

KINETICS OF OS(VIII) CATALYSED OXIDATION OF BENZALDEHYDE ANILS BY PDC – STUDY OF THE EFFECTS OF SUBSTRATE

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ABSTRACT : The kinetic study of simple and substituted anils were studied under various conditions viz., solvent, oxidants, electrolyte, catalysts and mixture of catalysts and temperature; simple and substituted anils, pyridinium Chloro Chromate(PCC), Pyridinium Dichromate (PDC) were prepared in the laboratory and kinetic study were carried out using Pyridinium dichromate, substrate, perchloric acid and osmium tetroxide. The graphs were drawn at all relevant places and results were obtained in full satisfaction.

KEYWORDS: simple and substituted anils, Pyridinium Chlorochromate [PCC], Pyridinium Dichromate [PDC], Osmiumtetroxide, oxidation

INTRODUCTION

Among oxo derivatives of variable valence metals, Chromium compounds play the most important role, in oxidative reaction. A number of Chromium reagents are readily available. Almost every oxidisable functional group may undergo Chromium oxidation. Chromium (VI) containing reagents include Chromium acid, dichromate ion, Chromyl chloride, chromyl acetate, t-butyl Chromate, Chromyl nitrate and Co-ordination complexes of Chromium trioxide.¹⁻²

Chromium (VI) oxidations are usually performed under acidic conditions. Co-solvents like (Jones's reagent) benzene, methylenechloride (Two Phase system) are often added in order to deal with water insoluble organic complexes. The oxidation rate is generally high under acid catalysed conditions, however the low pH of the reaction medium and the presence of water favouring hydrolytic conditions exclude the use of this class of reagents for the oxidation of molecules containing acid sensitive groups.

Polymer supported Chromium (VI) reagents have also been developed. These reagents offer the advantage of reducing the work of procedure to more filtration. Recently some neutral or almost neutral Chromium (VI) reagents have been developed to effect oxidation under mild conditions.

Pyridinium Chlorochromate (PCC) introduced by Corey et al³ is widely used in the oxidation of alcohols. Pyridinium chlorochromate has a less pronounced acidity and is an effective agent for the oxidation of polycyclic organic substrates. PCC was developed by Bhattacharjee and coworker.⁴

In 1986, Narayanan and Balasubramanian⁵ introduced Pyridinium bromochromate. This is an efficient oxidant for alcohols and a brominating agent as well. Firouzahadi and Coworkers used Zinc chromates. This is a very useful reagent for the oxidation of variety of organic compounds, including alcohols, oximes, olefins and aromatic hydrocarbons.

They have also shown chromium peroxide complexes as versatile, mild and efficient oxidants in organic synthesis. Chlorotrimethyl Silane, Imidazolium dichromate, 2,2'-Bipyridyl chlorochromate (BPCC) and the corresponding chromate (BPC) have been proposed as oxidants for the oxidation of the hydroxyl group to the carbonyl group. 4-(dimethylamino)-pyridinium chlorochromate and chromyl nitrate have been shown to be efficient and mild oxidants in aprotic media. Naphthylidinium chlorochromate, Pyrazinium chlorochromate, tetrabutyl ammonium chlorochromate, dimethylpyrazole chromate, tetrabutyl ammonium dichromate, Pyridine complex of oxodiperoxochromium and the dipyridine complex of chromium trioxide are some of the Chromium (VI) oxidants introduced in the last twenty years. Quinolinium Chlorochromate was used by Bhavani et al to oxidize methyl phenyl sulphoxides. Quinolinium dichromate was introduced by Balasubramanian and Prathiba⁶ as an effective oxidant under non-aqueous conditions.

Pyridium Chlorochromate (PCC) is irreplaceable in the generation of functional groups in highly unsaturated carbinols and pyrrols. yet mildly acidic character of Pyridinium chlorochromate precludes its use with acid sensitive substrates or products. Pyridium Chlorochromate oxidation has been investigated under various condition on different substrates. Despite the introduction of numerous oxidants based on chromium (VI) not much work has been done to investigate the mechanism of oxidation in these

