

Kinetics and mechanism of the oxidation of some neutral and acidic α -amino acids by tetrabutylammonium tribromide

RAGHVENDRA SHUKLA^a, PRADEEP K SHARMA^a and KALYAN K BANERJI^{b*}

^aDepartment of Chemistry, JNV University, Jodhpur 342 005, India

^bFaculty of Science, National Law University, Mandore (NH 65), Jodhpur 342 304, India

e-mail: banerjikk@rediffmail.com

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Abstract. The oxidation of eleven amino acids by tetrabutylammonium tribromide (TBATB) in aqueous acetic acid results in the formation of the corresponding carbonyl compounds and ammonia. The reaction is first order with respect to TBATB. Michaelis–Menten type kinetics is observed with some of the amino acids while others exhibit second-order dependence. It failed to induce polymerization of acrylonitrile. The effect of solvent composition indicate that the rate of reaction increases with increase in the polarity of the medium. Addition of tetrabutylammonium chloride has no effect on the rate of oxidation. Addition of bromide ion causes decrease in the oxidation rate but only to a limiting value. The reaction is susceptible to both polar and steric effects of the substituents. A suitable mechanism has been proposed.

Keywords. Amino acid; tetraalkylammonium tribromide; kinetics; mechanism; oxidation.

1. Introduction

Tetraalkylammonium polyhalides are widely used as halogenating reagents in synthetic organic chemistry.^{1–3} Tetrabutylammonium tribromide (TBATB) has been used for the bromination of some selected organic substrates.⁴ There are, however, only a few reports regarding their use as oxidizing and brominating agents in synthetic chemistry.^{5–7} These compounds are more suitable than molecular halogens because of their solid nature, ease of handling, stability, selectivity and excellent product yields. We have been interested in the kinetic and mechanistic studies of the reactions of polyhalides and many reports including those on TBATB have already emanated from our laboratory.^{8–10} There seems to be no report on the oxidation of α -amino acids by TBATB. A study of the kinetics of the oxidation of amino acids is important both from a mechanistic point of view and its bearing on the mechanism of amino acid metabolism. Therefore, we have studied the oxidation of several neutral and acidic α -amino acids by TBATB in aqueous acetic acid solution, and the mechanistic aspects are discussed in this paper.

2. Experimental

2.1 Materials

All the amino acids were commercial products of highest degree of purity and were used as supplied. Perdeuterioglycine ($\text{ND}_2\text{CD}_2\text{COOD}$) was obtained from Sigma Chemicals (USA). TBATB was prepared by the reported method¹ and its purity was checked by an iodometric method. Acetic acid was refluxed with chromic oxide and acetic anhydride for 3 h and then fractionally distilling it.¹¹ All other reagents were commercial products and were purified by the usual methods.¹¹

2.2 Product analysis

The main products of the amino acids were the corresponding carbonyl compounds and ammonia. The presence of ammonia in the reaction mixture was detected by the test with *p*-nitrobenzenediazonium chloride.¹²

In a typical experiment, *a*-alanine (4.45 g, 0.05 mol) and TBATB (4.45 g, 0.01 mol) were made up to 100 ml in 1:1 (v/v) acetic acid–water. The mixture was allowed to stand for \approx 15 h in the dark to ensure

*For correspondence

completion of the reaction. It was then treated with an excess (250 ml) of a saturated solution of 2,4-dinitrophenylhydrazine in 2 mol dm⁻³ HCl and kept in a refrigerator for ≈ 10 h. The precipitated 2,4-dinitrophenylhydrazone (DNP) was filtered off, dried, weighed, recrystallized from ethanol and weighed again. The yields of DNP before and after recrystallization were 2.24 g (95%) and 1.97 g (83%) respectively, based on the consumption of TBATB. The DNP was found to be identical (m.pt. and mixed m.pt.) with the DNP of acetaldehyde. In similar experiments, with other amino acids, the yields of the DNP were in the range of 80–89% after recrystallization.

Stoichiometry, with an excess of amino acid could not be studied because of the difficulty in determining the concentration of amino acids. Stoichiometric determination with an excess of TBATB showed that two moles of TBATB were consumed per mole of the amino acid. This is because aldehydes, the initial products of the oxidation, were further oxidized to carboxylic acids.

