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Measurement of the anticancer agent gemcitabine and its deaminated metabolite at low concentrations in human plasma by liquid chromatography-mass spectrometry

Yan Xu  Bruce Keith  Jean L. Grem

Introduction

Gemcitabine (Gemzar®, Eli Lilly, Indianapolis, IN, USA; 2′,2′-difluoro-2′-deoxycytidine, dFdC) is an analog of deoxycytidine (Fig. 1), which exhibits a broad spectrum of anti-neoplastic activity [1–3]. dFdC enters cells through nucleoside transporters, where it is phosphorylated to a 5′-monophosphate (dFdCMP) by deoxycytidine kinase in a rate-limiting reaction (Fig. 2). The subsequent phosphorylations by UMP/CMP monophosphate kinase and nucleoside diphosphate kinase (NDK) generate the active 5′-diphosphate (dFdCDP) and 5′-triphosphate (dFdCTP) nucleosides. The cytotoxic effects of dFdC are attributed to the combined actions of dFdCDP, which inhibits ribonucleotide reductase, and dFdCTP, which is incorporated into DNA and results in chain termination [4]. dFdC is deaminated by cytidine deaminase to its inactive metabolite 2′,2′-difluoro-2′-deoxyuridine (dFdU).

Clinically, dFdC is used to treat a wide variety of solid tumors, both as a single agent and in combination with other cytotoxic agents. As a single agent, it is most commonly administered intravenously weekly at a dose of 1000 mg/m² by 30 min infusion. To measure dFdC and dFdU in plasma samples from cancer patients receiving dFdC with the usual
Gemcitabine (dFdC)  
MW = 263

2',2'-Difluoro-2'-Deoxyuridine (dFdU)  
MW = 264

5'-Deoxy-5-Fluorouridine (5'-DFUR)  
MW = 246

Fig. 1. The chemical structures of dFdC, dFdU and internal standard.

Fig. 2. Metabolism of gemcitabine.

Experimental

Chemicals and solutions

Gemcitabine (dFdC) was supplied by the Cancer Therapy Evaluation Program at the National Cancer Institute (Bethesda, MD, USA). It was formulated by Eli Lilly as a lyophilized product containing the equivalent of 200 or 1000 mg gemcitabine (LY188011) as the hydrochloride salt. 2',2'-Difluoro-2'-deoxyuridine (dFdU) was provided by Lilly Research Laboratories (Indianapolis, IN, USA) through a material transfer agreement with the National Cancer Institute. The internal standard, 5'-deoxy-5-fluorouridine (5'-DFUR), was obtained from Sigma (St. Louis, MO, USA). Ammonium acetate was from Aldrich (Milwaukee, WI, USA). HPLC grade water and methanol were from Fisher (Fair Lawn, NJ, USA). Tetrahydouridine (Lot No. 112907-3/22) was provided by the Drug Synthesis and
samples were subjected to solid phase extraction prior to the analysis. The Oasis® HLB cartridge was conditioned with 3 ml methanol and then 3 ml PBS. After loading the sample, the cartridge was washed with 2 ml PBS and dried with air.

Blank plasma, plasma standards and patient samples

Pooled donor plasma containing 1 µM tetratetrahydroxystamine was used as the blank plasma in this study. To prepare plasma standards, 50 µl of standard mixture working solutions of dFdC and dFdU at each concentration level and 50 µl of internal standard solution (5′-DFUR, 5.00 µg/ml) were added to 1.5-ml centrifuge tubes (Marsh Bio Products, Inc., Rochester, NY, USA). After the solutions were dried at 30 °C for 20 min in an Eppendorf Vacufuge™ (Brinkmann Instruments, Westbury, NY, USA), 500 µl of blank plasma was added to each tube. The resultant plasma standards (dFdC and dFdU) were at the concentrations of 5.00, 10.0, 50.0, 100, 500, 1000, 2000 and 5000 ng/ml with a fixed concentration of internal standard 5′-DFUR (500 µg/ml).

To prepare patient plasma samples, aliquots of 50-µl internal standard solution (5′-DFUR, 5.00 µg/ml) were added to 1.5-ml centrifuge tubes. After the solutions were dried at 30 °C for 20 min in the Eppendorf Vacufuge™, 500 µl of blank plasma was added to each tube.

The above blank plasma, plasma standards, and patient samples were subjected to solid phase extraction prior to the instrumental analysis.

