HIGH-PERFORMANCE LIQUID CHROMATOGRAPHIC ANALYSIS OF CEFOTETAN EPIMERS IN HUMAN PLASMA AND URINE

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SUMMARY

Cefotetan, a new broad-spectrum 7a-methoxycephalosporin antibiotic, was assayed in plasma and urine by means of reversed-phase high-performance liquid chromatography. Commercially available cefotetan exists in two epimeric forms. The procedure described allows the separation and quantitation of both epimers. For the first time a different pharmacokinetic behaviour ($t_{1/2} = 3$ h versus $4$ h) for each epimer after intravenous injection to healthy volunteers is demonstrated. It is assumed that one epimer is bound to a greater extent to serum proteins and is therefore responsible for the differences observed. As both epimers exhibit similar antibacterial activity, it seems doubtful whether these differences would have clinical significance. Iothalamic acid was determined simultaneously as a marker of kidney function.

INTRODUCTION

Cefotetan disodium is a new 7a-methoxycephalosporin with a wide spectrum of antibacterial activity and remarkable stability against various β-lactamases. Pharmacokinetic parameters in man have been studied in different races [1–4]. The commercially available product consists of a mixture of two epimers because of the asymmetric carbon atom at the dithietan ring. In this paper the $R$ and $S$ epimers are called A and B, respectively, because the absolute configuration was not known. In weakly alkaline media the two epimers are
in equilibrium with a third tautomeric form (Fig. 1). Under physiological conditions the proportion of the tautomer is small, as demonstrated by others [1, 4]. The aim of the present study was to elucidate the pharmacokinetic behaviour of both epimers of cefotetan after intravenous injection of the drug in healthy volunteers.

MATERIALS AND METHODS

Reagents
Cefotetan was provided by ICI-Pharma (Plankstadt, F.R.G.), Conray 70® (meglumine iothalamate) and iothalamic acid (3-(acyethylamino)-2,4,6-triiodo-5-[(methylamino)carbonyl]-benzoic acid) by Byk-Gulden (Konstanz, F.R.G.); acetonitrile and dichloromethane (ChromAR) were obtained from Promochem (Wesel, F.R.G.), the primary and secondary sodium phosphates (analytical grade) from Merck (Darmstadt, F.R.G.), and tetrabutylammonium bromide (purum) from Fluka (Neu-Ulm, F.R.G.). Water was purified with a Milli-Q water purification system (Millipore, Neu-Isenburg, F.R.G.).

The stock solutions of cefotetan and iothalamic acid were prepared in 50 mM sodium phosphate buffer (pH 6.2) to yield final concentrations of 1.0 mg/ml and 10.0 mg/ml, and stored in aliquots at −20°C. The stock solutions were then diluted with drug-free plasma to provide assay standards over a concentration range of 1.25–100 μg/ml for cefotetan and 12.5–1000 μg/ml for iothalamic acid. For urine samples the standard solutions were prepared in 50 mM phosphate buffer (pH 6.2).

Analysis of plasma and urine samples
A 200-μl volume of plasma was buffered with 200 μl of 0.1 M sodium dihydrogen phosphate and mixed for 5 sec with 400 μl of acetonitrile on a Reax 1 shaker (Heidolph, Kelheim, F.R.G.). After 15 min at 4°C the mixture was
centrifuged (1 min, 10,500 g) and the clear supernatant transferred to a 12 × 75 mm disposable polypropylene tube. Then 2.0 ml of dichloromethane were added, the tube was capped and mixed (5 min) on a Reax 2 Mixer (Heidolph). After centrifugation (10 min, 4800 g) an aliquot (5–50 μl) of the upper aqueous phase was injected onto the column.

Urine samples were centrifuged (10 min, 4800 g) and diluted 1:10 with 50 mM sodium dihydrogen phosphate (resulting pH 5.8–5.9).

All biological samples were stored at −70°C until required for analysis.

