LC-ESI/MS determination of xanthone and secoiridoid glycosides from *in vitro* regenerated and *in vivo* *Swertia chirayita*

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Abstract A rapid analytical method has been developed to determine xanthone and secoiridoid glycoside in *in vitro* and *in vivo* *Swertia chirayita* extracts. Ultra performance liquid chromatography–electrospray ionization mass spectrometry (LC-ESI/MS) was applied and validated for the analysis of xanthone and secoiridoid glycoside a potential active component isolated from methanolic extracts of *in vitro* and *in vivo* *Swertia chirayita* plantlets. Chromatographic separation was achieved on a RP-C18 column using gradient elution. Mangiferin (Xanthone), Amarogentin and Swertiamarin (Secoiridoid glycosides) were identified in both the extracts. In the LC/ESI-MS spectra, major $[M+H]^+$ and $[M+Na]^+$ ions were observed in positive ion mode and provided molecular mass information. An ultra-performance liquid-chromatography in combination with electrospray ionization tandem mass spectrometry involving metal cationisation was successfully utilized for the rapid identification of xanthone and secoiridoid glycosides. This method is suitable for the routine analysis, as well as for the separation and identification of known and novel secoiridoid glycoside and xanthone.

Keywords *Swertia chirayita* · LC-ESI/MS · Xanthone · Secoiridoid glycoside

Introduction

S. chirayita (Roxb. ex Fleming. H. Karst.) is used as herbal medicine for various health ailments and it has been used in Unani medicine (Joshi and Dhawan 2005). Extracts of *S. chirayita* have been shown to possess antioxidative, antihepatotoxic and hypoglycemic, anti-inflammatory, antimalarial, anticarcinogenic, and antimicrobial activities (Kar et al. 2003; Saha et al. 2004; Tripathi et al. 2005). Previous reports documented the presence of flavonoids, xanthenes, terpenoids, iridoid and secoiridoid glycoside, that are responsible for therapeutic properties in *S. chirayita* (Pant et al. 2000). The bioactive constituents include the xanthone and secoiridoid glycosides consisting of mangiferin, amarogentin and swertiamarin. Mangiferin has been reported to possess various biological activities like antitumour, antiviral, antioxidant, antidiabetic and immunomodulatory activity (Guha et al. 1996; Sanchez et al. 2000; Garcia et al. 2003). Swertiamarin, and Amarogentin compounds also possesses various biological activities such as chemopreventive, antibacterial, anticholinergic and antihepatitis activity and served as important chemotaxonomic markers (Saha et al. 2006; Yamahara et al. 1991; El-Sedawy et al. 1989). Mass spectrometry (MS), especially coupled with electrospray ionization (ESI), is an important physiochemical method for the analysis of bioactive compounds. Protonated ions are usually generated by ESI-MS in positive ion mode (He et al. 2013). Identification of compounds in botanical extracts is established by a combination of useful online system liquid chromatography (LC) photodiode array UV–vis detection and electrospray ionisation mass spectrometry (ESI-MS) (He et al. 1998; Cubbon et al. 2010). Swertiamarin, Gentiopicroin and Mangiferin were identified by TLC in *Swertia chirayita* and its content was determined by HPLC (Yuan-can et al. 2010). Determination of five active components in *Swertia chirayita* by HPLC is also reported by Lei et al. (2010). Very few reports have been documented the analysis of xanthone and secoiridoid glycosides

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Fig. 1 **a.** UPLC chromatogram of Mangiferin (Standard compound). **b.** UPLC chromatogram of Swertiamarin (Standard compound). **c.** UPLC chromatogram of Amarogentin (Standard compound). **D.** Total ion

chromatogram of *in vitro* plant extract of *S. chirayita*. **E.** Total ion chromatogram of *in vivo* plant extract of *S. chirayita*

Fig. 1 (continued)

Fig. 1 (continued)

composition of wild *S. chirayita* by using liquid chromatography/tandem mass spectrometry (Suryawanshi et al. 2006, 2007). Hence, the present investigation was undertaken with the objective of to present a LC-ESI/MS based technique for the identification and analysis of xanthone (mangiferin) and secoiridoid glycosides (swertiamarin, amarogentin) in methanolic extracts of *in vitro* regenerated and *in vivo* *S. chirayita*. The study was performed in positive ion mode.

