ELECTROCHEMICAL AND ELECTROPHORETIC STUDY OF SODIUM METAMIZOLE

LUIS A. BASÁEZ1, IVÁN M. PERIC1, PAOLA A. JARA1, CÉSAR A. SOTO1, DAVID R. CONTRERAS2, CAROLINA AGUIRRE3 and PETR VANYSEK3

1Department of Analytical and Inorganic Chemistry, Faculty of Chemical Sciences, University of Concepción, Concepción, Chile
2Faculty of Chemical Sciences, University of Santísima Concepción, Concepción, Chile
3Department of Chemistry and Biochemistry, Northern Illinois University, DeKalb, IL 60115, USA

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ABSTRACT

Electrochemical and electrophoretic behavior of sodium metamizole under conditions of variable pH has been studied. Voltammetric results indicate that sodium metamizole behaves as a quasi-reversible system over the potential window considered. These studies were complemented with capillary zone electrophoresis analysis in order to know the starting composition of solutions of sodium metamizole at different pH values and buffer composition. Among degradation products, the extent of the active metabolite 4-methylaminoantipyrine was found to be media dependent as well as was the appearance of different degradation products. Consequently, the extent of the oxidation process in voltammetric studies is dependent on the composition of the starting solutions. Information obtained by cyclic voltammetry and capillary zone electrophoresis has been concordant and conclusive in that both methods can be used in a qualitative study of sodium metamizole, under the experimental conditions assayed.

Keywords: sodium metamizole, NSAID, drugs, voltammetry, electrochemical determination, capillary zone electrophoresis.

INTRODUCTION

Pharmaceutical industry requires fast and reliable analytical methods in order to study intermediates as well as final products. The industry is always seeing new analytical techniques to obtain faster and more economical results and therefore there is always active and justified active research in this field. Dipyrone or sodium metamizole, [CA 68-89-3], sodium [(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)methylamino] methanesulphonate (Fig. 1) is a well known water soluble white crystalline powder with an effective analgesic and antipyretic activity. Because of its possibly severe side effect, agranulocytosis, it is not available in the USA. However, it is very popular preparation in Europe and Latin America. Sodium metamizole is usually determined by spectrophotometry1-3, titrimetry4-6, amperometry4, fluorometry4-8, and high performance liquid chromatography (HPLC)9-11. Recently, chemiluminescence coupled with flow-injection analysis (FIA)12,13, spectrophotometry coupled with FIA14-17, voltammetric18 and reflectometric19 determinations have been reported.

Figure 1. Chemical structure of Sodium Metamizole

Dipyrone is a non-opiate and non-steroidal anti-inflammatory analgesic drug (NSAID) which belongs to the group of pyrazolones20. It is actually a prodrug, which is effective only when it becomes hydrolyzed, after oral or intravenous intake, into its active metabolite21. In oral administration sodium metamizole spontaneously hydrolyzes in the gastric fluid by a non enzymatic mechanism to 4-methylaminoantipyrine, which is quickly and almost completely absorbed in the gastrointestinal tract. The 4-methylaminoantipyrine is metabolized in the liver by demethylation to 4-aminoantipyrine and by oxidation to 4-formylaminoantipyrine (inactive metabolite). 4-aminoantipyrine also converts by a polyformic acetylation to 4-acetylaminoantipyrin (inactive metabolite). It is necessary to point out that other non identified metabolites exist22.

Dipyrone has been studied in vivo and in vitro by such techniques as HPLC and HPTC either as monodrug or in mixtures. Determination of 4-aminoantipyrine, dipyrone and its metabolites in urine and 4-methylaminoantipyrine, 4-formylaminoantipyrine in plasma by micellar chromatography has been carried out23. HPLC has shown to be a technique of choice to be use in stability assays of pharmaceutical formulations of dipyrone (tablets and ampules) due to a good selectivity in reverse phase, detection limits and procedure speed24. Some of the degradation products have been detected and quantified by TLC25. Favorable repeatability and detection limits make this technique favorable for dipyrone stability assays in tablets and ampoules25. High performance thin layer chromatography (HPTLC) has been used to quantify sodium metamizole in suppositories at it has been shown that this method did not produce results significantly different from the official spectrophotometric method prescribed for its quantification by the German Pharmacopoeia26.

Dipyrone is often determined in pharmaceutical formulations with caffeine, paracetamol, or codeine using successfully spectroscopy27. Preparation with ascorbic acid was successfully accomplished electrochemically by polarographic determination28.

Dipyrone itself is not particularly stable and therefore routine assays must be performed quite regularly. Dipyrone has good stability when stored under completely anhydrous conditions and at low atmospheric pressure. In liquid formulations, an increased stability could be obtained by blanketing with an inert gas and by addition of a suitable antioxidant29.