Solid phase extraction

Oasis® HLB (3 cm³) solid phase extraction cartridges (Waters Corporation, Milford, MA, USA) were used for sample preparation. A 500-µl plasma sample containing 250 ng of internal standard was first diluted with an equal volume of calcium- and magnesium-free phosphate buffered saline (PBS, 1×, pH 7.4) (Invitrogen/Gibco, Grand Island, NY, USA). The Oasis® HLB cartridge was conditioned with 3 ml methanol and then 3 ml PBS. After loading the sample, the cartridge was washed with 2 ml PBS and dried with air. The analytes were eluted from the cartridge with 2 ml methanol. The eluate was collected in a glass tube (12 mm × 75 mm) and dried in the Eppendorf Vacufuge™ at 30 °C for 2 h. The residue was reconstituted in 500 µl HPLC-grade water and re-suspended. Reconstitution of the residue was carried out in 37 °C water bath for 5 min to insure complete hydration. After filtering with a GHP Acrodisc® syringe filter (0.45 µm × 13 mm) ( Pall Gelman Laboratory, Ann Arbor, MI, USA), the solution was transferred to an autosampler vial (Waters, Part No. 186000326) for the instrumental analysis.

Instrumentation

The instrumentation system consisted of a Waters 2690 separations module, a Micromass Platform LC mass spectrometric detector (electrospray ionization mode (ESI)-MS) (Waters, Milford, MA, USA), and a personal computer station with MassLynx NT (Version 3.4) software (Micromass, Manchester, UK). The analytical column was a Waters YMC ODS-AQ™ (5 µm, 120 Å, 2.0 mm × 150 mm, Part # AY125051502WT) column, which was preceded by a SecurityGuard guard column (Part # KJO-4282) (Phenomenex, Torrance, CA, USA). The eluate from the analytical column was diverted to the ESI-MS and to the waste via a PEEK microvolume connector (Cat. # MT1XCPK) (Valco Instruments, Houston, TX) with a post-column split ratio of 1:2. The smaller flow went to the ESI-MS and the larger one to the waste. High pressure PEEK tubing was used for all connections. The tubing prior to the inlet of the analytical column was 1/16 in. o.d. and 0.01 in. i.d., and the tubing after the analytical column was 1/16 in. o.d. and 0.005 in. i.d.

Chromatographic conditions

The Waters 2690 separations module were operated under the following conditions: sample temperature, 4 (±1) °C; analytical column temperature, 30 (±10) °C; sample injection volume, 50 µl. A gradient elution method was used with two solvents: solvent A was 5 mM ammonium acetate at pH 6.8 and solvent B was HPLC-grade methanol. The initial composition of the mobile phase was 98% solvent A/2% solvent B (v/v). An isocratic gradient was used at 0.2 ml/min for 3 min, followed by a linear gradient over 4 min to 70% solvent A/30% solvent B. The gradient returned to the initial conditions by a linear gradient over 2 min and remained there for 6 min prior to the next injection. The analytical column was equilibrated with the initial separation conditions prior to the initial injection. The total run time for a LC-MS analysis was 15 min per sample.

ESI-MS detection

The mass spectrometer was operated in the negative electrospray ionization mode. It was tuned by the infusion of an analyte mixture (100.0 µg/ml each in 5 mM ammonium acetate, pH 6.8) with a Harvard syringe pump (Harvard
Apparatus, South Natick, MA, USA, Cat. No. 55–1111) at flow rate of 5.0 μl/min via a 0.005 in. i.d. PEEK tubing. The tuning sample was conveyed with the incoming LC mobile phase (98% of 5 mM ammonium acetate at pH 6.8 + 2% methanol at a flow rate of 0.2 ml/min before the post-column split) in a sample tee prior to the MS detection. The optimized ionization conditions were: analytical vacuum, <1.0 × 10⁻⁶ mbar; nitrogen gas, 400 l/h, capillary, 3.00 kV; cone, −25 V (dFdC and dFdU), −35 V (5’-DFUR); ion source temperature, 140 °C; low- and high-mass resolution, 15.0; ion energy, 0.8 V, and multiplier, −650 V.

Full-scan spectra were acquired in the continuum mode over the mass range of 230–270 amu at a scan rate 200 amu/s. Single-ion-monitoring (SIM) mode was used for analyte quantitation by simultaneously monitoring quasi-molecular ions, m/z 245 for [5’-DFUR–H]⁺, m/z 262 for [dFdC–H]⁺, and m/z 263 for [dFdU–H]⁺. Data acquisition was carried out with a dwell time of 0.50 s, a span of 0.002 s, a repeat of 1, and an inter-channel delay of 0.005 s. The cone voltages were 35, 25, and 25 for m/z 245, 265, and 263, respectively.