Experimental

Reagents and standards

Acetonitrile (HPLC grade) was purchased from Thomas Baker (Mumbai, India). Ultra-pure water was obtained from a MilliQ PLUS purification system (Millipore, USA). All the three standards (Mangiferin, Amarogentin and Swertiamarin) were procured from Chromadex™.

Plant material

Young and mature *S. chirayita* plants were collected from natural habitat of Darjeeling, West Bengal during the month

of November. Following the procedure described by Kumar et al. (2014) leaf explants excised from young plants were decontaminated in 0.2% Bavastin solution followed by (5–6 drops/100 mL) Tween 20 (20 min) and 0.1% mercuric chloride (8 min). The leaf explants were rinsed three times in sterile distilled water, cut into approximately 1 cm³ cubes and inoculated onto Murashige and Skoog (MS) (1962) medium supplemented with 30 g L⁻¹ sucrose, 0.1 g L⁻¹ myo inositol and 1.0 mg L⁻¹ 6- Benzyladenine (BA), 100 mg L⁻¹ Adenine sulphate (Ads), 0.1 mg L⁻¹ Indole-3-acetic acid (IAA), and 0.8 % agar, pH 5.8. *In vitro* shoots were maintained in a growth chamber under a 16 h-photoperiod at the light intensity of 40 μ mol s⁻¹ m⁻² provided by white fluorescent tubes and temperature 25 $\pm$ 2 °C. The shoots were subcultured every 4 weeks.

Preparation of plant extracts

In vitro regenerated and wild plants were freeze-dried and lyophilized. The ground materials (100 mg/mL) were homogenized with methanol. The resultant extracts were centrifuged for 10 min at 3000 rpm and the supernatant was retained for analysis.

Fig. 2 **a.** LC/ESI-MS/MS spectra of Mangiferin (m/z 423) in *in vitro* *S. chirayita* extract. **b.** LC/ESI-MS/MS spectra of Amarogentin (m/z 587) in *in vitro* *S. chirayita* extract. **c.** LC/ESI-MS/MS spectra of Swertiamarin (m/z 375) in *in vitro* *S. chirayita* extract. **d.** LC/ESI-MS/MS spectra of

Mangiferin (m/z 423) in *in vivo* *S. chirayita* extract. **e.** LC/ESI-MS/MS spectra of Amarogentin (m/z 587) in *in vivo* *S. chirayita* extract. **f.** LC/ESI-MS/MS spectra of Swertiamarin (m/z 375) in *in vivo* *S. chirayita* extract

Fig. 2 (continued)

Fig. 2 (continued)

Table 1 The ions detected (*m/z*) in LC/ESI-MS spectra with positive ion mode at constant cone voltage in *in vitro* regenerated *S. chirayita* extract

Mode	Compound	[M+H] ⁺	[M+Na] ⁺	[M+K] ⁺	[M+NH ₄] ⁺
ESI+	Mangiferin	423	445	461	–
	Amarogentin	587	–	–	604
	Swertiamarin	375	–	413	–

Separation of secondary metabolites using UPLC

Analytes were separated using a Waters Acquity UPLC system (Waters ACQUITY QSM) consisting of an Acquity UPLC binary systems manager, an Acquity UPLC sample manager, a PDA detector and an H/T column heater containing an Acquity UPLC BEH C₁₈ reverse phase column (2.1 mm×50 mm; 1.7 μm particle size). Several mobile phases (acetonitrile, methanol, water with and without additives like 0.1% formic acid and 5 mM ammonium acetate pH 4.2) and different gradient elution programs were tested to choose and optimise the best analytical conditions for simultaneous detection of mangiferin, amarogentin and swertiamarin. The binary mobile phase optimised consisted of (A) acetonitrile and (B) 5 mM ammonium acetate pH4.2. A linear gradient (curve no. 6) elution program was applied as follows: Initial: 10% A, 90% B; 0–4 min: 40% A, 60% B; 4–6 min: 70%A, 30%B; 6–8: 80%A, 20%B; 8–10: 10%A, 90%B; 10–15: 10%A, 90%B. The flow rate was maintained at 0.5 ml min⁻¹ and the temperature of the column and sample manager were set at 40 and 5 °C respectively. Injection volumes were 2 μl for standards as well as for samples. The pressure limits were set at 0 psi low and 15,000 psi high, the highest pressure observed was 9346 psi during the elution process. The PDA (Photo Diode Array) detector was set at 254 nm and instrument operations, data acquisition and processing were performed using EmPower2 chromatographic data software. The Mangiferin, Amarogentin and Swertiamarin peaks from samples were identified by the comparison of retention times with the corresponding retention times of standards.

