Rapid, Sensitive, and Quantitative LC/MS/MS Determination of Digitoxin and Digoxin in Plasma Using a Simple Procedure for Simultaneous Cleanup of Phospholipids and Proteins

Introduction
Digitoxin and digoxin are cardiac glycosides derived from the digitalis purpurea (Foxglove) plant. They have been in use for centuries for treatment of various heart conditions. Because of their narrow therapeutic range and high toxicity, their levels in patients taking digitoxin or digoxin are monitored. Immunoassay-based methods for digitoxin or digoxin exist, but are both time consuming and labor intensive. Moreover, the reported immunoassay methods are not selective toward digitoxin or digoxin due to the similarities in their chemical structures, which differ from each other in only one hydroxyl group (Figure 1). The present work was aimed at developing a rapid and selective LC/MS/MS method for the determination of digoxin and digitoxin in biological fluids. In addition, a simple sample cleanup technique for simultaneous removal of proteins and phospholipids using a zirconia-based sorbent, HybridSPE®-PLus, was explored.

Experimental
A 100 µL sample of spiked rat plasma was added to the well of a HybridSPE®-PLus 96-well plate followed by 300 µL of 1% formic acid in acetonitrile to precipitate the proteins. The plate was sealed with plastic film and agitated by vibration at 1,000 rpm for 2 minutes on a digital shaker. The plate was transferred to a vacuum manifold and the seal was removed. Vacuum (10 in. Hg) was applied for 4 minutes. The flow-through eluate was analyzed by LC/MS/MS. A Titan™ C18 column packed with 1.9 µm monodisperse, totally porous silica particles was chosen for the UHPLC separation of digitoxin and digoxin. Different mobile phase systems were studied.

LC/MS/MS Conditions
- sample prep: HybridSPE®-PLus 96-well plate
- column: Titan™ C18, 10 cm × 2.1 mm I.D., 1.9 µm particles
- mobile phase: 10 mM ammonium formate in water:methanol (20:80)
- flow rate: 0.2 mL/min
- detector: MS, ESI(+), MRM
- instrument: Shimadzu® LCMS-8030 Triple Quadrupole Mass Spectrometer

Results and Discussion
Effect of Mobile Phase Composition on MS Spectra
The effect of mobile phase composition was dramatic. Figure 2A shows digitoxin and digoxin are ionized primarily as sodium adducts in the 0.1% formic acid in acetonitrile:water (50:50) mobile phase. Because sodium ions have much higher affinity to digoxin and digitoxin in the gas phase than protons (H+), the sodium adducts are the primary ion in the 0.1% formic acid in acetonitrile-water mobile phase. Further experiments showed the sodium adduct ions are scarcely fragmented in MS/MS regardless of collision energy and gas pressure, thus not amenable to being monitored in MS/MS mode. This issue of forming sodium adducts can be overcome by replacing the mobile phase with water:methanol containing ammonium formate. Digitoxin and digoxin generate primarily ammonium adducts when ammonium formate replaces the formic acid in the mobile phase (Figure 2B). The ammonium adducts of digitoxin and digoxin are shown to be readily fragmented under mild collision conditions (Figure 3). The MRM transitions 782.5/635.5 and 798.5/651.5 were chosen for LC/MS/MS quantification of digitoxin and digoxin, respectively.

![Figure 1. Digitoxin and Digoxin](image)

**Digitoxin**
- Molecular Formula = C_{47}H_{54}O_{13}
- Monoisotopic Mass = 764.434692 Da

**Digoxin**
- Molecular Formula = C_{47}H_{54}O_{14}
- Monoisotopic Mass = 780.429607 Da
Linearity and LOD of the LC Method

Figure 4 shows the two analytes are well resolved by the Titan™ C18 column within four minutes. The limit of detection of the LC/MS/MS method was approximately 0.01 ng/mL of the analyte standard. The calibration curve of the method is linear over a range of three orders of magnitude, with a linearity ($r^2$) > 0.999 (Figure 5).

Comparison of Sample Prep Techniques

Protein precipitation is widely used for sample preparation of biological matrices prior to LC/MS analysis. The approach is effective in eliminating proteins, but not in removing phospholipids. Indeed, multiple species of phospholipids were present in high amounts in the rat plasma after protein precipitation as reflected by their intense MS signals (Figure 6A). This study exploited HybridSPE®-Plus which specifically and effectively eliminated both precipitated proteins and phospholipids from the plasma samples (Figure 6B). The mechanism behind the effectiveness of HybridSPE®-Plus technology is based on the unique Lewis acid/Lewis base interaction between the zirconia and the phosphate group of phospholipids.

