview of Equilibrium Constants in Chemical Reactions

Before delving into the specifics of the reaction involving mercury oxide, it is essential to understand what an equilibrium constant is, its significance, and the general rules for formulating it.

What Is an Equilibrium Constant?

The equilibrium constant, denoted as K , quantifies the ratio of the concentrations (or partial pressures) of products to reactants at equilibrium. It provides a snapshot of the reaction's position—whether it favors products or reactants—under specific conditions.

Mathematically, for a general reaction:



the equilibrium constant is expressed as:

$$K = \frac{[\text{C}]^c [\text{D}]^d}{[\text{A}]^a [\text{B}]^b},$$

where square brackets denote molar concentrations.

Depending on the phase of the species involved, different forms of the equilibrium constant are used, such as K_c (based on concentrations) and K_p (based on partial pressures).

Key Principles Governing K_c Formulation

- Pure solids and pure liquids do not appear in the expression because their activities are considered constant and incorporated into the equilibrium constant.
- Gases are included via their partial pressures or concentrations, depending on the form

of the equilibrium constant used.

- The equilibrium constant is temperature-dependent but independent of initial concentrations or pressures.

Analyzing the Reaction: $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$

This specific reaction involves the decomposition of solid mercury(II) oxide into liquid mercury and gaseous oxygen. To formulate the equilibrium constant, we need to analyze each phase involved.

Phases Involved in the Reaction

- Solid mercury oxide (HgO(s))
- Liquid mercury (Hg(l))
- Gaseous oxygen ($\text{O}_2\text{(g)}$)

The key aspect here is understanding how these phases influence the formulation of K_c .

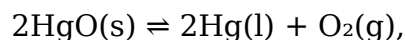
Why Do Solids and Liquids Usually Not Appear in K_c Expressions?

In equilibrium expressions, the activities of pure solids and liquids are generally considered constant and are set to unity. This is because their activities do not change significantly with concentration or pressure, given their pure state.

Therefore, only gases and aqueous species (if any) are included in the expression for K_c .

Formulating the Equilibrium Constant K_c for the Reaction

Given the reaction:



the first step is to write the general expression for K_c based on the activities of the species involved.

Activities of the Species

- Solid HgO : Activity = 1 (since it is a pure solid)
- Liquid Hg : Activity = 1 (assuming pure liquid mercury)
- Gaseous O_2 : Activity = partial pressure or molar concentration, depending on the formulation

Because the solids and liquids are pure, their activities are taken as constant (set to 1), simplifying the expression.

Expression for K_c

The equilibrium constant in terms of concentrations (K_c) is expressed as:

$$K_c = \frac{a_{\text{Hg}}^2 \times a_{\text{O}_2}}{a_{\text{HgO}}^2}$$

However, since activities of pure solids and liquids are 1, and only gases contribute to the equilibrium expression, this simplifies to:

$$K_c = a_{\text{O}_2}$$

or, in terms of molar concentrations:

$$K_c = [\text{O}_2]$$

Note: The activities of the liquid mercury and solid HgO are omitted because they are constant and incorporated into the equilibrium constant.

Understanding the Final Form of K_c

The simplified form of the equilibrium constant for this reaction is:

$$K_c = [\text{O}_2]$$

This indicates that the equilibrium position depends solely on the concentration (or partial pressure) of oxygen gas at equilibrium.

Alternatively, if partial pressures are used, K_p (pressure-based equilibrium constant) can be expressed as:

$$K_p = P_{\text{O}_2}$$

where P_{O_2} is the partial pressure of oxygen at equilibrium.

Implications of the Form of K_c

- The reaction's equilibrium position can be predicted based on the concentration or partial pressure of oxygen.
- Since only the gas phase appears in the expression, changes in pressure or oxygen concentration can shift the equilibrium.

Practical Considerations in Measuring K_c

When experimental chemists seek to determine the value of K_c for this reaction, they typically measure the concentration or partial pressure of oxygen at equilibrium under specific conditions and then calculate K_c accordingly.

