Determination of fluconazole in serum and amniotic fluid of rats by gas-chromatography/mass spectometry (GC/MS)

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INTRODUCTION

Fluconazole, (2,4-difluoro-α,α'-bis(1H-1,2,4-triazol-1-ylmethyl)benzyl alcohol, is a broad-spectrum triazole antifungal agent, which has shown to be active against systemic and superficial infections (Richardson et al., 1985). Studies on the pharmacokinetics of fluconazole showed: rapid absorption after oral administration, no metabolization, approximately 80% is excreted unchanged in the urine, and a wide distribution in several organs in the body (Ripa et al., 1993; Debruyne and Ryckelynck, 1993). Recently, some reports pointed out the adverse effects of fluconazole on pregnancy (King et al., 1998). Cases of newborns with skeletal and cranium malformations, and heart problems suggested that high doses of this drug (400 – 800 mg/day) might be teratogenic (Mastroiacovo et al., 1996; Pursley et al., 1996; Polifka, Friedman, 1999). Despite the observed effects on the development of the foetus, there is no evidence until now about the transference of fluconazole through placenta. Although several analytical chromatographic methods have been developed to analyse fluconazole in biological fluids (Wood, Tarbit, 1986; Beijnen et al., 1991; Koks et al., 1995; Ng et al., 1996; Majcherczyk et al., 2002; Eerkes et al., 2003), there are no reports of fluconazole quantification in amniotic liquid. The objective of this study was to develop a simple assay for determination of fluconazole in serum and amniotic fluid of rats treated with oral dose of 100 mg/kg of the drug during pregnancy, which will be useful in further teratogenic studies.

EXPERIMENTAL

Chemicals

Fluconazole was obtained from Pfizer (São Paulo, Brazil) and the internal standard, tioconazole, from Galena...
Ethyl acetate (HPLC grade) was from Burdick & Jackson, USA and reagent grade water (Milli-Q®, Millipor Corp.) was used.

**Chromatography**

Analyses were performed on a GC-MS Shimadzu QP5050A instrument employing the following conditions: Column: CBP-5 (Shimadzu) fused silica capillary column (30 m long × 0.25 mm i.d. × 0.25 mm film thickness composed of 5% phenylmethylpolysiloxane) connected to an ion trap detector operating in EI mode at 70 eV; carrier gas: He (1 mL/min); injector and ion-source temperatures were 300 °C and a split ratio of 1:5. Injection volume was 1 mL (ethyl acetate) and the oven temperature was programmed from 160 °C (isothermal for 1 min), with an increase of 35 °C/min, to 270 °C, then isothermal for 10 min. Tioconazol, 10.0 µg, was used as internal standard. Concentrations of fluconazole in the maternal serum and amniotic fluid samples were derived by interpolation of the ratio between peak areas in the calibration curve.

**Sample collection**

Eight pregnant Wistar rats were treated by gavage with a 20 mg/ml suspension of fluconazole at a dose of 100 mg/kg, during organogenesis (from the 6th to the 15th gestation day). On the 15th day, two hours after the last dose of fluconazole, the rats were anaesthetized by ethyl ether inhalation and the peritoneal and thoracic cavities were opened. Maternal blood was collected from the heart and immediately centrifuged at 2500 g for 5 min. Then serum was removed and frozen until assay. Uteri were dissected longitudinally and the amniotic fluids from each maternal-fetal unit were collected and frozen until analysis. The approval for the use of animals and for the procedures required for the experiments was obtained by the Ethical Committee for Animal Use in Experimental Studies of the Universidade Federal de Goiás.

**Sample preparation**

To both 0.5 mL of maternal serum and amniotic fluid, 1.0 mL of water was added. The samples were then extracted with 6 mL of ethyl acetate by vortex-mixing for one minute. After 10 min centrifugation at 1400 g, 25 ºC, 5.0 mL of the organic layer were transferred to a clean tube and evaporated to dryness under N₂ (Kim et al., 1994). The internal standard, 20.0 µL tioconazol (5000.0 µg/mL in ethyl acetate), was added to the residue and the volume was made up to 200.0 µL with ethyl acetate. An aliquot of 1.0 µL of the mixture was injected in the column.