We and others have reported earlier electrochemical techniques in which drug assay is performed by means of the transport across the interface between two immiscible electrolytes (ITIES). For example betalactamic antibiotics like ampicillin and cephalixin30 and amoxicillin31 were studied by this method. ITIES studies of the degradation products for acid and alkaline hydrolysis of penicillin V, amoxicillin in pure drugs as well as in pharmaceutical formulations, showed that the obtained results are comparable to the standard HPLC methods (XXII USP) and statistical evaluation showed no significant differences between the two methods. Voltammetric methods can be used for simple and fast determinations for some drugs such as gestodene, insulin and dopamine in presence of ascorbic acid32.

In this study we will use a variant of the cyclic voltammetry where the applied potential generates a potentiodynamic perturbation between two preset limits inside the potential window of the solvent33. Cyclic voltammetry, a type of potentiodynamic electrochemical measurement where a potential is applied to the system34-35, and the faradaic current response is measured is not alone very suitable analytical method for drug determination because it lacks the needed specificity.

On the other hand, capillary electrophoresis (CE) separation has been used frequently in the development and validation of a method for separation and
determination of components in different pharmaceutical formulations. If all the components of interest are in ionic form, then capillary zone electrophoresis (CZE) can be chosen as the better electrophoretic mode with by Capillary Electrophoresis Analysis Combined with Infra-red Spectroscopy. The high resolution and efficiency that can be achieved allow to apply this method not only for the determination of the active substance but also for stability control of pharmaceutical formulations. To the best of our knowledge, there are no communications on the use of capillary zone electrophoresis for the analysis of sodium metamizole in pure drug as well as pharmaceutical formulations. The main attractiveness of CE is that it uses small quantities of sample and reagents, it is fast, and also very versatile in separating analytes of small and large molecular mass, both neutral and charged.

**EXPERIMENTAL**

**Apparatus**

Cyclic voltammetric measurements were carried out with the Voltammetry Analyzer BAS CV-50W in a classic cell equipped with three electrodes (platinum working, platinum auxiliary (area 5 cm², approximately 10 times larger than the working electrode) and an Ag/AgCl reference electrodes. The capillary electrophoresis separations were performed on a PrinCE 450 capillary electrophoresis system (Prince Technologies, Essen, Netherlands) equipped with a Lambda 1010 UV-VIS detector (Bishoff, Leonberg, Germany). Fused silica capillaries (Polymer Technology, Phoenix, AZ, USA) of 50 µm i.d. and 45/55 cm were used. The system DAX v. 5.0 software was used for data acquisition.

**Reagents and solutions**

All the solutions were prepared with ASTM type 1 reagent grade water obtained by passing deionized water through a Pura-Filter™ 4-channel water purification system (Barnstead Thermenly Corp., Dubuque, USA). All chemicals of analytical grade and used without further purification. The supporting electrolytes used for experiments were a 0.10 mol L⁻¹ KNO₃ or HNO₃. Different buffers solutions were prepared in the range of pH from 1 up to 11. Na₂HPO₄/NaH₂PO₄ and NaH₂PO₄/H₃PO₄ systems for pH 1, 2, 3, 6 and 7. NaAc/HAc system for pH 4 and 5; and Na₂CO₃/NaHCO₃ system for pH 8, 9, 10 and 11. A 0.10 mol L⁻¹ NaOH or HCl were used to adjust the pH of the buffer solutions. A 0.010 mol L⁻¹ sodium metamizole (Industrial Kern, Spain; a gift from Laboratorio Pasteur, Chile) was prepared by dissolution of an appropriate amount of this reagent in a supporting electrolyte or buffer solution and used for voltammetric measurements. Capillary electrophoresis determinations were carried out with 0.050 mol L⁻¹ sodium metamizole samples obtained by further dilution of the 0.010 mol L⁻¹ solution, prepared as before, with the required supporting electrolyte or buffer solution. The running buffer consisted of 0.050 mol L⁻¹ boric acid adjusted to pH 9.2 with 0.01 mol L⁻¹ NaOH. All solutions were filtered through a 0.45 µm syringe filter and degassed by sonication for 5 minutes before injection to the CE system.

**Procedure**

Before the voltammetric measurements, the 0.010 mol L⁻¹ sodium metamizole solution was purged with nitrogen for 20 minutes to remove dissolved oxygen. The cell was thermostated at 20°C. The working electrode was polished with alumina (BAS CF-1050) on an alumina polishing pad (BAS MF-1040) for 2 min and then rinsed with Type 1 reagent grade water before each measurement. The voltammetric experiments were performed in stirred solutions.

