Mechanism of Oxidation of L-Histidine by Heptavalent Manganese in Alkaline Medium

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Abstract: The kinetics of oxidation of L-histidine by manganese(VII) in aqueous alkaline medium at a constant ionic strength of 0.05 mol dm⁻³ was studied spectrophotometrically. The reaction between permanganate and L-histidine in alkaline medium exhibits 2:1 stoichiometry (KMnO₄ : L-histidine). The reaction is of first order in [KMnO₄], less than unit order in [L-histidine] and [alkali]. Decrease in the dielectric constant of the medium decreases the rate of reaction. Effect of added products and ionic strength of the reaction medium have been investigated. The main products were identified by spot test and I.R. A mechanism involving the free radical has been proposed. In composite equilibrium step L-histidine binds to MnO₄⁻ species to form a complex(C). The reaction constants involved in the different steps of mechanism are evaluated. The activation parameters with respect to slow step of the mechanism are computed and discussed and thermodynamic quantities are also determined.

Key words: kinetics, mechanism, oxidation, permanganate, L-histidine

Introduction

Oxidations by permanganate ion find extensive applications in organic syntheses¹⁻⁷ especially since the advent of phase transfer catalysis^{3,4,6}, which permits the use of solvents like methylene chloride and benzene. Kinetic studies are important sources of mechanistic information on these reactions, as demonstrated by result referring to unsaturated acids in both aqueous¹⁻⁸ and non-aqueous media⁸.

As is known, in aqueous alkaline medium the permanganate ion oxidizes a number of organic compounds, which are not, or only very slowly, attacked in acidic or neutral medium⁹⁻¹¹. If [OH⁻] ≥ 0.1 mol dm⁻³, permanganate is reduced in most case to manganate only, the reaction of which further proceeds at a much lower rate^{9,12,13}. Nevertheless, since the process is very fast¹²⁻¹⁴, it cannot be decided, in general, whether permanganate is reduced in one-electron steps or hypomanganate is first formed in a two-electron step followed by a fast reaction⁹.

Amino acids not only act as the building blocks in protein synthesis but also play a significant role in the metabolism. Histidine is a semiessential amino acid. It is found abundantly in hemoglobin; has been used in the treatment of rheumatoid arthritis, allergic diseases, ulcers and anemia. A deficiency can cause poor hearing. The importance of the amino acid, histidine lies in the fact that the body uses it to manufacture histamine, and histamine is responsible for a wide range of physiological processes. Apart from its sexual functions, histidine is involved in many other physiological processes. It is necessary for the production of red and white blood cells and supports the activity of suppressor T cells. Histidine is used as a supplement for sufferers of rheumatoid arthritis, since it has been shown that in these patients, histidine levels are low. Histidine is like many other amino acids, important for growth and general tissue repair.

Amino acids have been oxidized by variety of oxidizing agents¹⁵. Although much variety of organic¹⁶ and inorganic¹⁷ substrates are oxidized in aqueous alkaline medium, there are only a few reports¹⁸ on the oxidation of amino acids by aqueous alkaline permanganate. No work, however, has been reported on the oxidation of L-histidine by this oxidant in aqueous alkaline medium and so the present study deals with title reaction in order to understand the redox chemistry of permanganate in such media and to arrive a plausible mechanism.

Experimental

Materials

All chemicals used were of reagent grade. Double distilled water was used throughout the work. Stock solution of L-histidine (Sisco-Chem. Ltd.) was prepared by dissolving the appropriate amount of sample in distilled water. The purity of the sample was checked by TLC and its m.p. was 283 °C (Literature 285 °C). The solution of potassium permanganate (BDH) was prepared and standardized against oxalic acid¹⁹. Potassium manganate solution was prepared as described by Carrington and Symons²⁰ as follows: an aqueous solution of potassium permanganate was heated to boiling > 100°C in 8.0 mol dm⁻³ KOH solution. Using the required amount of recrystallised sample, a stock solution of potassium manganate was then prepared in aqueous KOH. The solution was standardized by measuring the absorbance on a Hitachi 150-20 spectrophotometer with a 1 cm Quartz cell at 608 nm ($\epsilon = 1530 \pm 20 \text{ dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$). All other reagents were of analytical grade and their solutions were prepared by dissolving requisite amounts of the samples in doubly distilled water. NaOH and NaClO₄ were used to provide the required alkalinity and to maintain the constant ionic strength respectively.

