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**THERMODYNAMICS, KINETICS AND MECHANISM OF ABSORPTION OF A MIXTURE
OF GASES CONSISTING OF SULFUR OXIDES INTO ALKALINE SOLUTIONS**

Abdirashid Khasanov

Deputy Chief Engineer for Science
at "Almalyk MMC" JSC, doctor of technical Science, Professor,
Republic of Uzbekistan, Tashkent
E-mail: a.hasanov@agmk.uz

Shokhrukh Khojiev

Associate professor of department of Metallurgy, PhD,
Tashkent State Technical University,
Republic of Uzbekistan, Tashkent
E-mail: hojiyevshohruh@yandex.ru

Shokhrukh Munosibov

Doctoral student
of the National University of Uzbekistan,
Republic of Uzbekistan, Tashkent
E-mail: shokhrukhmunosibov@gmail.com

Vasliddin Khatamkulov

Student
of master course of department of Metallurgy,
Tashkent State Technical University,
Republic of Uzbekistan, Tashkent
E-mail: hojiyevshohruh@yandex.ru

**ТЕРМОДИНАМИКА, КИНЕТИКА И МЕХАНИЗМ АБСОРБЦИИ СМЕСИ ГАЗОВ,
СОСТОЯЩЕЙ ОКСИДОВ СЕРЫ, В ЩЕЛОЧНЫЕ РАСТВОРЫ**

Хасанов Абдирашид Салиевич

зам. главного инженера по науке
АО «Алмалыкский ГМК»,
д-р техн.х наук, профессор,
Республика Узбекистан, г. Ташкент

Хожиев Шохрух Тошпулатович

и.о. доцент
кафедры Металлургия, PhD,
Ташкентский государственный технический университет,
Республика Узбекистан, г. Ташкент

Муносибов Шохрух Мухиддин угли

докторант
Национального университета Узбекистана,
Республика Узбекистан, г. Ташкент

Хатамкулов Василюдин Хатамкулович

студент магистратуры,
кафедры Металлургия,
Ташкентский Государственный Технический Университет,
Республика Узбекистан, г. Ташкент

ABSTRACT

In the article, the thermodynamics, kinetics and mechanism of absorption of a mixture of sulfurous gases in alkaline solutions are considered. The result of thermodynamic analysis showed that the value of Gibbs energy of absorption of sulfur dioxide and trioxide into sodium hydroxide solution under standard conditions was negative. It was determined that the limiting step is the absorption of sulfur dioxide, and ways to increase the partial pressure of the gas were proposed to increase the reaction yield.

АННОТАЦИЯ

В статье рассмотрены термодинамика, кинетика и механизм абсорбции смеси сернистых газов в щелочных растворах. Результат термодинамического анализа показал, что значение энергии Гиббса поглощения диоксида и триоксида серы раствором гидроксида натрия в стандартных условиях было отрицательным. Установлено, что лимитирующей стадией является поглощение диоксида серы, и предложены пути увеличения парциального давления газа для увеличения выхода реакции.

Keywords: sulfur dioxide, sulfur trioxide, absorption, alkaline medium, thermodynamics, kinetics, mechanism.

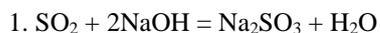
Ключевые слова: диоксид серы, триоксид серы, абсорбция, щелочная среда, термодинамика, кинетика, механизм.

The process of absorption of SO₂ and SO₃ in an alkaline environment has a wide range of practical applications, from the removal of pollutants in flue gas streams to the production of various chemicals. This process plays a crucial role in reducing air pollution and ensuring the sustainability of industrial processes. The method of gas chromatography was used to determine the composition of gases in the air, and the method of High-Performance Liquid Chromatography was used to determine the purity of the liquid absorbent. In the research, sodium hydroxide was used as a substance that creates an alkaline environment. In order to conduct a thermodynamic analysis of the reactions in the solution system, it is necessary to calculate the standard Gibbs free energy change (ΔG°) and the equilibrium constant (K) at a certain temperature [1-5]. The equations for ΔG° and K are:

$$\Delta G^\circ = -RT \ln K \quad (1)$$

Where: R is the universal gas constant (8.314 J/mol*K), T is the temperature in Kelvin, $\ln K$ is the natural logarithm of the equilibrium constant.