2.3 Kinetic measurements

The reactions were studied under pseudo-first-order conditions by keeping an excess (× 15 or greater) of the substrate over TBATB. The solvent was 1:1 (v/v) acetic acid–water, unless mentioned otherwise. Tribromide ion is known to dissociate to a large extent to bromine and bromide ion. The value¹³ of the dissociation constant in 1:1 (v/v) acetic acid–water is ≈ 0.02 mol dm⁻³. To suppress the dissociation, all kinetic runs were carried out in the presence of an excess (0.3 mol dm⁻³) of potassium bromide. The reactions were studied at constant temperature (± 0.1 K) and were followed by monitoring the decrease in the concentration of TBATB at 394 nm for up to 80% reaction. Pseudo-first-order rate constants, k_{obs} , were evaluated from linear plots ($r > 0.995$) of log[TBATB] against time. Duplicate kinetic runs showed that the rate constants are reproducible to within ± 3%. Simple and multivariate regression analyses were carried out by the least-squares method.

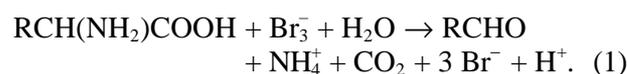
3. Results

Preliminary experiments showed that the reactions are not sensitive to changes in the ionic strength. Hence no attempt was made to keep the ionic strength constant.

The rate laws and other experimental data for all the eleven amino acids studied. Since the results are similar, only the representative data are reproduced here.

3.1 Stoichiometry

The overall reaction may be represented as under.



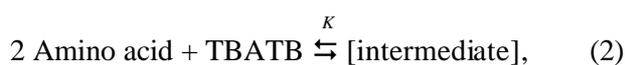
3.2 Rate laws

The reaction is first order with respect to TBATB. The individual kinetic runs obeyed first-order kinetics. Further, the values of k_{obs} are independent of the initial concentration of TBATB. The order is more than one but less than two in respect of glycine (Gly), α -alanine (Ala), 2-aminobutanoic acid (ABA), norleucine (NLE), norvaline (NVA) and phenylalanine (Phe) whereas the order with respect to valine (Val), leucine (Leu), isoleucine (Ile), aspartic acid (Asp) and glutamic acid (Glu) is two. The rate constants with respect of Ala and Leu are recorded in table 1. Plots of $1/k_{\text{obs}}$ against $1/[\text{amino acid}]^2$ for Gly, Ala, ABA, NLE, NVA and Phe are linear ($r^2 > 0.995$) with an intercept on the rate ordinate. Thus Michaelis–Menten type kinetics are observed with respect to these amino acids. This leads to the postulation of the following overall mechanism and rate law.

Table 1. Rate constants for the oxidation of α -alanine and leucine by TBATB at 308 K.

$10^3[\text{TBATB}]$ mol dm ⁻³	[Ala] mol dm ⁻³	[Leu] mol dm ⁻³	$10^4 k_{\text{obs}} \text{ (s}^{-1}\text{)}$	
			Ala	Leu
1.0	0.10	0.10	1.87	1.85
1.0	0.20	0.20	6.70	7.51
1.0	0.40	0.30	19.0	16.6
1.0	0.75	0.40	33.5	30.1
1.0	1.00	0.50	38.9	46.5
1.0	1.50	0.75	43.3	107
1.0	2.00	1.00	46.3	190
2.0	0.40	0.40	18.7	30.5
4.0	0.40	0.40	19.1	29.5
6.0	0.40	0.40	19.4	30.3
8.0	0.40	0.40	19.2	30.0
1.0	0.20	0.20	6.75*	7.61*

*Contained 0.005 mol dm⁻³ acrylonitrile



$$\text{rate} = K k_2 [\text{amino acid}]^2 [\text{TBATB}] / (1 + K [\text{amino acid}]^2). \quad (4)$$

It is proposed that the oxidation of Val, Leu, Ile, Asp and Glu also follow similar mechanism but in their case, the equilibrium constants have very small values. Thus in these cases, $K [\text{amino acid}]^2 \ll 1$ and no Michaelis–Menten type kinetics is observed but a second-order dependence is obtained. The reason for this must be steric. The amino acids exhibiting second-order dependence have bulkier substituents as compared to the ones showing Michaelis–Menten kinetics. The variation in amino acid concentration was studied at different temperatures and the values of K and k_2 evaluated from the double reciprocal plots. The thermodynamic and activation parameters

for the formation and disproportionation of the intermediate were calculated from the values of K and k_2 respectively, at different temperatures (tables 2 and 3). The reaction (2) is likely to take place in two steps. Therefore, the equilibrium constant, K , might be a composite quantity.