Factors Affecting Kc

- Temperature: As with all equilibria, Kc is temperature-dependent. Increased temperature can shift the equilibrium, changing the value of Kc.
- Pressure: Changes in external pressure influence the partial pressure of oxygen, thereby affecting the equilibrium position.

Comparison With Other Types of Equilibrium Constants

It's crucial to differentiate Kc from other equilibrium constants such as Kp or Kw (ionic product of water), as their formulations depend on the phases involved and whether concentrations or pressures are used.

Type of Constant	Definition	Applicable Phases	Expression
Kc	Based on concentrations	Gases, aqueous solutions	$\frac{[\text{products}]}{[\text{reactants}]}$ (excluding pure solids/liquids)
Kp	Based on partial pressures	Gases	$\frac{P_{\text{products}}}{P_{\text{reactants}}}$
Kw	Ionic product of water	Aqueous	$[\text{H}^+][\text{OH}^-]$

In this reaction, Kc is the appropriate constant because it involves gaseous oxygen, and the phase activities are well-defined.

Summary and Key Takeaways

- The reaction: $2\text{HgO}(\text{s}) \rightleftharpoons 2\text{Hg}(\text{l}) + \text{O}_2(\text{g})$, involves phases with different behaviors in equilibrium expressions.
- Pure solids and liquids do not appear in the formulation of Kc because their activities are constant and incorporated into the equilibrium constant.
- The equilibrium constant Kc for this reaction simplifies to the molar concentration (or activity) of oxygen gas, i.e., $K_c = [\text{O}_2]$.
- The form of Kc provides insights into how the reaction shifts under changing conditions, such as temperature and pressure.
- Understanding the formulation of Kc is essential for predicting reaction behavior, optimizing industrial processes, and conducting laboratory analysis.

Conclusion

In summary, the form of the equilibrium constant Kc for the reaction $2\text{HgO}(\text{s}) \rightleftharpoons 2\text{Hg}(\text{l}) + \text{O}_2(\text{g})$ is primarily determined by the gaseous oxygen component. Since pure solids and liquids are assigned activities of unity, they do not appear in the expression. Therefore, $K_c = [\text{O}_2]$, where $[\text{O}_2]$ is the molar concentration of oxygen at equilibrium, or equivalently, $K_p = P_{\text{O}_2}$, the partial pressure of oxygen. Recognizing these nuances allows chemists to accurately interpret the reaction's equilibrium state and manipulate conditions to favor desired outcomes, whether in industrial mercury processing or research settings.

Frequently Asked Questions

What is the form of the equilibrium constant K_c for the reaction $2 \text{HgO(s)} \rightleftharpoons 2 \text{Hg(l)} + \text{O}_2\text{(g)}$?

The equilibrium constant K_c is expressed in terms of the concentrations of the gaseous species: $K_c = [\text{O}_2]$, since pure solids and liquids are omitted from the expression.

Why are the concentrations of solids and liquids omitted when writing K_c for this reaction?

Because the activities of pure solids and liquids are considered constant and are incorporated into the equilibrium constant, only gaseous and aqueous species are included in the expression.

How would you write the expression for K_c for the reaction $2 \text{HgO(s)} \rightleftharpoons 2 \text{Hg(l)} + \text{O}_2\text{(g)}$?

$K_c = [\text{O}_2]$, since only the gaseous oxygen concentration appears in the equilibrium expression.

Does the presence of mercury in liquid form affect the calculation of K_c for this reaction?

No, because liquids are omitted from the equilibrium expression; only gases are included, so the mercury liquid phase does not appear in K_c .

Can the equilibrium constant K_c be used to determine the extent of the reaction at equilibrium?

Yes, the value of K_c indicates the position of equilibrium; if K_c is large, the reaction favors products, and if small, it favors reactants.

Is the equilibrium constant K_c dimensionless for this reaction?

Typically, yes, because concentrations are expressed in molarity, and the way the expression is set up results in a dimensionless value or one with units depending on the expression conventions.

