Kinetics of Oxidation of Tryptophan by Sodium Hypochlorite

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Introduction

Several kinetic investigations on amino acid oxidation have been performed [1–3]. Depending on the nature of the oxidant these authors report different products involving different reaction mechanisms. In this study sodium hypochlorite was used for the oxidation of the aromatic amino acid tryptophan. Under slight alkaline conditions 3-indoleacetaldehyde is the major oxidation product [4]. Under our experimental conditions this reaction was highly selective, so it was possible to investigate the reaction kinetics.

Materials and Methods

Materials: DL-tryptophan (side-chain-3-14C), specific activity 48 mCi/mmol (New England Nuclear); 3-indolyl-(2-14C)-acetic acid, specific activity 25 to 35 mCi/mmol (CEA); 3-indoleacetaldehyde × NaHSO3 (Sigma); indole-3-ethanol (Sigma); 3-indoleacetaldoxime, synthesized according to [5]; 5.2% NaOCl solution (Merck).

General procedure: 500 µl of an aqueous 10 mM try solution (adjustment to desired pH by either 1 N HCl or 1 N NaOH) including 200 nCi of 14C-labelled try are overlayered with 10 ml of redistilled benzene. The cylindrical reaction vessel (25 × 100 mm) is fixed in a water bath and the two layers are thoroughly mixed by a magnetic stirrer. The reaction is started by the addition of 4 µl of a 0.5% NaOCl solution of the same pH as the reaction mixture; the same amount of NaOCl solution is supplied at one minute intervals throughout the reaction time. Every 5 min a 0.5 ml sample of the benzene layer is taken for determination of its radioactivity and replaced by fresh benzene. At the end of the reaction the benzene layer is pipetted off.

Identification and quantification of the reaction product: For determination of purity the benzene phase is dried on Na2SO4 and its volume reduced to dryness by RFE. Aliquots equivalent to 50 nmol IAAld are submitted to TLC in 2 different systems:

1. Silicagel F254 on alufoil (Merck); solvent system chloroform: methanol (96:4).
2. Cellulose F254 on alufoil (Merck); solvent system water: methanol (89:1).

The product is identified by cochromatography with authentic IAAld and localized by both its fluorescence quench at 254 nm and its colour reaction with Ehmann’s reagent [6]. To determine the distribution of radioactivity non sprayed chromatograms are cut horizontally into 0.5 cm stripes and their radioactivity is determined by liquid scintillation counting.

Derivatization of the product: For the conversion to indole-3-ethanol an aliquot of the dried benzene phase is evaporated by RFE over 1 ml 1 M NaBH4 and the reaction is allowed to proceed for additional 10 min. The reaction mixture is extracted twice with 10 ml benzene and the combined benzene phases are dried on Na2SO4. After volume reduction by RFE aliquots equivalent to 50 nmol indole-3-ethanol are chromatographed in the TLC systems already described. Identification with authentic indole-3-ethanol and quantification follow the procedure described for IAAld.

The conversion to 3-indoleacetic acid is carried out by oxidation of IAAld with Ag2O. Aliquots of the dried benzene phase containing additionally varying amounts of unlabelled IAAld are evaporated to dryness by RFE. 2 ml 20 mM AgNO3 of different...
pH values (range 8.0 to 13.0) are added. The reaction time is varied between 0.5 and 90 min. The reaction is stopped by adjusting the reaction mixture to pH 2.5 with 1 M H₂SO₄. The reaction products are extracted twice with 10 ml of diethylether. TLC, identification and quantification follow the procedure described for IAAlld.

To form the bisulfite addition product the crude benzene phase is washed with 0.1 M potassium phosphate buffer, pH 7.0 and evaporated by RFE over 1 ml 50 mM NaHSO₃, pH 7.0. Non-bound radioactive compounds are reextracted with fresh benzene. Liberation of IAAlld from the bisulfite addition product is performed by adding 1 M sodium carbonate, pH 10.0 followed by extraction with fresh benzene.

