Stability-indicating capillary zone electrophoresis assay for the analysis of linezolid in tablets

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A simple and fast alternative methodology using capillary zone electrophoresis (CZE) to analyze linezolid and its cationic photodegradation products in tablets has been developed. The separation was carried out on fused silica capillary and conducted using 100 mM formic acid (pH 3.0) and by applying 30 KV voltage. Detection was performed at UV 254 nm. The optimized method was validated in terms of linearity, limits of detection and quantification, precision (repeatability), stability studies (selectivity) and accuracy. Good linearity (8-20 mg L⁻¹) was obtained and the limit of detection was 0.95 mg L⁻¹. The greatest advantages of the CZE method were the rapid set-up of instrumentation and capillary equilibration, short analysis time (12 min), low running cost and low waste generation. The method showed good stability in determining linezolid submitted to degradation by light and to a climatic chamber and can be used as an alternative for evaluation in stability studies of linezolid in tablets, as well as for the analysis of the drug in raw materials and finished products.

Keywords: Capillary-electrophoresis/assay. Linezolid/tablets/stability. Pharmaceutical-drugs.

INTRODUCTION

Linezolid, N-(((S)-3-(3-Fluoro-4-morpholino-phenyl)-2-oxo-5-oxazolidinyl)methyl)acetamide (Figure 1), is a member of a new class of antibiotics, the oxazolidinones (Marchese, Schito, 2001), and it was the first oxazolidinone to be developed and approved for clinical use (Di Paolo et al., 2010). Linezolid (LNZ) is currently the only antimicrobial agent which can be administered orally, being rapidly absorbed following oral administration and eliminated via renal and non-renal routes (Boak et al., 2014). It is particularly important in cases where intravenous access is difficult and the patients require outpatient therapy (Itani et al., 2005; Kalil, Puumala, Stoner, 2006). LNZ has a wide spectrum of activity against gram-positive organisms and appears to be an effective option of treatment when compared to vancomycin(Patel, Gallagher, 2015; Silva, Monteiro Neto, Sejas, 2007).

Analysis of LNZ and its degradation products are usually assayed by high-performance liquid chromatography (HPLC) and spectrometry (Erlich et al., 2001; Krishna-Reddy et al., 2002; Patel et al.,...
2007; Merli et al., 2011; La Marca et al., 2012; Cios, Kus, Szymura-Oleksia, 2013; Chavda et al., 2016); however, this technique has important limitations such as the high cost of the analysis and the excessive generation of residues due to the use of organic solvents. Capillary electrophoresis (CE) has also been described as an alternative to HPLC for LNZ assays. The range of pharmaceutical applications of CE is similar to HPLC and its advantages include appreciably short analysis times, low residue generation, use of aqueous solvents, high resolution and specificity (Landers, 1997). CE presents several modes of operation, with zone capillary electrophoresis (CZE) and micellar electrokinetic capillary chromatography (MEKC) being the most commonly used.

MEKC has been used to determine LNZ in human serum samples (Kitahashi, Furuta, 2002). The authors used 25 mM borate buffer (pH 10.0) containing 80 mM sodium dodecyl sulphate (SDS) as running buffer. They compared CZE and MEKC and concluded that MEKC enabled a separation (with a migration time of 14 min) without overlapping with the peaks of impurities in serum and improvement in the theoretical plate number. An improvement of the method was published by the same authors in 2004 (Kitahashi, Furuta, 2004) lowering the pH of borate buffer (pH 8.0) and the concentration of SDS (50 mM). There was the addition of 7 M urea as an additive to increase the LNZ dissolution into the running buffer. The advantages included direct serum injection without pretreatment and lower migration time (5.5 min). CE was also used for the separation of enantiomers from LNZ (Michalska, Pajchel, Tyski, 2008a; Bednarek, Bocian, Michalska, 2008). Michaska, Pajchel, and Tyski (2008a) used MEKC to perform the LNZ enantiomeric separation. Charged β-cyclodextrins (β-CD) was added to the running buffer to carry out the separation with a migration time of 15 min.