In the CE experiments the capillary was routinely rinsed with 0.10 mol L⁻¹ NaOH (15-20 min), water (10 min) and running buffer (10 min) at the beginning of each working day. The samples were injected hydrodynamically at pressure 5000 Pa for 6 s. The running voltage was 18 kV and the temperature was set at 25°C. UV detector was used with detection wavelength 200 nm. Between individual runs the capillary was rinsed with 0.10 mol L⁻¹ NaOH (10 min), water (5 min) and running buffer (5 min).

**RESULTS AND DISCUSSION**

The effect of the scan rate variation from 1 to 100 mV s⁻¹ on the cyclic voltammery response behavior was investigated. 10 mmol L⁻¹ sodium metamizole analtye in 100 mmol L⁻¹ KNO₃ or HNO₃ polarized in the range of from +0.30 to -0.75 V on right platinum electrode was used. Sodium metamizole shows a distinct anodic peak. The current of this peak varied linearly with the scan rate when KNO₃ was used as the supporting electrolyte. Based on these results (not shown), a scan rate of 100 mV s⁻¹ was chosen for further studies as it provided voltammograms with acceptable peak resolution.

The effect of pH on the voltammetry of the 0.010 mol L⁻¹ sodium metamizole solution at a platinum working electrode was also studied. Cyclic voltammograms at pH 2 and pH 3 (Figs. 2 and 3) indicates a clear evidence of a redox reaction, with an incipient sign of oxidation in the Na₂HPO₄/NaH₂PO₄ buffer. These results are in agreement with the corresponding electropherograms (Figs. 6 and 7) which show that hydrolysis at pH 2 was greater than at pH 3 when the samples were prepared. The greater the extent of hydrolysis, the smaller is the oxidation peak in the corresponding voltammogram.

The behavior of the sodium metamizole in acidic media does not have notable interference on the working electrode, possibly because the electrode is not sensitive enough to capture differences in the structure of the drug and degradation products. However, dynamics apparently plays an important role in the behavior of this analyte as is corroborated by the presence of a quasi-reversible behavior clearly manifested by the Eₚ - Eₛₚ differences close to 0.1 V vs. Ag/AgCl.

Systems at higher pH values become more irreversible (Figs. 4 and 5), which could be due to the appearance of different degradation products shown at the corresponding electropherograms (Fig. 8 and 9) in Na₂CO₃/NaHCO₃ buffer. The appearance of the active metabolite 4-methylaminoantipyrine is clearly shown by the presence of a peak at the left side of the electropherogram. Sodium metamizole itself is the peak at the right side of the electropherograms. Consequently, the peaks in between can be attributed to unidentified degradation products.

The effect of the media matrix is also remarkable, since at pH 8 in the Na₂CO₃/NaHCO₃ buffer the amount of 4-methylaminoantipyrine is small and the extent of the degradation product eluted close to sodium metamizole is important (Fig. 8), but the amount of 4-methylaminoantipyrine becomes significant at pH 11 as well as does the appearance of a new degradation product observed between sodium metamizole and 4-methylaminoantipyrine, a peak which is almost imperceptible at pH 8 (Fig. 9). On the contrary, Fig. 6 (pH 2) shows a important degradation product close to sodium metamizole, which is almost negligible in Fig. 7 (pH 3), all this is when the same Na₂HPO₄/NaH₂PO₄ buffer was used, at pH 2 and pH 3 respectively.

![Figure 2. Voltammogram of sodium metamizole in a buffered solution at pH 2](image-url)
Figure 3. Voltammogram of sodium metamizole in a buffered solution at pH 3

Figure 4. Voltammogram of sodium metamizole in a buffered solution at pH 8

Figure 5. Voltammogram of sodium metamizole in a buffered solution at pH 11

Figure 6. Electropherogram of sodium metamizole in a buffered solution at pH 2

Figure 7. Electropherogram of sodium metamizole in a buffered solution at pH 3

Figure 8. Electropherogram of sodium metamizole in a buffered solution at pH 8
CONCLUSIONS

Cyclic voltammetry and capillary zone electrophoresis have been assessed for the development of analytical methodologies for sodium metamizole as possible alternative to the quality control needs of the pharmaceutical industry. The information obtained by either the cyclic voltammetry or the capillary zone electrophoresis is concordant and lead to the conclusion that both methods can be used in the qualitative study of sodium metamizole. Voltammetric results indicate that sodium metamizole behaves as a quasi-reversible redox system. Voltammetric results of the Universidad de Concepción, DIUC (Grants N°203.021.017-1.0, N°207.021.025-1.0 and 208.021.025-1.0).

The extent of this oxidation process is determined by the composition of the starting samples, the different pH values as well as the buffer composition, all of which was established from the results of the capillary zone electrophoresis experiments. The corresponding electropherograms show peaks attributable to sodium metamizole, 4-methylaminoantipyrine and some additional degradation products, which are generated depending on pH and buffer composition. However, the chemical nature of the degradation products was not identified in this study.

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REFERENCES