Data analysis
Micromass MassLynx NT (Version 3.4) software was used for acquisition of the LC-MS data, construction of the internal calibration curves, and derivation of the regression equations. The dFdC and dFdU concentrations in plasma samples were determined by the regression equations after obtaining the peak area ratios of the unknowns to the internal standard over the mass range of 230–270 amu at a scan rate 200 amu/s. Single-ion-monitoring (SIM) mode was used for analyte quantitation by simultaneously monitoring quasi-molecular ions, m/z 245 for [5’-DFUR–H]⁺, m/z 262 for [dFdC–H]⁺, and m/z 263 for [dFdU–H]⁺. Data acquisition was carried out with a dwell time of 0.50 s, a span of 0.002 s, a repeat of 1, and an inter-channel delay of 0.005 s. The cone voltages were 35, 25, and 25 for m/z 245, 265, and 263, respectively.

Method validation
The recoveries were determined by comparing peak areas of the plasma samples prepared by spiking the blank plasma with known concentrations of dFdC, dFdU, and 5’-DFUR to the mean the peak areas of the control samples prepared by spiking dFdC, dFdU, and 5’-DFUR at the same levels in 5 mM ammonium acetate solution (pH 6.8). The intra- and inter-assay precisions were determined by analyzing the plasma samples with known concentrations of dFdC and dFdU within run and from different runs. The linearities of the calibration curves were assessed by the correlation coefficients. The limits of quantitation were calculated by 10 times of signal-to-noise ratio.

Patient blood sampling
Eight adult patients with advanced cancer were enrolled in an Institutional Review Board-approved Phase I trial involving dFdC as 24-h infusion weekly for three of 4 weeks. All patients provided written, informed consent. Blood samples were collected prior to (0 h) the infusion and at the steady state during the infusion (22- and 23-h) in 10 ml green-top heparinized Vacutainer® tubes (Becton Dickinson and Company, Franklin Lakes, NJ, USA) containing 10 nmol tetrahydrodoruridine as a cytidine deaminase inhibitor. The samples tubes were immediately placed on ice and centrifuged for 10 min at 800 g (4 °C). The plasma was then transferred into three labeled Nalgene® cryogenic vials (Nalge Nunc International, Rochester, NY, USA) and stored at −70 °C until analysis.

Results and discussion
ESI-MS detection
Fig. 3 shows the full-scan ESI mass spectra of dFdC, dFdU and internal standard 5’-DFUR, which indicated that detection of these compounds could be achieved by monitoring quasi-molecular ions [dFdC–H]⁺ at m/z 245, [dFdU–H]⁺ at m/z 263, and [5’-DFUR–H]⁺ at m/z 245. Hence, these ions were selected for SIM detection in the subsequent quantitative analysis. Although, the difference of m/z between [dFdC–H]⁺ and [dFdU–H]⁺ was only by 1, the specificity of the method could be further secured by choosing the proper chromatographic conditions.

Chromatographic conditions
A YMC ODS-AQ™ reversed-phase column and a gradient elution method were used for the separation of dFdC, dFdU and 5’-DFUR in human plasma. As shown in our previous studies [13], the column is well suited for the separation of nucleosides. In the current study, dFdC, dFdU and 5’-DFUR were completely resolved from each other and from endogenous compounds by the YMC ODS-AQ™ column with retention times of 11.46, 12.63 and 13.58 min, respectively (Fig. 4).

Solid phase extraction and recovery of analytes
We had previously reported that liquid-liquid extraction with two sequential acetoneitrile applications yielded the recovery of ≥70.0% for dFdC, and ≥81.8% for dFdU [6]. To simplify the procedure, the Waters polymer-based Oasis® HLB (3 cm³) extraction cartridge was evaluated for plasma sample preparation. Using this preparatory method, the absolute recoveries of dFdC, dFdU and internal standard 5’-DFUR from human plasma were found to be ≥86.8, 82.2 and 97.5%, respectively (Table 1), and the recovery data were rather consistent throughout the concentration range studied (50.0, 500, 5000 ng/ml). As compared with our liquid-liquid extraction procedure, the solid-phase extraction with Oasis® HLB cartridge is simple, easy to handle, and highly reproducible.
Fig. 3. The full-scan mass spectra of dFdC, dFdU and 5′-DFUR (I.S.). Experimental conditions were described in Section 2.6.

Fig. 4. Representative SIM chromatograms of dFdC, dFdU and 5′-DFUR in human plasma. Experimental conditions were described in Sections 2.5 and 2.6. Analyte concentration: 500 ng/ml.

Table 1
Recovery of dFdC, dFdU and I.S. in human plasma (n = 3)

<table>
<thead>
<tr>
<th>Analyte</th>
<th>50 (ng/ml)</th>
<th>500 (ng/ml)</th>
<th>5000 (ng/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>dFdC</td>
<td>% Recovery</td>
<td>% Recovery</td>
<td>% Recovery</td>
</tr>
<tr>
<td></td>
<td>%CV</td>
<td>%CV</td>
<td>%CV</td>
</tr>
<tr>
<td>dFdU</td>
<td>89.0</td>
<td>7</td>
<td>98.4</td>
</tr>
<tr>
<td>I.S.*</td>
<td>84.6</td>
<td>1</td>
<td>98.3</td>
</tr>
<tr>
<td></td>
<td>102</td>
<td>1</td>
<td>12.03</td>
</tr>
</tbody>
</table>

* I.S. (5′-DFUR) is at the fixed concentration of 500 ng/ml.

