Preparation of Carboxylic Acids from Aldehydes (Including Hydroxylated Benzaidehydes) by Oxidation with Chlorite

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During the oxidation of aldehydes to acids with sodium chlorite, chlorine dioxide is simultaneously formed and may react with starting material and products in undesirable side-reactions. By performing the reaction in the presence of a chlorine scavenger, e.g. sulphamic acid, the formation of chlorine dioxide is avoided, and high yields of the acids from the corresponding aldehydes (even hydroxylated benzaldehydes) are obtained.

The oxidation of vanillin (3-methoxy-4-hydroxybenzaldehyde) with chlorite and with chlorine dioxide has previously been studied by Purves and co-workers,¹ Sarkanen et al.,² and Ishikawa et al.³ They obtained complicated oxidation mixtures from which a few compounds could be isolated in low yields. Neither chlorine dioxide nor chlorite oxidation yielded any vanillic acid.

We reported recently that the chlorite oxidation of vanillin⁴ (as well as of some other phenolic lignin model compounds and some lignin materials⁵) is retarded when a chlorine scavenger is added. Further, the reaction is accelerated by chlorine addition. From these observations we concluded that the oxidation proceeds by a chain mechanism with the following propagation reactions:

\[ \text{Cl}_2 + 2 \text{ClO}_2^- \rightarrow 2 \text{ClO}_2 + 2 \text{Cl}^- \]
\[ \text{vanillin} + 2 \text{ClO}_2 \rightarrow \text{products} + \text{ClO}_2^- + \frac{1}{2} \text{Cl}_2 + \text{H}^+ \]

(The reaction can also be formulated with hypochlorous acid instead of chlorine.)

The way in which the chain process is initiated was not elucidated, but was suggested to be caused by the chlorine formed by a slow reaction between chlorite and vanillin. This proposal is confirmed in this paper.

RESULTS AND DISCUSSIONS

According to the proposed mechanism, the chlorite oxidation of vanillin in the presence of a chlorine scavenger should give the reaction products of
the initial reaction. When such an oxidation was performed with sulphamic acid or resorcinol as scavenger under proper conditions (room temperature, pH 3.5), vanilllic acid was obtained in good yield (81 – 84 %). In the experiment with resorcinol addition, 4-chloro-1,3-dihydroxybenzene was formed in almost theoretical amounts. When phenol was used as scavenger, the vanilllic acid formed was difficult to isolate from the reaction mixture.

These results indicate that the initiating reaction between chlorite and vanilllic gives hypochlorite and vanilllic acid. The oxidation probably starts with an attack by a chlorite ion on the protonized aldehyde group:

\[
R-\text{CHO} \xrightarrow{H^+} R-C^+\text{H(OH)} \xrightarrow{\text{ClO}_2^-} R-\text{CH(OH)}(\text{OClO}) \xrightarrow{\text{HOCl}} R-\text{COOH}
\]

This assumption is supported by the relative reaction rates between different aldehydes (see below) and by the fact that the reaction is catalysed by acids.

Chlorite seems to be a rather selective agent for oxidation of aldehydes to carboxylic acids and it has been used repeatedly for this purpose in the carbohydrate field.\(^6,7\) Its use is hampered, however, by the formation of chlorine dioxide. This gas is unpleasant and may cause side reactions which by the above mentioned chain mechanism may predominate as in the oxidation of vanilllin. These complications may be avoided by adding a chlorine scavenger.

The synthesis of vanilllic acid from vanilllin has earlier been carried out by oxidation with silver oxide and by dehydrogenation in alkali hydroxide solution, but no convenient method has been published for the oxidation of hydroxylated benzaldehydes (such as vanilllin and o-vanilllin) to their corresponding acids. Chlorite oxidation in the presence of a scavenger now seems to be a suitable and inexpensive method.

The disagreeable formation of chlorine dioxide was avoided also when the chlorite oxidations of benzaldehyde, cinnamic aldehyde, p-nitrobenzaldehyde and cyclohex-3-ene aldehyde were performed in the presence of a scavenger. The corresponding acids were obtained in good yields.

The reaction rate (determined by the chlorite consumption) increased in the order vanilllin, benzaldehyde, and p-nitrobenzaldehyde, as should be expected for a reaction involving a nucleophilic attack by chlorite, as postulated above.

