A simplified method for therapeutic drug monitoring of mitotane by gas chromatography-electron ionization-mass spectrometry

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Abstract
Mitotane is a key drug for the treatment of adrenal cortical carcinoma. Due to its narrow therapeutic window, 14–20 μg/mL, monitoring its concentration is crucially important. In this study, a simplified method for measuring mitotane in plasma using gas chromatography-electron ionization-mass spectrometry (GC-EI-MS) was developed. Through deproteination and liquid–liquid extraction, mitotane and an internal standard (IS) were extracted from plasma samples. GC-EI-MS yielded retention times of 8.2 and 8.7 min, for mitotane and the IS, respectively, with a total run time of 12 min. Selectivity and intra-/inter-batch accuracy and precision analyses provided a lower limit of quantification of 0.25 μg/mL, and a calibration curve between 0.25 and 40 μg/mL had good linearity (coefficient of determination = 0.992). The matrix effect factor and percent recovery of the method had good precision. Additionally, long-term sample stability was observed below 4°C. In a clinical setting, mitotane levels in plasma from an adrenal cortical carcinoma patient were within calibration range. Therefore, this simplified method can be applied to routine therapeutic drug monitoring of mitotane, which may contribute to improved treatment of adrenal cortical carcinoma.

Keywords
adrenal cortical carcinoma, electron ionization, gas chromatography, mitotane, therapeutic drug monitoring

1 | INTRODUCTION

Mitotane, a key drug for the treatment of adrenal cortical carcinoma (ACC), has a narrow therapeutic window between 14 and 20 μg/mL; therefore, it is desirable to routinely perform therapeutic drug monitoring (TDM). Although several quantitative methods for enantiomers and major metabolites of mitotane have been developed (Cantillana, Lindström, Eriksson, Brandt, & Bergman, 2009; Inouye, Mio, & Sumino, 1987), the lack of antitumor effects of a major metabolite, 1,1-(o,p’-dichlorodiphenyl) acetic acid, was recently demonstrated (Hescot et al., 2014), suggesting the low importance of TDM of these compounds compared to mitotane. Therefore, a simplified and rapid quantitative method for TDM of mitotane is clinically desired, compared to those of metabolites and chiral bodies.

Although validated liquid chromatography (LC) methods for the measurement of blood (including plasma and serum) mitotane levels...
have been abundantly reported (Andersen, Warren, Nome, Vesterhus, & Slerdal, 1995; Benecke, Vetter, & De Zeeuw, 1987; De Francia et al., 2006; Garg, Sakoff, & Ackland, 2011; Mooienaar, Niewint, & Oei, 1977; Sinsheimer et al., 1996), gas chromatography (GC) methods are rare (Inouye et al., 1987), despite GC being a major analytical method along with LC. Inouye et al. (1987), previously reported a measurement method for plasma mitotane levels using GC-electron ionisation-mass spectrometry (GC-EI-MS); however, this method required a 1 h heating of plasma-absorbed filter paper to liberate mitotane. Considering the fast-paced clinical requirements, these conditions are not applicable for routine TDM of mitotane. Furthermore, a measurement method involving a simplified sample preparation procedure has been recently reported by Feliu et al. (2017); however, this method involves the use of an ultra-high performance LC (UPLC) system, which may currently be less common than LC or GC systems, although UPLC may be more available in future decades. In addition, this procedure does not involve the use of an internal standard (IS), which creates the difficulty of judging whether the observation of a low mitotane peak is due to a low rate of extraction or a low presence of mitotane. Therefore, for the implementation of routine and on-site analyses, we determined that a validated method using a common chromatography system, an internal reference, and a simplified and user-friendly sample preparation procedure was required.

In this study, we aimed to develop a widely usable method for the measurement of plasma mitotane levels by GC-EI-MS involving a simplified pretreatment procedure in order to facilitate routine TDM of mitotane.

### EXPERIMENTAL

#### 2.1 Materials

Details are provided in Supporting Information.

#### 2.2 Samples for the calibration curve and quality control

The concentrations of mitotane in pooled human plasma for the calibration curve were 0.25, 0.5, 1, 10, 20, 30, and 40 μg/mL. Quality control (QC) samples (n = 5 for each concentration) for testing intra- and inter-batch variability were 0.25, 20, and 40 μg/mL.

#### 2.3 Sample pretreatment

For deproteination, 40 μL of IS (12.5 μg/mL)-containing methanol was added to each patient plasma or prepared plasma sample used for the calibration curve and QC. To each sample, 150 μL of ethyl acetate was added, followed by vortexing for 10 min. The samples were then centrifuged at 10,000 rpm for 5 min, and the supernatants (120 μL) were collected for GC-EI-MS analysis.

### RESULTS AND DISCUSSION

#### 3.1 Method validation

#### 3.1.1 Selectivity

Two major chromatographic peaks with retention times of 8.2 and 8.7 min, respectively, were observed while the total runtime was 12 min (Figure S1). Based on their mass spectra, these peaks were identified as mitotane and the IS, respectively (Figure S2). In testing selectivity, plasma containing 0.25 μg/mL of mitotane showed a peak having a signal-to-noise ratio 5 for six individual blank plasma samples (Figure S3); no interference from the IS occurred (Figure S3).
3.1.2 | Intra- and inter-batch variability

As shown in Table 1, acceptable intra- and inter-batch variability was observed. Good linearity of the calibration curve (coefficient of determination = 0.992) was also confirmed between 0.25 and 40 μg/mL (Figure FIGURE 1). Based on selectivity, calibration curve linearity, and variability, 0.25 μg/mL was established as the lower limit of quantification (LLOQ) of the method. It should be noted that the calibration range and LLOQ obtained are superior to those of previously reported methods (Andersen et al., 1995; Benecke et al., 1987; De Francia et al., 2006; Feliu et al., 2017; Garg et al., 2011; Inouye et al., 1987; Mornar, Sertić, Turk, Nigović, & Koršić, 2012).

3.1.3 | Matrix effect factor and percent recovery

The matrix effect factor and percent recovery had acceptable precision. An unexpectedly large recovery rate (mean = 127.5%) was observed at 40 μg/mL (Table S1), although this did not have a significant effect on calibration linearity or intra-/inter-batch variability.

3.1.4 | Stability

Acceptable accuracy and precision were observed for all stability-test conditions, with the exceptions of 40 μg/mL mitotane samples stored at 25°C for 14 and 28 days (Table S2). Therefore, if sample storage is required, it should be performed at ≤4°C.

3.2 | Clinical applicability of the established method

Clinical samples were obtained from an ACC patient, and the plasma mitotane levels were quantitated (Figure S1). The average mitotane concentration was 10.7 μg/mL (median = 9.1 μg/mL), with minimum and maximum detected levels of 8.6 and 14.5 μg/mL, respectively (Table S3). All clinical mitotane levels were within the calibration range of the established method. Thus, this newly developed, validated, and simplified quantitative method for mitotane in plasma is considered applicable to routine TDM, because our method requires only 15 min for deproteination compared to the 1 h pretreatment time required in the method by Inouye et al. (1987). We believe that our method with a simplified and sped-up procedure for sample pretreatment may be more compatible with a clinical setting.

4 | CONCLUSIONS

A simplified, validated, and clinically applicable GC-EI-MS method for mitotane in plasma was developed and determined to be applicable for routine TDM. Treatment involving TDM of mitotane will likely contribute to improved success of treatment of ACC patients.

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CONFLICT OF INTEREST

The authors declare no conflicts of interest.

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SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section at the end of this article.

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