Due to its chemical structure, LNZ is highly unstable under alkaline conditions and several degradation studies have been done to evaluate the compounds obtained under conditions of alkaline stress. Alkaline degradation and photodegradation products have been investigated by chromatography and spectrophotometry in LNZ commercial (tablets) and injectable samples (Bebawy, 2003; Lopes, Salgado, 2009; Lopes, Salgado, 2010; Kawy, Weshahy, Shokry, 2012; Hegazy et al., 2014). Chiral impurities from LNZ have been analyzed by MEKC in linezolid solution for infusion (Michalska, Pajchel, Tyski, 2008b).

CZE has never been used in LNZ tablet assay until now. The few scientific papers addressing CE for linezolid applied MEKC to LNZ samples in the injectable form. The aim of this work was to develop a stability-indicating CZE/UV assay method to analyze linezolid in the cationic form (under pH 4.0), which can be applied to studies of stability and determination of content of linezolid in the presence of degradation products. This is the first paper that uses CZE to analyze linezolid in its pharmaceutical form of tablets.

MATERIAL AND METHOD

Chemicals

All chemicals were of analytical grade and used as received. LNZ reference substance was obtained from “Synfine research” Chemical Company (Canada) and HPLC analysis showed that it contained 99.3 % of LNZ. Linezolid tablets (Zyvox®) were commercially obtained and were claimed to contain 600 mg of active drug (Pfizer, Brazil). Imidazole (purity 99.0%) from Merck was used as internal standard. Formic acid 85% was purchased from Synth. Ultrapure water was deionized by using a Milli-Q gradient system A10 (Millipore, Bedford, MA, USA) and was used to prepare all buffers.

Equipment

CE analyses were performed on a model PA800 Plus, Beckman Coulter, equipped with a UV/Vis detector. Bare fused silica capillaries with 75 μm i.d. were purchased from Polymicro Technologies, Phoenix, AZ, U.S.A. The capillary detection length was 50 cm and the total length 60 cm. The samples were injected by pressure (10s at 0.5 psi) and the detection was performed at 256 nm. Stock solutions of formic acid (1 M) were prepared in water and stored under refrigeration. Working background electrolyte consisted of 100 mM formic acid (pH 3.0) and the applied voltage was 30 kV.

Pre-treatment of capillary

The capillary was conditioned with 1 M NaOH for 30 min, followed by the passage of ultra pure water (Milli-Q) for 10 min and a running buffer for 20 min. At the beginning of each working day, the capillary was conditioned with a flush of NaOH (5 min), ultrapure water (5 min) and electrolyte (10 min). Between each running, the capillary was washed with 1 M NaOH followed by ultrapure water for 2 min and with background electrolyte for 5 min.
Preparation of the solutions

Preparation of standard solution and internal standard

Stock solutions of LNZ (reference substance) and imidazole (internal standard) were prepared in deionized water. Linezolid stock solution was prepared by accurately weighing 10 mg of linezolid in a 100 mL volumetric flask, and then dissolving this quantity in deionized water. The solution was sonicated for 10 min and brought to volume with water. For the calibration curve, working standard solutions were prepared by dilution of the linezolid stock solution to obtain six different concentrations within the range of interest, in this case, 8–20 mg mL⁻¹. All the solutions were prepared in triplicate.

Preparation of sample solution

An amount of powder equivalent to 500 mg LNZ was transferred to a 1000 mL volumetric flask with deionized water and shaken for 20 min, followed by the dilution to volume with deionized water (500 mg L⁻¹). A 10 mL aliquot of this solution was transferred to a 100 mL volumetric flask (50 mg L⁻¹). Lastly, an aliquot of this solution was dissolved in the deionized water to obtain a solution with a final concentration of 12 mg L⁻¹. This solution was used in the stability studies and in the assay of commercial tablets of linezolid.

Method validation

The method validation assays were carried out according to the International Conference on Harmonization Guidelines (ICH, 2005). The analyzed parameters were: linearity, limits of detection and quantification, precision (repeatability), selectivity, accuracy.