1.1 Reactions of Pyridinium Dichromate

Pyridium dichromate (PDC) in N,N – dimethyl formamide or dichloromethane is nearly neutral non – aqueous chromium (VI) oxidant. This species probably present in the Sarret and Conferth oxidizing mixtures. Although specific reference has been made to oxidation of alcohols by Pyridium dichromate itself in a short note, there is no indication of any unusual effectiveness, utility, or advantage to be gained by the preformed pyridium dichromate as a discrete oxidizing species before Corey and Schmidt¹ prepared it in 1979. Solutions of Pyridium dichromate in DMF rapidly oxidize allylic alcohols⁴ to alpha, beta – unsaturated carbonyl compounds at 0 °C. Kinetics and mechanism of oxidation of cyclic ketoximes by Pyridium dichromate and Quinolinium dichromate was investigated by Chidambaranathan et al.². Oxidation kinetics of Cyclanols by Pridiniumdichromate was studied by Padma et al.³. Kinetics and mechanism of oxidation of some *ortho*- substituted benzyl alcohol by PDC was studied by Kabilan and Thenmozhi.⁴ Kinetics and mechanism of oxidation of some substituted phenylmethyl sulphoxide using PDC was reported by Meenakshisundaram et al.⁵ Oxidation of S – phenylmercaptoacetic acid and phenoxyacetic acid by PDC was studied by Kabilan and Krishnasamy.⁶ Gurumurthy et al.⁷ investigated a comparative study on the kinetics of oxidation of some secondary alcohols by 1 – chlorobenzotriazole and PDC.

1.2 Reactions of Anils

Several oxidation of anils have been studied :

1.2.1 Effect of varying PCC concentration

The reaction was investigated by varying the concentration at constant substrate concentration. The reaction was found to be first order with respect to PCC [Table – 1.2]

Table -1.2

[Anil] = 1.5 x 10⁻² M ACOH :H₂O = 70:30 (V/V)
 [H₂C₂O₄] = 5.0 x 10⁻² M [HClO₄] = 3.87 x 10⁻³ M
 Temp. = 35 °c

[PCC] x 10 ³	Kobs. X 10 ⁴ s ⁻¹
1.2	1.932
1.5	1.921
2.00	1.811

1.2.2 Effect of varying the concentration of Anil

The reaction was carried out under pseudo – first order conditions. In this study, the concentration of PCC, Oxalic acid, Perchloric acid and percentage of oxalic acid were kept constant and the concentration of the anil varied. The reaction was found to be first order with respect to anil. [Table – 1.3]

Table – 1.3

[PCC] = 1.5 x 10⁻³ M ACOH :H₂O = 70:30 (V/V)
 [H₂C₂O₄] = 5.0 x 10⁻² M Temp. = 35 °c
 [HClO₄] = 3.87 x 10⁻³ M

[ANIL] x 10 ²	Kobs. X 10 ⁴ s ⁻¹
1.00	1.08
1.50	1.923
2.50	2.785
3.00	3.512

1.2.3 Effect of varying solvent Composition

The reaction was studied by the different composition of acetic acid under constant [Reactants]. It was found that as the percentage of acetic acid increased, the rate decreased. [Table – 1.4]

Table – 1.4

$$\begin{aligned}
 [\text{ANIL}] &= 1.5 \times 10^{-3} \text{ M} & [\text{HClO}_4] &= 3.87 \times 10^{-3} \text{ M} \\
 [\text{H}_2\text{C}_2\text{O}_4] &= 5.0 \times 10^{-2} \text{ M} & \text{Temp.} &= 35^\circ \text{ c}
 \end{aligned}$$

% AcOH	Kobs. X 10 ⁴ s ⁻¹
40	3.921
50	3.207
60	3.256
70	1.932

It was found that the rate decreases with increase of percentage of acetic acid

1.2.4 Effect of varying Oxalic acid

The reaction was carried out under by varying , the concentration of , Oxalic acid and by keeping other [reactants] constant. The result indicate that as the concentration of oxalic acid increased , the rate constant also increased [Table – 1.5].

Table – 1.5

$$\begin{aligned}
 [\text{ANIL}] &= 1.5 \times 10^{-2} \text{ M} & \text{ACOH :H}_2\text{O} &= 70:30 \text{ (V/V)} \\
 [\text{HClO}_4] &= 3.87 \times 10^{-3} \text{ M} & \text{Temp.} &= 35^\circ \text{ c}
 \end{aligned}$$

[H ₂ C ₂ O ₄] x 10 ⁴	Kobs. X 10 ⁴ s ⁻¹
5.0	1.932
7.5	2.872
10.0	3.871
12.	5.965

1.2.5. Effect of varying Perchloric acid

The reaction was studied by varying , the concentration of perchloric acid and by keeping other [reactants] constant. . The result indicate that as the concentration of Perchloric acid increased , the rate constant also increased. [Table – 1.6]