What role does temperature play in determining the value of K_c for this reaction?

Temperature affects the value of K_c because equilibrium constants are temperature-dependent; increasing temperature may shift the equilibrium and change K_c .

How does the reaction $2 \text{HgO(s)} \rightleftharpoons 2 \text{Hg(l)} + \text{O}_2\text{(g)}$ illustrate the concept of equilibrium in phase changes?

It shows how a solid decomposes into a liquid and a gas, reaching a balance where the rates of formation and consumption of each phase are equal, and the equilibrium constant reflects the gaseous component.

Would the equilibrium constant K_c change if the reaction was carried out at a different temperature?

Yes, since K_c is temperature-dependent, changing the temperature will alter the equilibrium position and the value of K_c .

Why is it important to understand the form of K_c for this reaction in industrial or laboratory settings?

Understanding K_c helps predict how the reaction will shift under different conditions, which is essential for controlling processes like mercury production or decomposition in industrial applications.

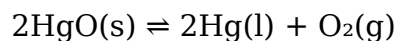
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Understanding the equilibrium constant, often denoted as K_c , is fundamental in grasping chemical equilibria. When dealing with the reaction $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$, it becomes essential to analyze how the equilibrium constant is expressed and what factors influence it. This reaction involves a solid (mercury(II) oxide), a liquid (mercury), and a gas (oxygen), which introduces specific considerations in formulating the equilibrium constant. In this article, we will explore the nature of the equilibrium constant K_c for this specific reaction, analyze its formulation, and discuss key features, advantages, and limitations associated with it.

Understanding the Reaction: An Overview

Before delving into the form of K_c , it's vital to understand the reaction at hand:



This is a decomposition reaction where solid mercury(II) oxide decomposes into liquid mercury and oxygen gas upon heating. The reaction is reversible, and at equilibrium, the rates of the forward and reverse reactions are equal.

Key Points:

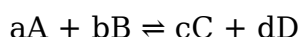
- The reaction involves a solid, a liquid, and a gas.
- The equilibrium position depends on temperature and pressure.
- The reaction is used industrially and historically in mercury extraction and thermodynamic studies.

What Is the Equilibrium Constant K_c ?

The equilibrium constant, K_c , quantifies the ratio of product concentrations to reactant concentrations at equilibrium for a given reaction at a specific temperature. It provides a measure of the extent to which a reaction proceeds before reaching equilibrium.

General Definition:

For a reaction:



the equilibrium constant K_c is expressed as:

$$K_c = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

where brackets denote molar concentrations (mol/L).

Form of K_c for the Reaction $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$

When formulating K_c for this particular reaction, the key is understanding which phases are included in the expression and how their activities or concentrations are represented.

Involving Solids and Liquids:

- The activities of pure solids and pure liquids are conventionally taken as unity (1) because their activities do not change significantly with concentration or pressure.
- This simplifies the expression for K_c by excluding solids and liquids from the concentration ratio, focusing only on gases.

Therefore, the expression for K_c in this reaction is:

$$K_c = [\text{O}_2]$$

Why?

- The activity of the solid HgO and the liquid Hg are considered constant and set to 1.
- The only gaseous species, O₂, appears in the numerator, and its molar concentration determines the equilibrium state.

Resulting Expression:

$$K_c = [O_2]$$

where [O₂] is the molar concentration of oxygen gas at equilibrium.

Why Is Only O₂ Included in the Expression?

The exclusion of solids and liquids from the equilibrium expression hinges on the concept of activity. Activities of pure solids and liquids are defined as 1 because:

- Their concentrations do not significantly change during the reaction.
- They serve as a reference state.

This simplifies the equilibrium expression and emphasizes the role of gaseous components in the reaction's equilibrium position.

Alternative Forms of the Equilibrium Constant

Depending on the conditions and what data is available, other forms of the equilibrium constant can be used:

- K_p (Partial Pressure Form): When dealing with gases, it is often more convenient to express the equilibrium in terms of partial pressures.