**RESULTS**

**Specificity**

The specificity of the assay was examined by comparing the chromatogram of the drug-free serum sample with that of standard solution of fluconazole and tioconazole. Retention times (RT) of fluconazole and tioconazole were approximately 5.37 and 11.7 min, respectively. Typical chromatograms are shown in Figure 1. The total run time for each sample injection was about 15 min. There were no interfering peaks in the blank serum at the retention times of fluconazole and the internal standard.

**Linearity**

Standard solutions were prepared by dissolving fluconazole in ethyl acetate, within a concentration range of 10.0 – 300.0 µg/mL (n=6). At each concentration level internal standard, tioconazol (10.0 µg), was added and the solutions were analysed in triplicate. Calibration curve was derived by plotting the peak-height ratios of fluconazole

![FIGURE 1 - GC chromatogram of fluconazole (concentration 50 mg/ml, RT 5,37 min) and tioconazole (concentration 10 mg/mL, RT 11,7 min).](image-url)
and internal standard in standard solutions against the known concentrations of fluconazole, using linear regression analysis. The equation obtained was $y = 0.19754x - 1.39985$, correlation coefficient $r = 0.9997$.

**Limit of quantification**

Limit of quantification defined as the lowest concentration with an R.S.D. = 20% was 0.1 µg/mL.

**Recovery**

For recovery studies, serum samples were spiked with fluconazole at three different concentrations (50, 100 and 200 µg/mL) and three samples from each concentration, were extracted using the assay procedure. After dryness, 20.0 µL of the internal standard, (5000.0 µg/mL in ethyl acetate), was added to the residue and the volume was made up to 200.0 µL with ethyl acetate, 1.0 µL aliquot of the mixture was analysed. The recovery was calculated from the peak areas of the spiked samples and the peak areas of standard solutions in the same concentration ranges. The efficiency of the extraction was greater than 90% in the three concentrations (Table I).

**Precision and accuracy**

Blank serum samples were spiked with fluconazole at three concentration levels (50, 100 and 200 µg/mL), at each concentration internal standard was added. Six replicates of each solution were analysed and the coefficients of variation (C.V.) were calculated using the mean of the concentrations found and S.D. For this study the variances were acceptable, with C.V. values below 5% (Table I). Accuracy (bias) for fluconazole was expressed as the percentage deviation of observed concentration from theoretical concentration; the greatest bias was equal to 3.8% (Table I).

**Maternal serum and amniotic fluid samples**

Typical GC chromatograms of serum and amniotic fluid samples of rats treated with fluconazole are shown in Figure 2.

**TABLE I** - Results obtained from method validation: precision (C.V.), accuracy (bias) and the recovery of fluconazole from the spiked serum samples

<table>
<thead>
<tr>
<th>Spiked serum samples (µg/mL)</th>
<th>Conc. found* (µg/mL) mean ± S.D.</th>
<th>C.V. (%)</th>
<th>Bias (%)</th>
<th>Mean recovery (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>50</td>
<td>49.4±1.8</td>
<td>3.6</td>
<td>-1.2</td>
<td>92.5±6.7</td>
</tr>
<tr>
<td>100</td>
<td>96.2±4.6</td>
<td>4.8</td>
<td>-3.8</td>
<td>93.2±4.9</td>
</tr>
<tr>
<td>200</td>
<td>195.3±7.5</td>
<td>3.8</td>
<td>-2.4</td>
<td>91.8±7.3</td>
</tr>
</tbody>
</table>

* Calculated from calibration curve for spiked serum samples ($n = 6$).

**FIGURE 2** - GC Chromatograms of fluconazole (5.37 min) and internal standard (tioconazole, 11.7 min). (A) serum sample and (B) amniotic liquid sample, obtained from rats treated with fluconazole.
Table II presents the concentrations of fluconazole determined in maternal serum and amniotic fluid after two hours of the last oral administration of the drug. The treatment lasted ten days (from 6th to 15th day of gestation) and the mean concentrations of fluconazole in maternal serum and in amniotic fluid were 206.01 ± 105.25 µg/mL and 125.34 ± 65.24 µg/mL, respectively.