Table -1.6

$$\begin{aligned}
 [\text{ANIL}] &= 1.5 \times 10^{-2} \text{ M} & \text{ACOH :H}_2\text{O} &= 70:30 \text{ (V/V)} \\
 [\text{H}_2\text{C}_2\text{O}_4] &= 5.0 \times 10^{-2} \text{ M} & \text{Temp.} &= 35^\circ \text{ c}
 \end{aligned}$$

[HClO ₄] x 10 ⁴	Kobs. X 10 ⁴ s ⁻¹
3.90	1.932
5.80	2.785
7.70	2.952
9.70	4.609

1.2.6. Effect of varying Sodium Perchlorate

To follow the Primary Salt effect , the reaction was studied with varying concentration of Sodium Perchlorate by keeping other [reactants] constant. It was found that as the concentration of Sodium Perchlorate increased , the rate constant also increased. This shows that the participation of ions which are similar in their sign or dipole –ion interaction in the rate determining step. [Table 1.7]

Table – 1.7

$$\begin{aligned}
 [\text{ANIL}] &= 1.5 \times 10^{-2} \text{ M} & \text{ACOH :H}_2\text{O} &= 70:30 \text{ (V/V)}
 \end{aligned}$$

$$\begin{aligned}
 [\text{H}_2\text{C}_2\text{O}_4] &= 5.0 \times 10^{-2} \text{ M} & [\text{HClO}_4] &= 3.87 \times 10^{-3} \text{ M} \\
 [\text{PCC}] &= 1.2 \times 10^{-3} \text{ M} & \text{Temp.} &= 35^\circ \text{ c}
 \end{aligned}$$

Table 2.5

[NaClO ₄] X 10 ³	K _{obs} x 10 ⁴ S ⁻¹
0.00	1.932
2.50	3.548
5.00	4.657
7.50	5.086
10.00	5.799

1.2.7.Effect of MnSO₄

The reaction rate decreases tremendously with the increase in the concentration of MnSO₄ [Table - 1.8] This may be due to the formation of Cr(VI) in the rate determining step.

Table - 1.8

$$\begin{aligned}
 [\text{ANIL}] &= 1.5 \times 10^{-2} \text{ M} & \text{AcOH} : \text{H}_2\text{O} &= 70 : 30 \text{ (V/V)} \\
 [\text{H}_2\text{C}_2\text{O}_4] &= 5.0 \times 10^{-2} \text{ M} & [\text{PCC}] &= 1.2 \times 10^{-3} \text{ M} \\
 [\text{HClO}_4] &= 3.87 \times 10^{-3} \text{ M} & \text{Temp.} &= 35^\circ \text{ C}
 \end{aligned}$$

[MnSO ₄] X 10 ⁴	K _{obs} x 10 ⁴ S ⁻¹
0.00	1.932
1.00	1.053
2.00	0.723
3.00	0.575

1.2.8 Effect of temperature

The rate of oxidation of some *meta* and *para* substituted anils had been studied at four different temperatures viz., 35⁰ c , 40⁰ c, 45⁰ c and 50⁰ c . It was observed , as we expected ,that the rate increases very much with increase of temperature .[Table – 1.9]

Table – 1.9

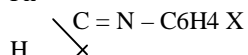
$$\begin{aligned}
 [\text{Substrate}] &= 1.5 \times 10^{-2} \text{ M} & \text{ACOH} : \text{H}_2\text{O} &= 70:30 \text{ (V/V)} \\
 [\text{H}_2\text{C}_2\text{O}_4] &= 1.0 \times 10^{-4} \text{ M} & [\text{PCC}] &= 1.2 \times 10^{-3} \text{ M} \\
 [\text{HClO}_4] &= 4.0 \times 10^{-3} \text{ M} & \text{Temp.} &= 35^\circ \text{ c}
 \end{aligned}$$

S.No.	Substituents	Kobs. X 10 ⁴ s ⁻¹			
		35 ⁰	40 ⁰	45 ⁰	50 ⁰
2	H	1.932	3.335	4.127	6.735
3	P -CH ₃	3.91	6.123	8.315	10.371
4	P - O CH ₃	6.22	9.132	12.612	16.431
5	p- Cl	1.023	2.425	3.671	4.923
6	m - Cl	0.705	1.412	2.213	3.824

II.EXPERIMENTAL

2.1. General method of preparation of Anils

2.1. Ph



Where X = H , p - CH₃ , p - OCH₃ , p - Cl , p - Br , P - COOH ,
 p - NO₂ , m - CH₃ , m - Cl , m - NO₂

The anils were prepared 87a by refluxing equimolar quantities of benzaldehyde and aniline or substituted anilines in alcohol for about 2 to 3 hrs. The resulting solution was cooled and poured into the cold water . The precipitated anil was filtered , washed , dried and recrystallised from alcohol. The purity of anils were checked by determining their melting points (Table 2.1).