Results and Discussion

1. Reaction order

As the oxidant was supplied continuously the reaction order was expected to be first order. The graphical determination of the reaction order confirmed this assumption (Fig. 1). For all experiment variants more than 90% of the reaction products cochromatographed with authentic IAAlld in two TCL systems demonstrate that the aldehyde is the only benzene soluble product.

![Fig. 1. Determination of reaction order. Initial tryptophan concentration: 10 mM; temperature: 45°C; pH 10.0. a, initial radioactivity of tryptophan (4.4 × 10⁸ dpm); x, actual radioactivity of indoleacetaldehyde.](image)

2. pH Dependence of the reaction

Varrying the pH in the range from 8.5 to 11.0 did not alter the selectivity of the reaction as concluded from TLC of the dried benzene phase. The initial reaction rate is directly proportional to the concentration of the unprotonated try suggesting that only the unprotonated amino acid is oxidized (Fig. 2).

![Fig. 2. Dependence of initial rate on unprotonated tryptophan concentration. Initial tryptophan concentration (total) 10 mM. Unprotonated tryptophan concentration at different pH values calculated from the pK value of the amino group (9.39).](image)

3. Temperature dependence of the reaction

The selectivity of the reaction was not impaired when varying the temperature between 298 K and 318 K. The activation energy is 35 ± 2.2 (SE) kJ·mol⁻¹ as calculated from the Arrhenius plot (Fig. 3).

![Fig. 3. Arrhenius plot. Initial tryptophan concentration: 10 mM; pH 10.0.](image)

4. Evidence against a radical mechanism

The oxidant NaOCl is in a pH dependent equili- brium with HOCI. The latter may readily dispro- portion to HCl and oxygen radicals. If oxygen or OH-radicals (which are formed subsequently in aqueous solution) were the oxydizing agents the addition of compounds known to scavenge these radicals should decrease the reaction rate [3]. However, adding mannitol in excess (1 M) did not influence the reaction rate significantly (control: rate 320 nmol·min⁻¹; + 1 M mannitol = rate 350 nmol·min⁻¹). This result and the high selec- tivity of the reaction suggest that OCl⁻ ions are the more likely oxidant.

5. Identification of the reaction products

The identity of the aldehyde was confirmed as follows:
a) More than 90% of the radioactivity of the benzene phase could be reversibly bound to NaHSO₃ indicating the formation of the bisulfite addition product,
b) in 2 TLC systems more than 90% of the radioactivity cochromatographed with authentic commercially available IAAld,
c) after derivatisation with NaBH₄ 95% of the radioactivity cochromatographed with authentic commercially available indole-3-ethanol,
d) the derivatisation to 3-indoleacetic acid was attempted by oxidation with Ag₂O yielding a variety of labelled compounds one of which cochromatographed with authentic 3-indoleacetic acid (varying the parameters reaction time and pH did not improve the conversion to 3-indoleacetic acid; however, the spectrum of oxidation products was the same when commercially available unlabelled IAAld or 3-indolyl-(2-¹⁴C)-acetic acid were submitted to the same procedure).

25% of the major impurity (accounting for up to 6% of the radioactivity in the benzene phase) could be converted to IAAld simply by stirring the compound in benzene over sodium carbonate buffer, pH 10.0 at 45 °C. The compound cochromatographed with authentic indoleacetaldoxime which is readily hydrolyzed to the aldehyde under these conditions. Hence 3-indoleacetaldoxime is a postulated intermediate in the proposed reaction mechanism.

6. Proposed reaction mechanism
The initial step could be a nucleophil substitution between the unprotonated amino group and 2 OCI⁻ ions similar to the initial step proposed by Gowda and Mahadevappa [2] who used Chloramine-T as the oxidant. After dehydration and decarboxylation the aldoxime would be formed which under our experimental conditions hydrolyzes to the aldehyde.