**TABLE II - Fluconazole concentration detected in serum and amniotic liquid samples from rats treated at a dose of 100 mg/kg, during organogenesis**

<table>
<thead>
<tr>
<th>Treated Rats (n=8)</th>
<th>Fluconazole concentration in serum (mg/mL)</th>
<th>Fluconazole concentration in amniotic liquid (mg/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rat 1</td>
<td>273.02</td>
<td>170.34</td>
</tr>
<tr>
<td>Rat 2</td>
<td>321.41</td>
<td>100.57</td>
</tr>
<tr>
<td>Rat 3</td>
<td>126.70</td>
<td>91.55</td>
</tr>
<tr>
<td>Rat 4</td>
<td>170.94</td>
<td>157.44</td>
</tr>
<tr>
<td>Rat 5</td>
<td>48.55</td>
<td>25.04</td>
</tr>
<tr>
<td>Rat 6</td>
<td>107.42</td>
<td>60.70</td>
</tr>
<tr>
<td>Rat 7</td>
<td>296.18</td>
<td>194.56</td>
</tr>
<tr>
<td>Rat 8</td>
<td>303.86</td>
<td>202.52</td>
</tr>
<tr>
<td>mean±S.D.</td>
<td>206.01±105.25</td>
<td>125.34±65.24</td>
</tr>
</tbody>
</table>

**DISCUSSION**

Quantification of fluconazole in serum and amniotic fluid of Wistar rats was possible using the present method. The preparation of sample is easy followed by GC/MS analysis with a simple capillary column. Previous GC assays for fluconazole quantitative determination described a pre-treatment of the packed column with benzoyl chloride (Wood, Tarbit, 1986), or indicated the use of a semi-capillary column (Beijnen et al., 1991) with low efficiency. In both methods an electron capture detector was used, which is less selective than the ion trap detector. In the present method, column deactivation and derivatization of fluconazole were unnecessary; even so the limit of detection was 0.1 µg/mL. This fact shows the sensitivity of the proposed method. HPLC analysis of fluconazole using UV detector (Koks et al., 1995; Ng et al., 1996; Majcherczyk et al., 2002) showed to be less sensitive than GC/MS method. Recently a very sensitive HPLC/MS/MS method (Eerkes et al., 2003) has been published, however, unfortunately these instruments have not become commonly available in all laboratories yet.

**CONCLUSION**

A simple, specific and rapid GC/MS method for the measurement of fluconazole concentrations, in serum and amniotic fluid, is described. It offers a suitable alternative to the existing GC and HPLC techniques and it can be used to quantify fluconazole and tioconazole in biological fluids for teratogenic and pharmacokinetic studies.

**ACKNOWLEDGEMENTS**

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**RESUMO**

Determinação de fluconazol em soro e líquido amniótico de ratas por cromatografia a gas/espectrometria de massas (CG/EM)

Soro e líquido amniótico de ratas tratadas com fluconazol (dose oral de 100 mg/kg) durante a prenhez foram quantificados para este fármaco usando cromatografia gasosa acoplada à espectroscopia de massas (CG/EM). O fluconazol foi extraído das amostras com acetato de etila e analisado empregando-se um cromatógrafo CG/EM Shimadzu QP5050A com coluna capilar de sílica fundida CBP-5. O tioconazol foi utilizado como padrão interno. A curva padrão foi linear no intervalo das concentrações de 10,0 a 300,0 µg/mL. O limite de quantificação foi de 0,1 µg/mL e não foi observada interferência no branco de soro e líquido amniótico. As concentrações médias do fármaco no soro e líquido amniótico foram 206,01 ± 105,25 µg/mL e 125,34 ± 65,24 µg/mL, respectivamente. Este procedimento mostrou-se sensível e eficiente para ser usado em estudos de teratogenicidade do fluconazol e outros azóis.


**REFERENCES**


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