Table – 2.1

S.No.		m.pt. 0 C

	R'	R'	Observed	Literature
1	H	H	54.5	54
2	p - CH ₃	H	36.0	35
3	p -OCH ₃	H	61.0	60
4	p - Cl	H	62.5	62
5	p -Br	H	57.0	56 - 57
6	PCOOH	H	134.0	135
7	p - NO ₂	H	121.0	120
8	m - Cl	H	58.5	58
9	m - CH ₃	H	114.0	115
10	m- NO ₂	H	65.0	66

2.2.PREPARATION OF PYRIDINIUM DICHROMATE [PDC]

Pyridine (80.6 ml) was gradually added to a cooled solution of 100 gm (1 mol) of Cr₂O₃ in 100 ml water at 30 C . The solution was distilled with 400 ml of acetic acetone and cooled to 20 c . After 3 hrs. the orange crystals were collected , washed with acetone and dried in vacuuo.

2.3.PRODUCT ANALYSIS

The reactants in the ratio anil : PDC (1 : 10) in the 70 % acetic acid were mixed under kinetic conditions . The products of oxidation were identified as the corresponding benzaldehyde and nitrosobenzene . benzaldehyde was characterized as the 2, 4 - dinitrophenyl hydrozone derivative . The remaining solution on evaporation yielded nitrozobenzene , which was identified by its characteristic UV spectrum .

2.4. STOICHIOMETRY

The stoichiometry of the reaction was detmrined by carrying out several sets of experiments with varying amounts of [PDC] largely in excess over [anil]. The estimation of unreacted PCC showed that one mole of anil reacts with two mole of PDC [ie.1 : 1)

III.RESULT AND DISCUSSION

Kinetics and mechanism of several organic substrates by PDC were documented ; but there are no systematic kinetics reports of the oxidation of Osmium (VIII) . Catalysed oxidation of benzaldehyde anils by Cr (VI) Complexes. This prompted us to undertake the title for investigation.

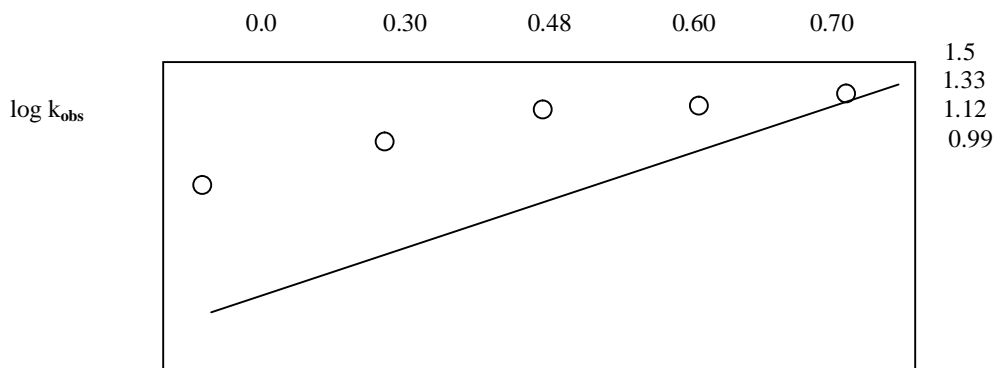
3.1.Effect of varying the of substrate

The concentration of Anil was varied 1.0 - 5.0 /10⁻² M at fixed concentrations other reaction components. The rate of oxidation increases with increase of [Anil] [Table 3.2] Plots of log k_{obs} versus log [Anil] gave straight line withslope equal to 1.09 [Fig.1]. hence the order with respect to [substrate] is uity.

Table 3.2

[PDC] = 1.0 X 10⁻³ M AcOH : H₂O = 70 : 30 (V/V)
 [HClO₄] = 7.0 X 10⁻³ M [OsO₄] = 1.0 X 10⁻⁵ M
 Temp. = 40⁰ C

[Anil] X 10 ² M	k _{obs} x 10 ³ S ⁻¹
1.0	4.45
2.0	9.87
3.0	15.70
4.0	21.76
5.0	28.14



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