For this reaction:

$$K_p = p_{O_2} \text{ (partial pressure of oxygen at equilibrium)}$$

- Relation between K_c and K_p:

The two are related via the ideal gas law:

$$K_p = K_c (RT)^{\Delta n}$$

where:

- R = universal gas constant
- T = temperature in Kelvin
- Δn = change in moles of gases (moles of gaseous products - moles of gaseous reactants)

In our reaction:

- Gaseous products: 1 mole of O_2
- Gaseous reactants: 0 (since solids and liquids are omitted)

Change in moles:

$$\Delta n = 1 - 0 = 1$$

Therefore,

$$K_p = K_c RT$$

This relationship allows conversion between concentration and pressure-based equilibrium constants.

Key Features and Considerations

Understanding the form of K_c for this specific reaction involves recognizing several features:

Features:

- Solid and Liquid Phases Are Excluded: Their activities are set to 1, simplifying the expression.
- Gaseous Species Are Included: Only gases appear explicitly in the expression, either as concentrations or partial pressures.
- Temperature Dependence: The value of K_c depends heavily on temperature, often increasing or decreasing as the reaction temperature changes.
- Relation to Thermodynamics: The equilibrium constant relates to the standard Gibbs free energy change (ΔG°) via the relation:

$$\Delta G^\circ = -RT \ln K$$

Pros:

- Simplifies calculations by excluding solids and liquids.
- Enables focus on gaseous components, which are often easier to measure.
- Facilitates the use of pressure or concentration data interchangeably through K_p and K_c .

Cons:

- Assumes ideal behavior, which may not hold at high pressures or for non-ideal gases.

- The activity of solids and liquids is set to 1, which is an approximation that may sometimes introduce errors.
- Changes in the physical state or purity of solids/liquids are not reflected in the equilibrium constant.

Implications in Industrial and Laboratory Settings

The form of the equilibrium constant K_c for this reaction has several practical applications:

Industrial Significance:

- Mercury production and purification processes often involve the decomposition of HgO .
- Understanding the equilibrium helps optimize temperature and pressure conditions for maximum efficiency.
- The gas-phase composition at equilibrium can be predicted and controlled.

Laboratory Analysis:

- Measuring partial pressures or concentrations of oxygen at equilibrium allows for the determination of K_c .
- The reaction serves as an example in teaching thermodynamics and equilibrium principles.

Features in Practice:

- The easy exclusion of solids and liquids simplifies experimental analysis.
- Use of partial pressures (K_p) is often more convenient in gas reactions, especially at high temperatures.

Conclusion: The Form of the Equilibrium Constant K_c for the Reaction

In summary, for the reaction $2\text{HgO(s)} \rightleftharpoons 2\text{Hg(l)} + \text{O}_2\text{(g)}$, the equilibrium constant K_c is expressed primarily in terms of the gaseous species involved, primarily oxygen. The standard form simplifies to:

$$K_c = [\text{O}_2]$$

where $[\text{O}_2]$ is the molar concentration of oxygen gas at equilibrium. This formulation stems from the principle that activities of pure solids and liquids are taken as unity, which streamlines the equilibrium expression. When considering partial pressures, the analogous

constant K_p can be used, related to K_c through the ideal gas law and the temperature-dependent relation involving RT and Δn .

Understanding this form aids chemists and engineers in predicting the behavior of the system, optimizing conditions for industrial processes, and conducting accurate thermodynamic calculations. The balance between simplicity and precision is critical—while the formula provides a practical approximation, awareness of its assumptions and limitations ensures proper application in real-world scenarios.

In conclusion, the form of the equilibrium constant K_c for the reaction $2\text{HgO}(s) \rightleftharpoons 2\text{Hg}(l) + \text{O}_2(g)$ is primarily based on the concentration or partial pressure of oxygen gas, with solids and liquids excluded. This approach exemplifies the core principles of chemical equilibrium and the importance of phase considerations in defining equilibrium constants.

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