Kinetics measurements

The oxidation of L-histidine by KMnO₄ was followed under pseudo-first order conditions where [L-Hist] was excess over MnO₄⁻ at 30.0 ± 0.1 °C unless stated otherwise. The reaction was initiated by mixing the required quantities of previously thermostatted solutions of MnO₄⁻ and L-histidine, which also contained definite quantities of NaOH and NaClO₄ to maintain the required alkalinity and ionic strength. The progress of the reaction was followed by measuring the absorbance of unreacted MnO₄⁻ in the reaction mixture in a 1 cm quartz cell located in the thermostatted compartment of a Peltier Accessory (temperature control) attached Hitachi 150-20 spectrophotometer at 526 nm; other constituents of the reaction mixture do not absorb significantly at this wavelength. Application of Beer's law to permanganate at 526 nm earlier had been verified, giving molar extinction coefficient 'ε' was found to be 2083 ± 50 dm³ mol⁻¹ cm⁻¹ (Literature ε = 2200 dm³ mol⁻¹ cm⁻¹). The course of the reaction was followed by monitoring the decrease in the absorbance of KMnO₄. The first order rate constants (k_{obs}) were evaluated by plots of $\log(A_t - A_\infty)$ versus time by fitting the data to the expression $A_t = A_\infty + (A_0 - A_\infty) e^{-k_{\text{obs}}t}$, where A_t , A_0 and A_∞ are the absorbances of KMnO₄ of time t , 0, and ∞ respectively. In almost all cases the reaction was followed up to 75% completion of the reaction and the k_{obs} values were reproducible within ± 5%.

The effect of dissolved oxygen on the rate of reaction was studied by preparing the reaction mixture and following the reaction in an atmosphere of nitrogen. No significant difference between the results was observed. In view of the ubiquitous contamination of basic solutions by carbonate, the effect of carbonate on the reaction was also studied. Added carbonate had no effect on reaction rate. However, fresh solutions were used during the experiments.

Results

Stoichiometry: Different sets of reaction mixtures containing L-histidine and an excess MnO_4^- with constant OH^- and NaClO_4 were kept for 6 h in closed vessels under N_2 atmosphere. The remaining concentrations of MnO_4^- were assayed spectrophotometrically at 526 nm. The results indicated 1:2 stoichiometry as given in equation (1).

The main reaction products were identified as manganate by its visible spectra and 2-imidazole acetaldehyde²¹ by a spot test, ammonia¹⁹ by Nessler's reagent. The CO_2 evolved is tested by bubbling nitrogen gas through a tube containing lime water²² after acidification. The quantitative estimation of aldehyde as its 2,4-DNP derivative was nearly 80%. The aldehyde was also confirmed by its I.R. spectrum, which showed²³ a band at 2919 cm^{-1} due to aldehydic C-H stretching and 1676 cm^{-1} due to carbonyl stretching. Tests for the corresponding acid were negative. It was also observed that the aldehyde does not undergo further oxidation under the present kinetic conditions.

Reaction order: The order with respect to [L-histidine] and [alkali] were found by $\log k_{\text{obs}}$ versus log concentration plots and were also confirmed by differential method by the plot log rate versus log concentration using the equation $\log \text{rate} = \log k + n \log c$; these orders were obtained by varying the concentration of L-histidine and alkali in turn while keeping others constant.

The concentration of MnO_4^- was varied in the range, 4.0×10^{-5} to $6.0 \times 10^{-4}\text{ mol dm}^{-3}$ at fixed [L-histidine], $[\text{OH}^-]$ and ionic strength. The non-variation in the pseudo-first order rate constants at various concentrations of MnO_4^- indicates the order in $[\text{MnO}_4^-]$ as unity (Table 1). This was also confirmed from the linearity of plots of log absorbance versus time ($r > 0.9984$, $S \leq 0.025$) up to 75% completion of the reaction. The substrate, [L-histidine] was varied in the range of 8.0×10^{-4} to $8.0 \times 10^{-3}\text{ mol dm}^{-3}$ at 30°C keeping all other reactants concentrations constant. The k_{obs} values were increase with increase in concentration of L-histidine and it was found to be less than unit order dependence on [L-histidine] (Table 1). The effect of [alkali] on the rate of reaction was studied at constant concentrations of L-histidine, MnO_4^- and ionic strength at 0.05 mol dm^{-3} . The rate constants increased with increase in [alkali] and the order was also found to be less than unity (Table 1).