The dependence of reaction on hydrogen-ion concentration could not be determined by an addition of a mineral acid like perchloric acid as TBATB decomposed rapidly in the presence of a mineral acid. Therefore, effect of sodium acetate on the reaction rate was studied. Results showed that addition of sodium acetate has no effect on the rate of the oxidation (table 4).

3.3 Induced polymerization of acrylonitrile

The oxidation of amino acids, in an atmosphere of nitrogen, failed to induce the polymerization of acrylonitrile. Further, the addition of acrylonitrile had no effect on the rate of oxidation (table 1).

Table 2. Formation constants of the TBATB-amino acid complexes and the thermodynamic parameters.

Substrate	K (dm ³ mol ⁻¹) at				ΔH	ΔS	ΔG
	298 K	308 K	318 K	328 K	kJ mol ⁻¹	J mol ⁻¹ K ⁻¹	kJ mol ⁻¹
H	3.20	2.72	2.21	1.75	-18.9 ± 0.8	-45 ± 3	-5.4 ± 0.6
Me	4.85	4.03	3.27	2.55	-19.8 ± 0.8	-45 ± 2	-6.5 ± 0.6
PhCH ₂	3.97	3.34	2.72	2.16	-19.0 ± 0.7	-44 ± 2	-6.0 ± 0.6
ABA	5.32	4.12	3.35	2.60	-21.6 ± 0.4	-51 ± 1	-6.6 ± 0.4
NLE	6.03	5.05	4.10	3.26	-19.2 ± 0.7	-41 ± 2	-7.0 ± 0.5
NVA	4.13	3.70	3.35	2.91	-12.8 ± 0.6	-20 ± 2	-6.0 ± 0.5

Table 3. Rate of the decomposition of the intermediate complexes and the activation parameters.

Substrate	$10^4 k_2$ (s ⁻¹) at				ΔH^*	ΔS^*	ΔG^*
	298 K	308 K	318 K	328 K	kJ mol ⁻¹	J mol ⁻¹ K ⁻¹	kJ mol ⁻¹
Gly	2.23	4.81	10.6	21.3	58.9 ± 0.5	-99 ± 2	88.2 ± 0.4
Ala	27.3	48.3	86.4	148	43.4 ± 0.4	-130 ± 1	82.0 ± 0.3
ABA	36.8	63.9	110	185	41.3 ± 0.5	-135 ± 2	81.2 ± 0.5
NVA	55.1	92.1	151	256	38.9 ± 0.7	-139 ± 3	80.2 ± 0.6
NLE	59.9	97.5	164	271	38.8 ± 0.6	-139 ± 2	80.1 ± 0.4
Phe	87.5	140	215	347	34.5 ± 0.7	-150 ± 2	79.1 ± 0.6
$10^4 k_2$ (dm ⁶ mol ⁻² s ⁻¹)							
Val	71.7	115	195	308	37.3 ± 0.6	-142 ± 2	79.6 ± 0.5
Leu	121	187	300	472	34.5 ± 0.7	-148 ± 2	78.3 ± 0.5
Ile	180	285	440	671	33.1 ± 0.6	-149 ± 2	77.3 ± 0.6
Asp	6.83	16.3	33.4	70.1	56.4 ± 1.0	-97 ± 4	85.3 ± 1.0
Glu	22.2	38.7	75.2	134	46.7 ± 1.1	-121 ± 4	82.5 ± 0.9

Table 4. Effect of sodium acetate on the rate of oxidation of leucine by TBATB.[TBATB] = 0.001 mol dm⁻³; [Leu] = 0.40 mol dm⁻³; Temp. = 308 K.

[NaOAc] (mol dm ⁻³)	0.00	0.10	0.20	0.30	0.40	0.50
10 ⁴ k _{obs} (s ⁻¹)	30.1	29.8	30.7	29.3	31.5	30.4

Table 5. Effect of tetrabutylammonium chloride on the oxidation of leucine by TBATB.[TBATB] = 0.001 mol dm⁻³; [Leu] = 0.40 mol dm⁻³; Temp. = 308 K.

10 ³ [TBACl] (mol dm ⁻³)	0.00	0.5	1.0	2.0	3.0	4.0
10 ⁴ k _{obs} (s ⁻¹)	30.1	30.3	29.5	29.9	30.5	30.0

Table 6. Dependence of rate of oxidation of leucine on the concentration of bromide ions.[TBATB] = 0.001 mol dm⁻³; [Leu] = 0.40 mol dm⁻³; Temp. = 308 K.