Chromatography

The chromatographic system consisted of a pump M 6000A, a fixed-wavelength detector M 440 allowing simultaneous measurement at 254 and 280 nm, an automatic injector WISP 710 B, a data module M 730 and a system controller M 720 (all from Waters Assoc., Königstein/Ts., F.R.G.). A Hibar® stainless-steel column (125 × 4 mm I.D.), prepacked with 5-μm LiChrosorb RP-18 silica (Merck), was used for separation. The flow-rate was maintained at 1.0 ml/min, the resulting back-pressure was 110–115 bar. The eluent was monitored at 254 and 280 nm, integrating the peak area of the signals at 280 nm. For plasma samples the mobile phase was prepared by combining 925 ml of water, 75 ml of acetonitrile, 5.50 g of sodium dihydrogen phosphate monohydrate, 1.80 g of disodium hydrogen phosphate dihydrate and 20 mg of tetrabutylammonium bromide. For urine samples the contents of acetonitrile and tetrabutylammonium bromide were changed to 45–50 ml/l and 22.5 mg/l, respectively. The apparent pH of the buffered solutions was about 6.4.

RESULTS

Sample clean-up

The major disadvantage of high-performance liquid chromatography (HPLC) in comparison with the microbiological assay is the need to remove proteins from the samples before injection onto the analytical column. In order to maintain the precision, a minimal number of clean-up steps (e.g. pipetting or extracting) is desirable. Deproteinization is usually performed by adding organic solvents or acids. Ultrafiltration seems to be a very fast and simple technique and was used for the determination of cefmenoxime in plasma; but addition of sodium dodecylsulphate (SDS), a highly protein-bound displacing agent, is necessary to recover the protein-bound fraction of the drug [5]. By adding 0.5% SDS we obtained nearly quantitative recoveries of both epimers of cefotetan from plasma using the sample clean-up procedure described under Materials and methods. However, the ion-pairing properties of SDS in reversed-phase HPLC caused chromatographic aberrations with peak splitting. Deproteinization by addition of perchloric acid resulted in low recoveries because of apparent co-precipitation of cefotetan. After sample clean-up by extraction of the cephalosporin and iothalamic acid into chloroform—pentanol as described in ref. 6, we obtained 90% recovery for both epimers of cefotetan and 70% recovery for iothalamic acid. Because of its simplicity for routine use, we preferred the procedure described under Materials and methods which is a
modification of other methods known from the literature [7–9]. Acetonitrile acts as an efficient protein precipitation agent. In the following extraction step with dichloromethane, neutral lipophilic substances as well as the acetonitrile used for protein precipitation, are removed. The cephalosporin remains in the undiluted aqueous layer. Using this procedure we achieved quantitative recoveries from biological fluids for the following acid compounds: cefadroxil, cefotaxime, desacetylcefotaxime, cefotiam, cefmenoxime, cefsulodin, ceftazidime and iothalamic acid. For cefotetan the recovery was 85% for epimer A and 98% for epimer B. On the other hand, the recovery for a neutral compound, namely desacetylcefotaxime lactone, was only 10%.

**Chromatography**

In Fig. 2 the chromatograms of an aqueous standard and plasma samples after simultaneous intravenous injection of 1 g of cefotetan and 20 ml of Conray 70 (containing 13.2 g of iothalamic acid) are depicted. The best compromise between run-time and separation of the isomers of cefotetan from each other and from endogenous compounds was obtained by adding small quantities (0.1 mM) of tetrabutylammonium salt to the buffered (pH 6.3–6.5) mixture of water and acetonitrile. For urine samples it was necessary to lower

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Fig. 2. Chromatograms of (1) an aqueous standard solution of cefotetan (CTN, commercially available mixture of epimers A and B) and iothalamic acid (ITS) (amount injected: ITS, 5 μg; CTN, 500 ng) and of plasma samples from a healthy volunteer 10 min (2), 2 h (3) and 6 h (4) after simultaneous intravenous injection of 1 g of cefotetan and 20 ml of Conray 70 (corresponding to 13.2 g of iothalamic acid). Concentrations (μg/ml): (2) ITS, 849; CTN-A, 56.4; CTN-B, 72.8. (3) ITS, 204; CTN-A, 29.5; CTN-B, 29.9. (4) ITS, 27.8; CTN-A, 10.2; CTN-B, 7.3.
Fig. 3. Chromatograms of (1) an aqueous standard solution of cefotetan (CTN) and iothalamic acid (ITS) (amount injected: ITS, 5 µg; CTN, 500 ng) and of urine samples from a healthy volunteer 0–2 h (2) and 4–6 h (3) after simultaneous intravenous injection of 1 g of cefotetan and 20 ml of Conray 70 (corresponding to 13.2 g of iothalamic acid). Concentrations (µg/ml): (2) ITS, 9870; CTN-A, 325; CTN-B, 502. (3) ITS, 2370; CTN-A, 376; CTN-B, 342.