Effect of ionic strength and solvent polarity: The effect of ionic strength was studied by varying the NaClO_4 concentration from 5.0×10^{-3} to $5.0 \times 10^{-2}\text{ mol dm}^{-3}$ at constant concentration of permanganate, L-Histidine and alkali. It was found that the rate of reaction increased with increasing the concentration of NaClO_4 ; the plot of $\log k_{\text{obs}}$ versus $I^{1/2}$ is linear with a positive slope (Fig. 1).

The relative permittivity effect was evaluated by variation of the *t*-butanol-water content while keeping all other conditions constant. Attempts to measure the relative permittivities failed; however, they were computed from the values of pure liquids^{24,25}. The inertness of the solvent with respect to the oxidant was checked under the experimental conditions. There was no reaction of the solvent with the oxidant. The rate constant k_{obs} decrease with decreasing dielectric constant of the medium.

Effect of Initially added products: The effect of addition of products of the reaction, such as manganate, ammonia and aldehyde on rate of reaction were investigated. Addition of these products did not affect the reaction rate significantly.

Test for free radicals: To test the intervention of free radicals, the reaction mixture was mixed with acrylonitrile monomer and kept for 24 hours under nitrogen atmosphere. On dilution with methanol, white precipitate of polymer was formed, indicating the intervention of free radicals during the reaction. The blank experiment of either KMnO_4 or L-histidine in with acrylonitrile alone did not induce polymerization under the same condition as those induce with reaction mixture. Initially added acrylonitrile decreases the rate indicating the free radical intervention, which is the case as observed in earlier work²⁶.

Effect of Temperature: The rate of reaction was measured at different temperatures under varying L-histidine and alkali concentrations. The rate of reaction increased with the increase of temperature. The rate constants, k of slow step of scheme 1 were obtained from intercepts of the plots of $1/k_{\text{obs}}$ versus $1/[\text{L-histidine}]$ ($r > 0.9978$, $S \leq 0.0162$) as 3.43×10^{-3} , 4.03×10^{-3} , $4.75 \times 10^{-3} \text{ s}^{-1}$ at 30, 35, 40, 45°C respectively. The activation parameters corresponding to these constants were evaluated from the plot of $\log k$ versus $1/T$ ($r > 0.9989$, $S \leq 0.0135$) and are tabulated in **Table 2**.

Discussion

Potassium permanganate is widely used as an oxidizing agent in synthetic as well as in analytical chemistry and also as a disinfectant. The reactions are governed by pH of the medium. Among six oxidation states of manganese from 2+ to 7+, permanganate, Mn(VII) , is the most potent oxidation state in acid as well as in alkaline media.

The reaction between permanganate and L-histidine in alkaline medium has a stoichiometry of 1:2 reductant to oxidant with first order dependence on $[\text{MnO}_4^-]$, apparent less than unit order in $[\text{OH}^-]$ and [L-histidine]. The active species of permanganate in aqueous alkaline medium may be deduced from the dependence of the rate on $[\text{OH}^-]$ in the reaction medium. The apparent order of less than unity in OH^- may be indication of formation of alkaline permanganate ion in prior equilibrium step²⁷. The formation of $\text{MnO}_4\text{OH}^{2-}$ species in alkaline medium further supported by the Michaelis-Menten plot, which is linear with a positive intercept (**Fig. 2**). It is also known that permanganate ion in alkaline medium exist as $\text{MnO}_4\text{OH}^{2-}$. Thus, it reacts with L-histidine and leads to the formation a complex (C) in a prior equilibrium step. Then this complex (C) decomposes in a slow step to form a free radical species derived from L-histidine. This free radical reacts with another molecule of $[\text{MnO}_4\text{OH}]^{2-}$ species in a fast step to yield the products. All the results indicate a mechanism as given in **Scheme 1**.