Table 2 The ions detected (*m/z*) in LC/ESI-MS spectra with positive ion mode at constant cone voltage in *in vivo* *S. chirayita* extract

Mode	Compound	[M+H] ⁺	[M+Na] ⁺	[M+K] ⁺	[M+NH ₄] ⁺
ESI+	Mangiferin	423	–	461	–
	Amarogentin	587	–	–	604
	Swertiamarin	375	397	413	–

Mass spectrometry conditions

To confirm the identification of secondary metabolites in *in vitro* and *in vivo* plantlets of *S. chirayita*, samples were subjected to UPLC and separations were performed. The TUV eluent was sent to Electrospray Ionisation Mass Spectrometer (ESI-MS) operated in positive ion mode. Probe and source conditions included capillary voltage 3.50 kV, 350 °C desolvation temperature, 50 L h⁻¹ cone gas, 650 L h⁻¹ desolvation gas and 110 °C block temperature. Mangiferin, Swertiamarin and Amarogentin were identified based on comparison of the mass fragmentation tandem MS analysis data with standard compounds.

Results and discussion

The capacity of *in vitro* cultures to accumulate secondary metabolites is an interesting biochemical phenomenon in plant cell biotechnology applications. *S. chirayita* plants has a diverse chemical profile (Joshi and Dhawan 2005). The xanthone (Mangiferin) and secoiridoid glycosides (Amarogentin and Swertiamarin) profiles of *in vitro* regenerated and wild plant extracts were analysed using LC-ESI/MS (Figs. 1 and 2; Table 5). For analysis, three reference compounds procured from chromadex™ viz. swertiamarin, mangiferin and amarogentin were used in the experiment. The results indicate similarity in presence of mangiferin, amarogentin and swertiamarin between *in vitro* and wild plant extracts. There was a variation in the concentration of mangiferin, amarogentin and swertiamarin between both the extracts. The concentration of Amarogentin was higher in wild plant extracts compared to a *in vitro* regenerated plant extracts (Table 5). The concentration of mangiferin and swertiamarin were higher in *in vitro* regenerated compared to wild plants (Table 5). This finding is concordant with the results of a similar study in *Centaurium erythraea* (Jankovic et al. 2011). He also reported higher amounts of xanthone compounds in *in vitro* raised plantlets in comparison to naturally growing plantlets. Similarly, higher concentration of mangiferin and swertiamarin in *in vitro* compared to wild plant extracts has been recently reported by Kumar et al. (2014).

Initially, full scan LC/ESI-MS spectra of *in vivo* and *in vitro* *S. chirayita* methanolic extracts was acquired in positive ion mode utilizing the LC gradient condition as described earlier with a mobile phase consisting of acetonitrile and 5 mM ammonium acetate pH4.2. LC/ESI-MS spectral data of *in vivo* and *in vitro* *S. chirayita* methanolic extracts are presented in Tables 1 and 2 and shows a series of peaks between *m/z* 200–1000. In positive ion mode of LC/ESI-MS experiments, all components in *in vitro* extracts showed highly

Table 3 LC/ESI-MS/MS characterization of components in *in vitro* regenerated *Swertia chirayita* extract

Compound	M.W.	Precursor ion	Product ion (m/z)
Mangiferin	422	423	405, 387, 369, 357, 327, 303, 284, 273, 199
Amarogentin	586	587	391, 373, 359, 311, 289, 247, 229, 197
Swertiamarin	374	375	357, 339, 303, 284, 273, 213, 195, 177, 167