The chemical processes involved in each gas component were thermodynamically analyzed [6-7]:



The standard Gibbs free energy change for this reaction can be calculated using the values of the standard Gibbs free energy of formation (ΔG_f°) of the reactants and products:

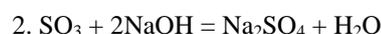
$$\Delta G^\circ = \Delta G_f^\circ(\text{Na}_2\text{SO}_3) + \Delta G_f^\circ(\text{H}_2\text{O}) - \Delta G_f^\circ(\text{SO}_2) - 2\Delta G_f^\circ(\text{NaOH}) \quad (2)$$

ΔG_f° values at 298 K: $\Delta G_f^\circ(\text{Na}_2\text{SO}_3) = -1095.8$ kJ/mol, $\Delta G_f^\circ(\text{H}_2\text{O}) = -237.2$ kJ/mol, $\Delta G_f^\circ(\text{SO}_2) = -300.4$ kJ/mol and $\Delta G_f^\circ(\text{NaOH}) = -470.1$ kJ/mol. If these values are replaced in the equation: $\Delta G^\circ = -40.8$ kJ/mol.

The equilibrium constant at 298 K can be calculated from the value of ΔG° using the following equation [8-9]:

$$K = e^{-\Delta G^\circ/RT} \quad (3)$$

Substituting the values gives: $K = 7.03 \cdot 10^6$



The standard Gibbs free energy change for this reaction can be calculated using the previous equation:

$$\Delta G^\circ = \Delta G_f^\circ(\text{Na}_2\text{SO}_4) + \Delta G_f^\circ(\text{H}_2\text{O}) - \Delta G_f^\circ(\text{SO}_3) - 2\Delta G_f^\circ(\text{NaOH}) \quad (4)$$

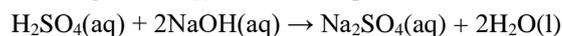
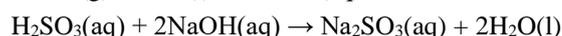
ΔG_f° values at 298 K: $\Delta G_f^\circ(\text{Na}_2\text{SO}_4) = -1385.1$ kJ/mol, $\Delta G_f^\circ(\text{H}_2\text{O}) = -237.2$ kJ/mol, $\Delta G_f^\circ(\text{SO}_3) = -371.1$ kJ/mol and $\Delta G_f^\circ(\text{NaOH}) = -470.1$ kJ/mol. Substituting these values into the equation, we get: $\Delta G^\circ = -36.6$ kJ/mol.

The equilibrium constant at 298 K can be calculated from the value of ΔG° using the previous equation: $K = 2.46 \cdot 10^6$

Negative values of ΔG° in both reactions indicate that the reactions are thermodynamically favorable and occur spontaneously [10].

The absorption of sulfur oxides such as sulfur dioxide (SO₂) and sulfur trioxide (SO₃) in alkaline media involves complex chemical reactions that occur in two stages. The first stage is the physical absorption of gas into the liquid phase, and then the chemical reaction of the absorbed gas with sodium hydroxide solution results in the formation of soluble sulfite or sulfate [11].

The absorption mechanism of sulfur oxides in an alkaline environment is usually described by the following equations:



The first equation shows the equilibrium between sulfur dioxide gas and water to form sulphitic acid, which then reacts with sodium hydroxide to form sodium sulphite and water in the second equation [12]. Similarly, the third equation shows the formation of sulfuric acid from sulfur trioxide and water, which in the fourth equation reacts with sodium hydroxide to form sodium sulfate and water. The kinetics of the gas absorption process depends on various factors, for example,

the gas concentration in the air, the temperature and pressure of the system, and the properties of the liquid absorbent. The rate of absorption is also affected by the reaction rate of chemical reactions that can be determined experimentally.

In alkaline environments, the rate of absorption of sulfur oxides generally follows first-order kinetics, where the rate of absorption is proportional to the concentration of the gas in the air. The rate constant, k , depends on various factors, such as the absorptivity of the liquid, the temperature, and the partial pressure of the gas [13].

The negative values of ΔG° in both reactions after absorption of SO_2 and SO_3 gases in alkaline solutions

indicate that the reactions are thermodynamically favorable and spontaneous.

The overall efficiency of the absorption process depends on the pH level of the alkaline solution, the concentration of the alkaline solution, and the surface area of the liquid gas absorption. Increasing the pH and concentration of an alkaline solution can increase absorption efficiency by increasing the rate of chemical reactions. The use of high-surface-area liquid gas absorbers, such as sprays or packed tower tanks, can also increase the efficiency of the absorption process by increasing the gas-liquid contact area.

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