The probable structure of complex (C) is

The spectral evidence for the complex (C) formation between oxidant and substrate was obtained from UV-VIS Spectra of the oxidant and mixtures of substrate and oxidant. Analogous effects upon complex (C) formation between a substrate and an oxidant have been observed earlier [28]. The formation of the complex (C) is also proved kinetically by the non-zero intercept of the plot of $1/k_{\text{obs}}$ versus $1/[\text{L-Hist}]$ ($r > 0.9998$, $S \leq 0.0152$) (Fig. 3). Since **scheme 1** is in accordance with the generally well-accepted principle of non-complementary oxidations taking place in sequence of one-electron steps, the reaction between the substrate and oxidant would afford a radical intermediate. A free radical scavenging experiment revealed such a possibility. This type of radical intermediate has also been observed in earlier work ²⁹.

Scheme 1 leads to the rate law equation (1),

$$-\frac{d[\text{MnO}_4^-]}{dt} = \text{rate} = \frac{kK_1K_2[\text{L-Hist}][\text{MnO}_4^-][\text{OH}^-]}{1+K_1[\text{OH}^-]+K_1K_2[\text{L-Hist}][\text{OH}^-]} \quad 1$$

$$k_{\text{obs}} = \frac{kK_1K_2[\text{L-Hist}][\text{OH}^-]}{1+K_1[\text{OH}^-]+K_1K_2[\text{L-Hist}][\text{OH}^-]} \quad 2$$

$$\frac{1}{k_{\text{obs}}} = \frac{1}{kK_1K_2[\text{L-Hist}][\text{OH}^-]} + \frac{1}{kK_2[\text{L-Hist}]} + \frac{1}{k} \quad 3$$

According to equation (3), the plots of $1/k_{\text{obs}}$ versus $1/[\text{OH}^-]$ ($r > 0.9989$, $S \leq 0.0121$) and $1/k_{\text{obs}}$ versus $1/[\text{L-Hist}]$ ($r > 0.9998$, $S \leq 0.0152$) should be linear as shown in figure 2 and 3. From the slopes and intercepts, the values of k , K_1 and K_2 could be derived as $3.43 \times 10^{-3} \pm 0.06 \text{ s}^{-1}$, $15.72 \pm 0.25 \text{ dm}^3 \text{ mol}^{-1}$ and $29.54 \pm 0.24 \times 10^2 \text{ dm}^3 \text{ mol}^{-1}$ respectively. Using these constants, the rate constants were calculated over different experimental conditions and there is a reasonable agreement between the calculated and experimental values (**Table 1**), which verifies the proposed mechanism.

The small effect of ionic strength on the reaction is presumably due to the fact that the reaction takes place between a neutral and charged species (**Scheme 1**). A plot of $\log k_{\text{obs}}$ against $I^{1/2}$ according to Debye-Huckel equation was linear with a positive slope. However, the present measurements, of necessity, lie far outside the Debye-Huckel region, covering a range over which the activity coefficient of many electrolytes are known to be fairly dependent on ionic strength ³⁰. The ionic strength dependence is qualitatively as expected when considering the charges involved ³¹. The effect of solvent on reaction rate was studied ³² by varying the t-butyl alcohol-water content in the reaction mixture with all other conditions being constant. Increasing the content of t-butyl alcohol in the reaction medium

leads to an increase in the rate of reaction. The plot of $\log k_{\text{obs}}$ versus $1/\varepsilon_T$ ($r>0.9998$, $S<0.0112$) is expected to be linear with a negative slope for a reaction between a negative ion and neutral molecule, where as positive slope results for a positive ion and neutral molecule. In the present study, a plot of $\log k_{\text{obs}}$ vs. $1/\varepsilon_T$ ($r>0.9998$, $S<0.0112$) is linear with negative slope (**Fig. 2**) which supports the involvement of negative ions as given in **Scheme 1**.

Table 1. Effect of variation of $[\text{KMnO}_4]$, $[\text{L-Hist}]$ and $[\text{OH}^-]$ on oxidation of L-histidine by alkaline KMnO_4 at 30°C , $I = 0.05 \text{ mol dm}^{-3}$.