3.4. Intra- and inter-assay precision

The intra- and inter-assay precisions of the method were determined by analyzing blank plasma samples spiked with known concentrations of dFdC and dFdU (50.0, 500, 5000 ng/ml). Three replicates were analyzed at each concentration level within a single run and from runs performed on three different days. The results for dFdC and dFdU, as well as 5′-DFUR, in human plasma were shown in Table 2.

The percent coefficients of variation (%CVs) for dFdC,
Table 2
The intra-and inter-assay precision of plasma samples (n = 3)

<table>
<thead>
<tr>
<th>Analyte</th>
<th>50.0 (ng/ml)</th>
<th>500 (ng/ml)</th>
<th>5000 (ng/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Peak area</td>
<td>S.D.</td>
<td>%CV</td>
</tr>
<tr>
<td>Intra-assay</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>dFdC</td>
<td>11173</td>
<td>31</td>
<td>0.3</td>
</tr>
<tr>
<td>dFdU</td>
<td>42502</td>
<td>412</td>
<td>1</td>
</tr>
<tr>
<td>I.S. a</td>
<td>117098</td>
<td>1064</td>
<td>0.9</td>
</tr>
<tr>
<td>Inter-assay</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>dFdC</td>
<td>12101</td>
<td>796</td>
<td>7</td>
</tr>
<tr>
<td>dFdU</td>
<td>42394</td>
<td>403</td>
<td>1</td>
</tr>
<tr>
<td>I.S. a</td>
<td>118517</td>
<td>1605</td>
<td>1</td>
</tr>
</tbody>
</table>

a I.S. (5′-DFUR) is at the fixed concentration of 500 ng/ml.

Table 3
The pharmacokinetic parameters of dFdC and dFdU during initial gemcitabine infusion

<table>
<thead>
<tr>
<th>Gemcitabine (mg/(m² 24 h))</th>
<th>Patients</th>
<th>dFdC (µM/ml min)</th>
<th>dFdU (µM/ml min)</th>
<th>Cₚ (ng/ml)</th>
<th>Ratio dFdU to dFdC</th>
<th>Linear clearance (ml/min m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>150</td>
<td>3</td>
<td>12.1 ± 2.9</td>
<td>2020 ± 399</td>
<td>18.9 ± 4.4</td>
<td>16.7 ± 4.7</td>
<td>2845 ± 558</td>
</tr>
<tr>
<td></td>
<td></td>
<td>dFdC</td>
<td>dFdU</td>
<td>3053 ± 741</td>
<td>160.9 ± 11.5</td>
<td>5964 (25%, 3968, 75%, 8703)</td>
</tr>
<tr>
<td>100</td>
<td>5</td>
<td>10.3 ± 5.8</td>
<td>916 ± 333</td>
<td>16.0 ± 9.0</td>
<td>17.6 ± 10.4</td>
<td>1266 ± 438</td>
</tr>
<tr>
<td></td>
<td></td>
<td>dFdC</td>
<td>dFdU</td>
<td>1477 ± 590</td>
<td>128.0 ± 90.9</td>
<td></td>
</tr>
</tbody>
</table>

* The 22- and 23-h values for dFdC and dFdU were averaged before calculating the ratio.
at dose levels of 150 and 100 ng/(m² 24 h) were 12.1 and 10.3 µg/ml min for dFdC, and 2020 and 916 µg/ml min for dFdU. The medium clearance was 5961 ml/min m².

Conclusions

An LC-MS method has been developed and validated for the determination of dFdC and dFdU at low concentrations in human plasma. This method uses solid-phase extraction for plasma sample preparation, reversed-phase LC for analyte separation, and electrospray ionization mass spectrometry for analyte quantitation. It employs an internal standard for calibration and offers shorter chromatographic separation time (15 min), and improved limits of quantitation for plasma dFdC and dFdU. This method has been proven useful in a Phase I trial of weekly dFdC given as 24-h infusion, where the concentrations of dFdC were lower than or near the limits of quantitation of the LC-UV methods. Although we do not recommend further evaluation of the 24-h infusion schedule of gemcitabine, this LC-MS method offers advantages over the LC-UV methods in terms of shorter run times and greater sensitivity, and may prove useful in the analysis of limited volume biological samples and tissue pharmacokinetics.
Acknowledgements

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References