Figure 7 compares the effectiveness of the two sample preparation techniques, standard protein precipitation and HybridSPE®-Plus, on the LC/MS/MS signals. The ion signals of both analytes of the samples prepared by standard protein precipitation were only one-third to one-fourth of those of the samples prepared by HybridSPE®-Plus, resulting in low recoveries (Table 1). Furthermore, the ion signals of both analytes following standard protein precipitation decreased with increasing number of injections, likely from build-up of phospholipids on the column, leading to irreproducible results and shortened column lifetime. In contrast, the samples prepared by the HybridSPE®-Plus exhibited high recoveries (>90%) with good reproducibility (<6% RSD). Matrix effects were essentially eliminated by using the HybridSPE®-Plus method.
Figure 2B. MS Spectra of Digoxin and Digitoxin in Ammonium Formate Mobile Phase

mobile phase: 10 mM ammonium formate in water:methanol (50:50)

![MS Spectra](image)

Table 1. Analyte Recovery from Plasma Using Standard Protein Precipitation and HybridSPE®-PLus Methods

<table>
<thead>
<tr>
<th></th>
<th>Protein Precipitation Method</th>
<th>HybridSPE®-PLus Method</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MRM Quantifier Ions</td>
<td>Recovery (%)</td>
</tr>
<tr>
<td>Digitoxin</td>
<td>782.5/635.5</td>
<td>27.0</td>
</tr>
<tr>
<td>Digoxin</td>
<td>798.5/651.5</td>
<td>35.4</td>
</tr>
</tbody>
</table>

n = 37
Figure 3. MS/MS Spectra of Digitoxin and Digoxin (Ammonium Adducts)

**mobile phase:** 10 mM ammonium formate in water:methanol (50:50)

**Digitoxin** (precursor ion 782.5 m/z)

- 113.1
- 243.3
- 277.2
- 336.9
- 373.5
- 419.6
- 505.5
- 617.7
- 635.4
- 782.7

**Digoxin** (precursor ion 798.5 m/z)

- 1002.0
- 1004.0
- 1006.0
- 1008.0

**Intensity (×10,000)**

Figure 4. LC/MS/MS of Digitoxin and Digoxin in Blank and Spiked (10 ng/mL) Rat Plasma on Titan™ C18 after SPE using HybridSPE®-PLus

**Peak**
1. Digoxin
2. Digitoxin

**MRM(m/z)**
- 798.5/651.5
- 782.5/635.5

**Column:** Titan™ C18, 10 cm × 2.1 mm I.D., 1.9 µm particles (577124-U)

**Mobile phase:** 10 mM ammonium formate in water:methanol (20:80)

**Flow rate:** 0.2 mL/min

**Pressure:** 4550 psi

**Column temp.:** 35°C

**Detector:** MS, ESI(+), MRM

**Injection:** 2 µL

**Sample:** Rat plasma, blank and spiked at 10 ng/mL, extracted with HybridSPE®-PLus

**Instrument:** Shimadzu® LCMS-8030
Figure 5. LC/MS/MS Calibration Curves for Digitoxin and Digoxin

![Digitoxin Calibration Curve](image1)

Digitoxin

![Digoxin Calibration Curve](image2)

Digoxin

$r^2 = 0.9996$

$r^2 = 0.9991$

Figure 6. Absence of Phospholipids in ESI-MS Spectra of Plasma Prepared by HybridSPE®-PLus Compared to Protein Precipitation

![ESI-MS Spectra Comparison](image3)

A. Protein Precipitation

<table>
<thead>
<tr>
<th>m/z</th>
<th>Intensity (×1,000,000)</th>
</tr>
</thead>
<tbody>
<tr>
<td>227.0</td>
<td>258.1</td>
</tr>
<tr>
<td>369.4</td>
<td>496.4</td>
</tr>
<tr>
<td>524.4</td>
<td>758.7</td>
</tr>
</tbody>
</table>

*Indicates phospholipids which are confirmed by MS/MS spectra with product ion of 184/104.

B. HybridSPE®-PLus

<table>
<thead>
<tr>
<th>m/z</th>
<th>Intensity (×1,000,000)</th>
</tr>
</thead>
<tbody>
<tr>
<td>229.2</td>
<td></td>
</tr>
</tbody>
</table>

![m/z Intensity Graph](image4)
Summary
A rapid and sensitive LC/MS/MS method has been developed for the determination of digitoxin and digoxin in plasma. A Titan™ C18 UHPLC column with monodisperse particles was employed to resolve the two analytes within four minutes. Severe matrix effects, including ion suppression and irreproducibility, that were observed using standard protein precipitation procedures were overcome by utilizing HybridSPE-PLus, an innovative technique for simple and quick sample cleanup of phospholipids and proteins in biological matrices.

References