Table 4 LC/ESI-MS/MS characterization of components in *in vivo* *Swertia chirayita* extract

Compound	MW	Precursor ion	Product ion (m/z)
Mangiferin	422	423	405, 387, 369, 357, 327, 303, 273, 239, 223, 209, 201
Amarogentin	586	587	391, 373, 359, 311, 281, 247, 229, 197
Swertiamarin	374	375	357, 327, 303, 284, 256, 247, 223, 213, 195, 183, 177, 167

abundant proton and sodium ion adducts but a relatively lower proportion of potassium and ammonium adducts was also detected (Table 1). In case of *in vivo* extracts all three components showed highly abundant proton ion adducts but a relatively lower proportion of sodium, potassium and ammonium adducts were detected (Table 2). Similar results were obtained by Suryawanshi et al. (2006, 2007) in *in vivo* methanolic extracts of *S. chirayita*. The Collision induced dissociation (CID) spectra and mass spectral data of *in vivo* and *in vitro* extracts of fragment ions of all three compounds in positive ion mode are given in Tables 3 and 4. The CID spectra of the $[M+H]^+$ ion at m/z 423 (Mangiferin) in both extracts (*in vivo* and *in vitro*) showed fragment ions at m/z 405 and 387 corresponding to subsequent loss of two water units, $[M+H-18]^+$ and $[M+H-36]^+$, respectively. A similar result was found by Suryawanshi et al. (2006). The CID spectrum of the $[M+H]^+$ ion of Amarogentin (see Fig. 2b and e) in *in vitro* and *in vivo* extracts contains fragment ions at m/z 391, 373, 359, 311, 289, 247, 229, 197 and m/z 391, 373, 359, 311, 281, 247, 229, 197 respectively. Highly abundant $[M+Na]^+$ ions completely replaced the protonated ion formation. Usually, $[M+Na]^+$ ions are found to be very stable and produce only negligible amounts of fragment ions (Suryawanshi et al. 2006). Similarly this unfavourable phenomenon did not occur with swertiamarin. As we observed LC/ESI-MS spectra in positive ion mode for swertiamarin in *in vivo* and *in vitro* extracts. The CID spectrum of the $[M+H]^+$ ion of swertiamarin (see Fig. 2c and f) in *in vitro* and *in vivo* extracts contains fragment ions at m/z 357, 327, 303, 284, 256, 247,

223, 213, 195, 183, 177 and 167. All three compounds (mangiferin, amarogentin and swertiamarin) were analyzed in *in vitro* and *in vivo* plantlets of *S. chirayita*. Table 5 shows the total mangiferin, amarogentin and swertiamarin, contents in *in vitro* and *in vivo* plantlets of *S. chirayita*. The amount of mangiferin ($11.48 \pm 8.41 \mu\text{g/mL}^{-1}$) and swertiamarin ($15.16 \pm 0.44 \mu\text{g/mL}^{-1}$) was higher in *in vitro* plantlets of *S. chirayita* whereas amarogentin ($30.45 \pm 0.68 \mu\text{g/mL}^{-1}$) was higher in a *in vivo* plantlets.

In conclusion, a rapid and convenient LC/ESI-MS technique was developed and validated for the extraction and identification of secoiridoid glycosides (Amarogentin and Swertiamarin) and xanthone (Mangiferin) from *in vitro* and *in vivo* methanolic extracts of *S. chirayita*, an important medicinal plant. Major $[M+H]^+$ and $[M+Na]^+$ ions were observed in LC/ESI-MS with positive ion mode and provided molecular mass information. More detailed study in relation of component identification and analysis is still awaited.

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Conflict of interest Authors declare no conflicts of interest. The authors alone are responsible for the content and writing of the article.

Table 5 Mangiferin, Amarogentin and Swertiamarin profile ($\mu\text{g/mL}$) of *in vitro* and *in vivo* *S. chirayita* plant extracts

Plant extract	Mangiferin	Amarogentin	Swertiamarin
<i>In vitro</i> plantlets	11.48 ± 8.41	28.96 ± 10.38	15.16 ± 0.44
<i>In vivo</i> plantlets	4.63 ± 0.39	30.45 ± 0.68	9.94 ± 4.19

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