Linearity

For the assay of linearity stock solutions of LNZ reference substance were prepared at six concentration levels from 8.0-20 mg L⁻¹ and injected in triplicate. The linearity was evaluated by the least square regression method.

Limits of detection and quantification

The limits of detection (LOD) and quantification (LOQ) were calculated according to the equations:

\[ \text{LOD} = 3 \frac{S_y}{b} \]

\[ \text{LOQ} = 10 \frac{S_y}{b} \]

Precision

The precision of the method was evaluated by the repeatability (intra-assay). Repeatability of the method was studied by analyzing samples of tablets, at the same concentration, within 1 day and under the same experimental conditions.

Stability studies (Selectivity)

Selectivity and forced degradation studies were conducted according to the ICH Guidelines by introducing a sample of linezolid sample solution to different forced degradation conditions using thermal and photolytic analysis. For the selectivity test of the proposed method, tablet powder of LNZ was prepared and exposed to light (254 nm) and climate chamber (40 °C/75% for 90 days) to determine the effects of irradiation and humidity/temperature on the stability of LNZ in its solid state and was subsequently analyzed.

Accuracy

The accuracy of the method was evaluated by comparing the results obtained in the simultaneous determination of linezolid tablets by HPLC and capillary electrophoresis. Upon test “t-paired” it was found that the average values obtained by the two techniques did not differ significantly at a confidence level of 95%.

RESULTS AND DISCUSSION

Optimization CE/UV separation

In capillary electrophoresis, the choice of buffer or background electrolyte has direct implications for the optimization of separation since the pH of the electrolyte determines the effective mobility by decisively affecting the degree of dissociation (Landers, 1997). For basic compounds, it is recommended that the pH of the background electrolyte is around 2.5. This would provide better selectivity since acidic or neutral interferents exhibit negative mobility or simply do not have mobility, respectively, thus not being detected. Although LNZ does not have basic character, it has a nitrogen atom, which can be ionized and migrated as cation in CZE. Michalska, Pajchel and Tyski (2008a) studied the influence of pH on the behavior of LNZ in CE and found that it is protonated under pH 4.0 (LNZ has pKa around 1.8).

In this study formic acid under pH 3.0 was used as background electrolyte. Formic acid showed a good baseline at the work concentration (100 mM). Figure 2 shows the electropherogram of LNZ standard and the blank sample. Imidazole was used as internal standard and methanol as a marker of electroosmotic flow (EOF). It can be observed that the LNZ peak migrates before the EOF, which ensures that it is still protonated at this pH; however,
as the EOF is smaller and the electrophoretic mobility is also low, it takes 12 minutes to migrate.

Method validation

The method was validated in respect to parameters including linearity, limits of detection (LOD) and quantitation (LOQ), precision, stability studies (selectivity) and accuracy.

Linearity, limits of detection and quantitation

Linearity studies were performed by linear regression standard curve. Lower values of parameters like standard error (S.E.) of slope and intercept (Table I) indicated high precision of the proposed methods. Also, the mean slope and intercept values are within the 95% confidence interval. Goodness of fit of the regression equations was supported by high regression coefficient values and lower calculated $F$-values (Table I). With data from the evaluation of linearity, we calculated the limits of detection (LOD), quantification (LOQ) and the determination coefficient ($r^2$), which are presented in Table I. This analysis was done at 95% confidence and 5% significance level.

Precision

The data obtained from precision experiments are given in Table II for intra-day precision studies. Each concentration was injected six times ($n = 6$). The % RSD values for intra-day precision study was < 4.0% confirming that the method was sufficiently precise (Table II).