$10^4 \times [\text{KMnO}_4^-]$ mol dm^{-3}	$10^3 \times [\text{L-Hist}]$ mol dm^{-3}	$10^3 \times [\text{OH}^-]$ mol dm^{-3}	$10^3 \times k_{\text{obs}} \text{ s}^{-1}$	
			Found	Calcd.*
0.4	2.0	5.0	1.04	1.03
0.6	2.0	5.0	1.03	1.03
1.0	2.0	5.0	1.05	1.03
2.0	2.0	5.0	1.03	1.03
4.0	2.0	5.0	1.02	1.03
6.0	2.0	5.0	1.03	1.03
2.0	0.8	5.0	0.50	0.50
2.0	1.0	5.0	0.61	0.61
2.0	2.0	5.0	1.05	1.03
2.0	4.0	5.0	1.59	1.58
2.0	6.0	5.0	1.88	1.93
2.0	8.0	5.0	2.18	2.17
2.0	2.0	2.0	0.52	0.52
2.0	2.0	4.0	0.90	0.89
2.0	2.0	5.0	1.05	1.03
2.0	2.0	8.0	1.35	1.36
2.0	2.0	10.0	1.53	1.53
2.0	2.0	20.0	2.27	2.01

The activation parameters of the reaction are calculated by a plot of $\log k$ versus $1/T$ as given in **Table 2**. The experimental values of $\Delta H^\ddagger$ and $\Delta S^\ddagger$ were both favorable for electron transfer processes. The high negative value of $\Delta S^\ddagger$ indicates that interaction of reacting ions of similar charges to form an activated complex and is more ordered than the reactants due to loss of degree of freedom. This also supports by a small value of frequency factor³³.

The hydroxyl ion concentration (as in **Table 1**) was verified at four different temperatures and the K_1 values were determined as $15.72 \text{ dm}^3 \text{ mol}^{-1}$, $17.77 \text{ dm}^3 \text{ mol}^{-1}$, $19.77 \text{ dm}^3 \text{ mol}^{-1}$, $21.99 \text{ dm}^3 \text{ mol}^{-1}$ at 30 , 35 , 40 and 45°C respectively (**Fig. 2**). Similarly L-histidine concentration as in **Table 1** was varied at four different temperatures and K_2 values were determined at each temperature (**Fig. 3**). The values of K_2 were obtained as $29.54 \times 10^2 \text{ dm}^3 \text{ mol}^{-1}$, $26.69 \times 10^2 \text{ dm}^3 \text{ mol}^{-1}$, and $23.64 \times 10^2 \text{ dm}^3 \text{ mol}^{-1}$, $21.12 \times 10^2 \text{ dm}^3 \text{ mol}^{-1}$ at 30 , 35 , 40 and 45°C respectively. A van't Hoff's plot was made for the variation of K_1 with temperature (i.e. $\log K_1$ versus $1/T$). Values of the enthalpy change of the first equilibrium step (K_1) of the reaction (ΔH), entropy of the reaction (ΔS) and free energy of reaction (ΔG) were calculated (**Table 2**) as $17.86 \pm 0.25 \text{ kJ/mol}$, $81.87 \pm 2.0 \text{ J K}^{-1} \text{ mol}^{-1}$ and -6.706 kJ/mol respectively. Similarly, thermodynamic parameters for second step (K_2) are calculated as $-18 \pm 2 \text{ kJ/mol}$, $7 \pm 1 \text{ J/mol}$, $-20 \pm 2 \text{ kJ/mol}$ for change in enthalpy (ΔH), change in entropy of the reaction (ΔS) and change in free energy of the reaction (ΔG) respectively (**Fig. 4**). The proposed mechanism is also

supported by the above thermodynamic parameters. The latter values with those obtained for slow step of the reaction shows that these values mainly refer to the rate limiting step, supporting the fact that the reaction before the rate determining step are fairly slow and involves high activation energy³⁴.

Figure 1. Plot of $\log k_{\text{obs}}$ versus $\log 1/D$ and $\log k_{\text{obs}}$ versus $I^{1/2}$.

Table 2. Activation parameters for the oxidation of L-histidine by alkaline permanganate $[\text{KMnO}_4] = 2.0 \times 10^{-4} \text{ mol dm}^{-3}$, $[\text{L-Hist}] = 2.0 \times 10^{-3} \text{ mol dm}^{-3}$, $[\text{OH}^-] = 0.005 \text{ mol dm}^{-3}$ and $I = 0.05 \text{ mol dm}^{-3}$.