**EXPERIMENTAL**

The sodium chlorite used was technical grade (about 80% pure). The chlorite concentrations were determined iodometrically. Besides the m.p. determination, all products were identified by their IR and/or NMR spectra.

2-Hydroxy-3-methoxybenzoic acid. To a solution of o-vanilllin (10 mmol) and sulphamic acid (13 mmol) in water (500 ml), sodium chlorite (13.4 mmol) in water (20 ml) was added. After 30 min the solution was extracted with ether. Evaporation of the ether and recrystallisation of the residue from water gave 2-hydroxy-3-methoxy-benzoic acid monohydrate (1.36 g, yield 80%; m.p. 151 – 152°C, lit. 151°C).

Vanilllic acid. (a) Sulphamic acid as scavenger. To a solution of vanilllin (98.6 mmol) and sulphamic acid (134 mmol) in water (2 l) a solution (150 ml) of sodium chlorite (102 mmol) was added. The precipitate of vanilllic acid (13.9 g, 84 %; m.p. 208 – 211°C, lit. 211°C) was filtered off after 1 h.

Resorcinol (b) as scavenger. Vanilllin (10 mmol) in t-butanol (19 ml) and resorcinol (13 mmol) in an acetate buffer (pH 3.93) and aqueous sodium chlorite solution (12 mmol, 13 ml) were mixed. The precipitate of vanilllic acid (yield 50 %) was filtered off. From the

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filtrate, a further amount (31 %, total yield 81 %) of vanillic acid was obtained: the filtrate was extracted with methylene chloride and from the extract the fraction soluble in a sodium hydrogen carbonate solution was isolated. The extract fraction soluble in a sodium hydroxide solution contained 4-chloro-1,3-dihydroxybenzene; the yield estimated by NMR was 87 %.

_Cyclohex-3-en carboxylic acid._ To a mixture of cyclohex-3-en aldehyde (100 mmol) in a phosphate buffer (pH 3.5, 100 ml) sodium chlorite (120 mmol) in water (40 ml) was slowly added at −5°C. The temperature rose rapidly to 20°C. After 15 min the main part of the dioxane was removed by evaporation. The residue was then acidified and extracted with ether from which the acidic product (13.1 g) was collected as above.

An aliquot (5 g) of the residue was distilled under nitrogen. The main fraction (138 – 140°C, 3 mmHg) consisted of cyclohex-3-en carboxylic acid (4.0 g, yield 80 %) indistinguishable by IR and NMR from an authentic sample.

_Cinnamic acid._ To a mixture of cinnamic aldehyde (10 mmol) in t-butanol (25 ml) and resorcinol (13 mmol) in an acetate buffer (20 ml, pH 3.92), sodium chlorite (12.4 mmol) in water (18 ml) was added. After 45 min the mixture was evaporated in order to remove most of the t-butanol and was then extracted with ethyl ether. From the ether solution the acidic fraction (1.15 g) was collected as above. It consisted of cinnamic acid contaminated with 4-chloro-1,3-dihydroxybenzene. Sublimation of an aliquot (202 mg) of the mixture gave cinnamic acid (171 mg), m.p. 136 – 137°C, which corresponds to an over all yield of 72 %.

_P-Nitrobenzoic acid._ To a mixture of p-nitrobenzaldehyde (10 mmol) in t-butanol (60 ml) and resorcinol (13 mmol) in an acetate buffer (50 ml, pH 4) an aqueous solution of sodium chlorite (12 mmol, 13 ml) was added. After 30 min the mixture was extracted with ether. Evaporation of the ether gave p-nitrobenzoic acid, m.p. 241 – 243°C, lit. 242°C, (yield 76 %).

_Benzoic acid._ To a mixture of benzaldehyde (10 mmol) in t-butanol (10 ml) and resorcinol (13 mmol) in an acetate buffer (20 ml, pH 3.93) a solution of sodium chlorite (12 mmol) in water (13 ml) was added. After 18 h the reaction mixture was extracted with methylene chloride, from which the acid fraction (0.955 g, m.p. 85 – 110°C) was collected as above. Fractionated sublimation of an aliquot (100 mg) gave benzoic acid (84.3 mg, over-all yield 86 %), m.p. 124 – 125°C.

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