Status of sennosides content in various Indian herbal formulations: Method standardization by HPTLC

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Abstract

Several poly-herbal formulations containing senna (Cassia angustifolia) leaves are available in the Indian market for the treatment of constipation. The purgative effect of senna is due to the presence of two unique hydroxyl anthraquinone glycosides sennosides A and B. A HPTLC method for the quantitative analysis of sennosides A and B present in the formulation has been developed. Methanol extract of the formulations was analyzed on a silica gel 60 GF254 HPTLC plates with spot visualization under UV and scanning at 350 nm in absorption/reflection mode. Calibration curves were found to be linear in the range 200-1,000 ng. The correlation coefficients were found to be 0.991 for sennoside A and 0.997 for sennoside B. The average recovery rate was 95% for sennoside A and 97% for sennoside B showing the reliability and reproducibility of the method. Limit of detection and quantification were determined as 0.05 and 0.25 µg/g respectively. The validity of the method with respect to analysis was confirmed by comparing the UV spectra of the herbal formulations with that of the standard within the same Rf window. The analysis revealed a significant variation in sennosides content.

Introduction

Cassia angustifolia (Family: Caesalpiniaceae), popularly known as senna, is a valuable plant drug in ayurvedic and modern system of medicine for the treatment of constipation (Atal et al., 1982; Das et al., 2003; Martindale, 1977; Sharma, 2004). Sennoside A and B are the two anthraquinone glycosides that are responsible for purgative action of senna. A variety of poly-herbal formulations containing senna leaves are available in India to relieve constipation and allied troubles. Senna is a strong purgative that should be taken in proper dosage otherwise it may lead to gripping and colon problem (Bhattacharjee, 2004). Different analytical techniques, viz, thin layer chromatography, spectrophotometry, column chromatography have been reported in the literature for estimation of sennosides (Azam et al., 2002; Bala et al., 2000; Habib et al., 1980; Lin et al., 1998). HPTLC method has also been reported (Shah et al., 2000). But these techniques are tedious and time consuming. The present paper reports a simple, rapid and accurate HPTLC method for analysis of sennoside A and B occurring in herbal formulations and to make an assessment on the status of sennoside A and B content in the laxative formulations.
Materials and Methods

Ten different branded herbal formulations (poly-herbal, powdered/microgranules) were collected from herbal shops of local markets.

Analytical standards of sennoside A and B were obtained from M/s. ChromaDex, Santa Ana, CA, USA. Solvents (methanol, 2-propanol, ethyl acetate, formic acid) used in entire study were from Merck, India.

Preparation of standard solutions: Standard solutions of sennoside A and sennoside B (1 mg/5 ml) were prepared in methanol.

Extraction of samples: Each 4 g of different branded formulation was sonnicated with 70% methanol (3 x 20 ml) for about 45 min. Then the extract was filtered in a Buchner funnel using Whatman No. 1 filter paper and was concentrated under vacuum in a rotary evaporator at 50°C, redissolved in methanol and finally reconstituted in 20 ml methanol prior to HPTLC analysis.

Chromatographic parameters: Chromatography was performed on glass-backed silica gel 60 GF\textsubscript{254} HPTLC layers (20 cm x 20 cm; 0.30 mm layer thickness) prepared using a Camag TLC plate auto coater (Camag, Switzerland). Samples and standard compounds 1 and 2 of known concentrations were applied as 8 mm wide bands using Camag Linomat 5 automated TLC applicator (Camag, Muttenz, Switzerland) with nitrogen flow providing delivery speed 150 nl/s from the syringe. These parameters were kept constant throughout the analysis.

Detection and quantification of sennosides: After completion of sample application, the plate was developed in a Camag Twin through glass tank presaturated with mobile phase of 2-propanol: ethyl acetate: water: formic acid (17:19:12:2). The wavelength 350 nm was found to give the highest sensitivity. Linearity of sennoside A and B was found in the concentration range 200-1000 ng. Regression analysis of the experimental data points showed a linear relationship with excellent correlation coefficient(r) of sennoside A and sennoside B of 0.991 and 0.997, respectively [Table I]. The average recovery rate was 95% for sennoside A and 97% for sennoside B [Table II] showing the reliability and reproducibility of the method. In this study the limit of detection (LOD) was 0.1 µg per gram of the formulations.

Results and Discussions

Figure 1 shows the distinct separate bands of sennoside A and B by TLC. The high resolution and reproducible peaks were obtained [Figure 2] by using the mobile phase 2-propanol: ethyl acetate: water: formic acid (17:19:12:2). The wavelength 350 nm was found to give the highest sensitivity. Linearity of sennoside A and B was found in the concentration range 200-1000 ng. Regression analysis of the experimental data points showed a linear relationship with excellent correlation coefficient(r) of sennoside A and sennoside B of 0.991 and 0.997, respectively [Table I]. The average recovery rate was 95% for sennoside A and 97% for sennoside B [Table II] showing the reliability and reproducibility of the method. In this study the limit of detection (LOD) was 0.1 µg per gram of the formulations.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Rf value</th>
<th>Regression equation</th>
<th>r</th>
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<tbody>
<tr>
<td>Sennoside A</td>
<td>0.84</td>
<td>Y=7.087X+107.744</td>
<td>0.991</td>
</tr>
<tr>
<td>Sennoside B</td>
<td>0.63</td>
<td>Y=3.589X+702.086</td>
<td>0.997</td>
</tr>
</tbody>
</table>
and limit of quantification of sennoside A and B were determined to be 0.05 and 0.25 µg/g. The result [Table III] showed that the relative amount of sennoside A and B in formulation-1 was highest and in formulation-10 it was lowest. The result reflected a 2.75 times higher concentration in formulation-1 compared to that of formulation-10. The minimum content of sennoside B (considering the highest content as 100%) was shown only by 5.41% in case of formulation-10 and less than 17.5% in case of formulation number ‘3’ to ‘9’. This result clearly revealed that there is a significant variation...
of sennoside A and B content in popular herbal formulations. Since the chemical contents are biologically active, variation in their daily dose may impact on health benefits. Therefore, it is necessary that herbal formulations should always be analyzed to maintain the quality of the herbs present in the formulation.

In conclusion, the proposed HPTLC method can be used for the determination of sennoside A and B in various commercial formulations for quality evaluation. Method is very simple, rapid and suitable for rapid screening of plant materials for genotypic assessment and can be performed without any special sample pretreatment. The method is less expensive than an LC method. The method is, thus, suitable for quality control laboratories, where economical use of time and space is essential.

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