The acetonitrile content and to change the concentration of the ion-pairing agent in order to maintain the separation from interfering compounds. As a consequence the overall run-time increased to 15 min compared to 8 min for plasma samples (Fig. 3).

Recovery and sensitivity

The recoveries of iothalamic acid and cefotetan were checked in the concentration range 12.5–1000 µg/ml and 1.25–100 µg/ml, respectively. Two sets of spiked plasma samples were prepared and each analysed two times on different days. The peak areas found were compared with those of standard aqueous solutions. The results are shown in Table I. The mean recoveries were 100.3 ± 2.4% for iothalamic acid, 84.6 ± 2.2% for isomer A of cefotetan and 97.6 ± 2.4% for isomer B. The recoveries from aqueous samples were 95.1 ± 2.0%, 93.7 ± 3.2% and 93.6 ± 3.7%, respectively. Obviously, the recovery of isomer A from plasma was substantially lower than the recoveries of the other two compounds. The recoveries of the individual spiked plasma samples fluctuated between 74 and 118%. For quantitation of the concentrations found in the unknown plasma samples, the mean value was used. The regression lines obtained in all cases had correlation coefficients better than 0.9996.
TABLE I

RECOVERY OF IOTHALAMIC ACID AND CEFOTETAN (EPIMERS A AND B) FROM HUMAN PLASMA

<table>
<thead>
<tr>
<th>Amount added (µg/ml)</th>
<th>Recovery* (%)</th>
<th>Iothalamic acid (ITS)</th>
<th>Cefotetan (CTN)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ITS/CTN</td>
<td></td>
<td>Epimer A</td>
<td>Epimer B</td>
</tr>
<tr>
<td>1000/100</td>
<td>100.7 ± 4.9</td>
<td>86.4 ± 1.1</td>
<td>97.4 ± 3.3</td>
</tr>
<tr>
<td>500/50</td>
<td>102.8 ± 3.8</td>
<td>87.0 ± 2.6</td>
<td>98.3 ± 1.6</td>
</tr>
<tr>
<td>250/25</td>
<td>101.7 ± 3.3</td>
<td>84.7 ± 2.6</td>
<td>97.0 ± 0.8</td>
</tr>
<tr>
<td>100/10</td>
<td>100.1 ± 3.9</td>
<td>85.5 ± 7.3</td>
<td>96.7 ± 4.3</td>
</tr>
<tr>
<td>50/5</td>
<td>102.2 ± 7.9</td>
<td>83.6 ± 5.1</td>
<td>102.5 ± 10.6</td>
</tr>
<tr>
<td>25/2.5</td>
<td>98.8 ± 3.8</td>
<td>80.4 ± 6.6</td>
<td>94.7 ± 1.5</td>
</tr>
<tr>
<td>12.5/1.25</td>
<td>95.9 ± 9.7</td>
<td>84.4 ± 11.3</td>
<td>96.6 ± 12.4</td>
</tr>
<tr>
<td>Mean ± S.D.</td>
<td>100.3 ± 2.4</td>
<td>84.6 ± 2.2</td>
<td>97.6 ± 2.4</td>
</tr>
</tbody>
</table>

*Mean ± S.D.; n = 4.

is low at 280 nm (λ_max = 240 nm; see also Figs. 2 and 3). For an injection volume of 50 µl this would correspond to a limit of detection of 20–40 ng/ml and 200–400 ng/ml, respectively. This analytical limit of detection could not be reached in biological fluids. In all plasma samples interfering compounds corresponding to up to 1 µg/ml of iothalamic acid were present, whereas in a few samples only an interfering peak corresponding to not more than 0.2 µg/ml of epimer B of cefotetan occurred. The blanks observed in urine were about ten times higher. Since up to 12 h after administration the concentrations of iothalamic acid were much higher than the blank values, no attempt was made to eliminate the unusually high blank value.