[Br ⁻] (mol dm ⁻³)	0.0	0.04	0.08	0.12	0.16	0.20	0.24	0.30
10 ⁴ k _{obs} (s ⁻¹)	30.1	26.9	22.6	19.8	16.0	15.8	15.7	15.8

Table 7. Dependence of the rate of oxidation of leucine by TBATB on solvents compositions.[TBATB] = 0.001 mol dm⁻³; [Leu] = 0.40 mol dm⁻³; Temp. = 308 K.

% AcOH (v/v)	25	40	50	60	70
10 ⁴ k _{obs} (s ⁻¹)	53.8	38.3	30.1	26.4	22.7

3.4 Effect of tetrabutylammonium ions

The rates of oxidation were not affected by the addition of tetrabutylammonium chloride (TBACl) (table 5).

3.5 Effect of bromide ions

The rate of oxidation decreases with an increase in the concentration of potassium bromide but reaches a limiting value at [KBr] ≈ 0.16 mol dm⁻³ (table 6).

3.6 Effect of solvent composition

The rate of oxidation was determined in solvents containing different amounts of acetic acid and water. It was observed that the rate increased with an increase in the amount of water in the solvent mixture (table 7).

4. Discussion

An isokinetic plot between activation enthalpies and entropies of oxidation of the eleven amino acids is

reasonably good ($r^2 = 0.9887$, s.d. = 0.97). The value of isokinetic temperature is 460 ± 16 K. The isokinetic relationship was verified and found genuine by applying Exner's¹⁴ criterion. The Exner plot between $\log k_2$ at 298 K and $\log k_2$ at 328 K is linear ($r^2 = 0.9969$, s.d. = 0.02; slope = 0.7571 ± 0.0140). The isokinetic temperature is 496 ± 14 K, which is in very good agreement with the value obtained from the activation parameter data. The linear isokinetic correlation implies that all the amino acids are oxidized by the same mechanism and the changes in the rate are governed by the changes in both the enthalpy and entropy of the activation.

4.1 Reactive oxidizing species

We have earlier⁸ carried out some conductivity measurements to determine the nature of TBATB in aqueous acetic acid solution. It was observed that acetic acid has very low conductivity. Addition of TBATB increases the conductivity of the solution. We measured the conductivity of TBATB in solvents containing different proportions of acetic acid (100–30%) and water. We found that the conductivity increases sharply as the water content is increased and

Table 8. Temperature dependence of the reaction constants.

Temp.(K)	r^*	d	R^2	s.d.	y
298	-0.77 ± 0.01	-0.58 ± 0.01	0.9997	0.01	0.02
308	-0.64 ± 0.02	-0.55 ± 0.01	0.9988	0.02	0.04
318	-0.57 ± 0.01	-0.51 ± 0.01	0.9994	0.01	0.03
328	-0.52 ± 0.02	-0.47 ± 0.01	0.9998	0.01	0.02

The rate constants of the aliphatic amino acids failed to give any significant correlation with Taft's polar and steric effects separately. The rates were therefore analysed in terms of Pavelich-Taft's¹⁶ dual substituent parameter (10).

$$\log k_2 = -1.12 \pm 0.29 S^* - 2.33, \quad (8)$$

$$r^2 = 0.6518; \text{ s.d.} = 0.36; y = 0.62; \\ n = 10; T = 298 \text{ K.}$$

$$\log k_2 = -0.77 \pm 0.16 E_s - 2.70, \quad (9)$$

$$r^2 = 0.7439; \text{ s.d.} = 0.31; y = 0.53; \\ n = 10; T = 298 \text{ K.}$$

$$\log k_2 = S^*r^* + dE_s + \log k_0. \quad (10)$$

Here n is the number of data points and y is Exner's statistical parameter.¹⁷

The correlations in terms of (9) are excellent (table 8). The reaction constants have negative values. The analysis showed that the reaction is susceptible to both the polar and steric effects of the substituents. The values of the reaction constants support the proposed mechanism. The negative polar reaction constant is in accordance with the net flow of electrons towards the oxidant. An increase in the electron density at the reaction centre facilitates the flow of the electrons from substrate towards the oxidant. The negative steric reaction constant indicates a steric acceleration. This may well be due to high ground state energy of the more substituted amino acids. The steric crowding is relieved in the product $(RCH=NH_2)^+$ and as well as in the transition leading to it. This coupled with the fact that there is not much difference in the transition state energy of the crowded and uncrowded molecules explain the steric acceleration.

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