Activation Parameters	Values
E_a (kJ mol ⁻¹)	29.0 ± 1.4
$\Delta H^\ddagger$ (kJ mol ⁻¹)	26.5 ± 1.3
$\Delta S^\ddagger$ (J K ⁻¹ mol ⁻¹)	-49.7 ± 2.2
$\Delta G^\ddagger$ (kJ mol ⁻¹)	41.5 ± 1.5
Log A	5.01

Variation in the rate within a reaction series may be caused by change in the enthalpy and or entropy of activation. Changes in the rate are caused by changes in both $\Delta H^\ddagger$ and $\Delta S^\ddagger$, but these quantities vary extensively in a parallel fashion. A plot of $\Delta H^\ddagger$ versus $\Delta S^\ddagger$ is linear according to equation, $\Delta H^\ddagger = \beta \Delta S^\ddagger + \text{constant}$, β is called the isokinetic temperature; it has been asserted that apparently linear correlation of $\Delta H^\ddagger$ with $\Delta S^\ddagger$ are some times misleading and the evaluation of β by means of the above equation lacks statistical validity³⁵. Exner³⁶ advocates an alternative method for the treatment of experimental data. If the rates of several reactions in a series have been measured at two temperatures and $\log k_2$ (at T_2) is linearly related to $\log k_1$ (at T_1) (Fig 5.) i.e. $\log k_2 = a + b \log k_1$, he proposes that β can be evaluated from the equation,

Figure 2. Rate law plots ($1/k_{\text{obs}}$ versus $1/[\text{OH}^-]$) of oxidation of L-Histidine by potassium permanganate in aqueous alkaline medium at different temperatures. (conditions as in Table 1)

Figure 3. Rate law plots ($1/k_{\text{obs}}$ versus $1/[\text{L-histidine}]$) of oxidation of L-Histidine by potassium permanganate in aqueous alkaline medium at different temperatures. (conditions as in Table 1)

Table 3. Activation parameters for oxidation of some amino acids by alkaline KMnO_4 .

Amino acid	$k_1 \times 10^2$ s^{-1} at 298 K	$k_2 \times 10^2$ s^{-1} at 303 K	$\Delta S^\ddagger$ $\text{J K}^{-1} \text{mol}^{-1}$	$\Delta H^\ddagger$ kJ mol^{-1}	$\Delta G^\ddagger$ kJ mol^{-1}	References
Rac-Serine	10.00	28.00	-28 ± 2.4	49 ± 1	58 ± 3	38
L(+)-Aspartic acid	2.22	8.84	-83 ± 10	40 ± 3	64.7 ± 3	39
Glutamic acid	4.00	6.13	-86 ± 4	45 ± 2.5	70.6 ± 1	40
L-histidine	0.27	0.34	-49.7 ± 2.2	26.5 ± 1.3	41.5 ± 2	Present work

$$\beta = T_1 T_2 (b-1) / (T_2 b - T_1)$$

We have calculated the isokinetic temperature, β as 277.7 K by plotting $\log k_2$ at 303 K versus $\log k_1$ at 298 K (Fig. 5) ($r \geq 0.979$ & $s \leq 0.0051$) for a series of α -amino acids like rac-serine, L-(+) aspartic acid, L-glutamic acid and present work. The value of β (277.7 K) is smaller than the experimental temperature (303K). This indicates that the rate is governed by the enthalpy of activation³⁷. The linearity and the slope of the plot obtained may confirm that the kinetics of these reactions follow similar mechanisms, as previously suggested (free radical intervention). However, the higher value of $S^\ddagger$ of L (+)-aspartic acid and L-glutamic acid is due to the stabilization of carbocation by successive CH_3 group(s) which is unlike in L-histidine.

Figure 4. Effect of temperature on KMnO_4 oxidation of L-histidine in aqueous alkaline medium for mechanism of step 1 and 2 as per Scheme 1.

Figure 5. Plot of $\log k_2$ at 303 K vs. $\log k_1$ at 298 K for a series of amino acids (Table 3)
1. Rac-Serine, 2. L-(+)-Aspartic acid, 3. L-Glutamic acid and 4. L-histidine.

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