![FIGURE 2 - CZE of (A) – Blank sample, (B) - linezolid (chemical reference). CE/UV conditions: 100 mM formic acid, pH 3.0, applied voltage +30 kV, injection by pressure (0.5 psi for 10s) and detection at 254 nm. Caption: methanol as eof marker and imidazole as IS (internal standard).](image)

### TABLE I - Statistical analysis of calibration graphs and analytical data in the determination of linezolid by electrophoresis method ($n=6$)

| Parameters                  | Proposed electrophoresis method |
|-----------------------------|----------------------------------|
| Regression analysis         |                                  |
| Slope (S.E.$^a$)            | 275.30 (6.27)                    |
| Intercept (S.E.$^a$)        | -226.46 (87.22)                  |
| Correlation coefficient ($r^2$) | 0.9943                        |
| Equation                    | $y = 275.30x - 226.46$           |
| Analysis of variance        |                                  |
| Linear regression$^b$       | 1926.1430                       |
| Validation parameters       |                                  |
| Linearity (mg L$^{-1}$)     | 8-20                             |
| LOD (mg L$^{-1}$)           | 0.95                             |
| LOQ (mg L$^{-1}$)           | 3.16                             |

$^a$ Standard error of mean. $^b$ Corresponding critical values for $F$ at $p < 0.05$

### TABLE II - Comparison of CE and HPLC methods for determination of linezolid in Tablets (600 mg)

|                        | CE assay (content, %) | HPLC method (content, %) |
|------------------------|-----------------------|--------------------------|
| Results                | 100.91                | 103.5                    |
|                        | 98.64                 | 100.32                   |
|                        | 99.14                 | 100.81                   |
|                        | 106.84                | 105.00                   |
|                        | 105.71                | 101.31                   |
|                        | 107.56                | 101.31                   |
| Mean                   | 103.13                | 102.04                   |
| Number of samples      | 6                     | 6                        |
| R.S.D. (%)             | 3.90                  | 1.77                     |

$^*$Student’s $t$-value for $p = 0.05, n = 6, t_{tabulated} = 2.447$

Stability studies (Selectivity)

Figure 3 presents the electropherograms of LNZ tablet submitted to degradation by light. The sample of LNZ was injected in high concentration to visualize the degradation products. The appearance of two major peaks (1 and 2) can be observed, corresponding to cationic degradation products in just 10 days of exposure to light (LNZ solid state). These degradation products probably contain nitrogen in their structure, which is protonated at this pH and migrate as cationic form during electrophoresis. It is noted that the elution of
these degradation products are very close, assuming that the chemical structures are very similar. During 90 days of exposure in the UVC light the degradation products primarily formed, increasing over the course of time, indicating that these products can be characterized as primary products of degradation of linezolid submitted to light exposure. Figure 4 shows the electropherogram of linezolid tablet subjected to degradation in a climatic

**FIGURE 3** - Electropherogram of photodegradation products of LNZ tablet submitted to light UVC (254 nm).

**FIGURE 4** - Electropherogram of degradation products of LNZ tablet submitted to climate chamber (40 °C/75 %).
chamber (40°C/75%). The degradation in the solid state is slow and time consuming, but it is consistent with the adverse conditions of temperature and humidity in different countries during transport and distribution of this drug. In this analysis, we can watch the beginning of the formation of degradation products after 30 days. This electropherogram is consistent with the data obtained earlier by HPLC (Lopes, Salgado, 2009).

**Accuracy**

With the statistics of the two methods analyzed, it was found that there is no statistical difference between the methods, confirming the accuracy of the method by capillary electrophoresis (Table II).

**Analysis of pharmaceutical tablets**

The validated method was applied for the assay of commercial tablets containing 600 mg of linezolid. Results of the determination were found to be 103.13 ± 3.90% (RSD). The method was statistically compared with the HPLC method (Table II). The averages of the validated methods were compared by t-test and statistical differences were not observed between them.

**CONCLUSION**

In this work the linezolid tablets and its cationic degradation products were determined by CZE in low pH. This simple and validated methodology provided good linearity, precision and detection limits. CZE method was shown to be specific because there was no interference of the excipients present in the formulation of the linezolid tablets, as well as showing the ability to distinctly determine the LNZ from its degradation products. It could be used as an alternative to HPLC in the determination of the content and stability in studies of the LNZ tablets.

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**CONFLICT OF INTEREST STATEMENT**

The authors declare that there are no conflicts of interest regarding this manuscript.

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