Sample stability

Especially if automatic injection systems are used for overnight runs, samples must be stable at room temperature. β-Lactam antibiotics are labile in weak basic and strong acid media. The pH of plasma samples kept at room temperature rises to 9 [10]. Therefore plasma samples containing cephalosporins should be buffered to a neutral or weak acid pH [5]. Following the preparation procedure described in this report, no degradation of cefotetan or iothalamic acid was measurable within 12 h. Moreover, at the resulting pH of 5.8–6.0 the ratio of the two epimers of cefotetan was constant too, whereas in unbuffered plasma samples (final pH 8.2–8.3 after sample clean-up) the amount of A increased and that of B decreased in samples with prevalent isomer B. Iothalamic acid was also stable in unbuffered samples.

Application

The analytical method described above was used to determine simultaneously both epimers of cefotetan and iothalamic acid in plasma and urine after intravenous injection of 1 g of cefotetan and 20 ml of Conray 70 (containing 13.2 g of iothalamic acid) for pharmacokinetic studies. Iothalamic acid is nearly
exclusively eliminated through the kidney by glomerular filtration and served in this study as marker for the function of the kidney [11]. A detailed report will be published elsewhere [12]. In Fig. 4 the mean plasma concentrations of cefotetan obtained from twelve healthy volunteers are shown. As can be seen, the time–concentration curve of isomer A is flatter. The terminal half-life was about 3 h for isomer B and 4 h for isomer A, respectively. These data are in agreement with average half-lives of cefotetan found by others [1–4].

Immediately after cessation of the injection the ratio A/B was 40:60 and the reversed after 12 h. The observed isomerisation in proteinaceous solutions was confirmed in vitro by incubation of cefotetan in phosphate buffer and 7% bovine serum albumin (BSA) at pH 7.4. As can be seen in Fig. 5, within 18 h the ratio shifted nearly to 50:50 in buffer, whereas in BSA solution the resulting ratio (70:30) was similar to that in plasma of volunteers.
DISCUSSION

Beside the microbiological assay, HPLC has been established as a standard method to determine cephalosporins in biological matrices. Because of its precision, high linear range and possibility to detect and assay active metabolites quantitatively, as in the case of the 3-acetoxycephalosporins [7, 13], HPLC is the preferred method for pharmacokinetic studies. In addition, after simultaneous application of different antibiotics, separation and quantitation of the various compounds can be achieved.

In the present paper a different pharmacokinetic behaviour of each epimer of cefotetan has been demonstrated. The reason for this effect is probably a higher ratio of protein binding of epimer A, which can be assumed from the observed lower recovery after deproteinisation with acetonitrile, and the isomerisation in BSA solution leading to another ratio of the epimers than in buffer. On the other hand, lower stability of epimer B could contribute to this effect too, because in BSA solution degradation of cefotetan is enhanced with respect to the buffer (Fig. 5). Similar results were obtained from work with moxalactam. Commercially available moxalactam preparations are also mixtures of two epimers with slightly different protein-binding ratios, biological half-lives and antimicrobial activities [14–19]. As both epimers of cefotetan have a similar antibacterial activity [20], probably no clinical consequences will result from the finding of different serum half-lives for the isomers. Whether accumulation of isomer A will occur in plasma or other body fluids after multiple doses of cefotetan seems questionable, since both epimers are presumably in equilibrium with each other. The analytical method described in the present paper could clarify this problem during further clinical trials.

Fig. 5. Degradation of cefotetan (mixture of epimers A and B) at 37°C, pH = 7.4, in 0.1 M sodium phosphate buffer (○, •) and 7% bovine serum albumin (BSA) solution (△, ▲). Increase in fraction of epimer A.
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