Distribution of Silymarin in the Fruit of *Silybum marianum* L.

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**Abstract**

The fruit of *Silybum marianum* or milk thistle is known for its rich contents of flavonolignan compounds known as silymarin. Silymarin has been recognized for centuries as “liver tonics” and is well known to prevent or reverse hepatotoxicity. Silymarin is usually extracted from the defatted fruits in methanol in a yield of less than 2%. Fluorescent microscopy, scanning electron microscopy, accurate mass spectrometry as well as infra-red and Raman spectroscopy were used for mapping silymarin in the fruit. Silymarin was found to be only located in the pericarp section of the fruit. Extraction of silymarin from the pericarp gave higher yields of more than 6% and did not require defatting.

**Keywords:** Antioxidants; Liver protection; Flavonolignan; Milk thistle; Raman spectral mapping; SEM

**Introduction**

*Silybum marianum* L. Gaertn (Asteraceae) is an annual to biennial herb originates from mountains of the Mediterranean and North African regions where it forms scrub on a rocky base [1]. Currently it grows wild throughout Europe, North Africa, the Americas, and Australia [2,3]. It has several common names such as milk thistle, blessed milk thistle, Marian thistle, Mary thistle, Saint Mary’s thistle, Mediterranean milk thistle, variegated thistle and Scotch thistle. The flower heads of the plant are mostly red Purple (some cultivars have white flower) the flower head-bud is about 4 cm long and 12 cm wide. *S. marianum* plants flower from June to August in the North or December to February in the Southern Hemisphere. *S. marianum* fruits vary from white to nearly black. The plant is cultivated on a large scale in Austria, Germany, Hungary, Poland, China and Argentina for the production of raw material for the pharmaceutical industry. The fruits of *S. marianum* are one of the most important herbal liver medicines. The plant is the most studied medicinal plant in the treatment of liver disease. Fruits of *S. marianum* have been used for more than 2000 years to treat a range of liver and gallbladder disorders, including hepatitis, cirrhosis, and jaundice, and to protect the liver against poisoning from chemical and environmental toxins, including snake bites, insect stings and mushroom poisoning. Extracts of *S. marianum* fruits have been recognized for centuries as “liver tonics” and are well known to prevent or reverse hepatotoxicity of reactive drug metabolites or naturally occurring toxins [4]. The plant is sometimes also used as a decorative element in gardens, and its dried flower heads may be used for the decoration of dry bouquets [5]. The fruits are harvested in July-August after blooming. The medicinal part of the plant is the fruits often referred to as “seeds”. Silymarin is the major bioactive constituent of the fruit extract of *Silybum marianum*, consisting of a mixture of one flavanonol (taxifolin) and six flavonolignans (silychristin, silydianin, silybin A, silybin B, isosilybin A and isosilybin B) [6-9], however, a very recent investigation by Hohmann group revealed the presence of up to 23 flavolignan components [10]. The fruit is also known to be rich in lipids with up to 25% weight/weight [11] which interfere with the extraction of silymarin and has to be removed from the fruits (defatting) prior to silymarin extraction using nonpolar organic solvents. Little is known about the location of silymarin in the fruit of *Silybum marianum*, but most of them relayed on using microscopic histology that were lacking structural information or spectral data [12], or were based on mechanical separation of fruit parts and extracting them with appropriate solvent which still did not show exactly the precise location of silymarin in the fruit [13-14].

The objective of this study was to use microscopic spectroscopy for mapping the distribution of silymarin in the fruit of *Silybum marianum* for the purpose of improved efficiency of extracting silymarin.

**Experimental Section**

**Fruits of *Silybum marianum***

Certified *Silybum marianum* fruit were purchased from Frontier Natural Products CO-OP (2012 Frontier, Norway, IA 52318 USA) patch # 912.3128. Seeds were washed with deionized water, ultrasonicated, and air dried prior to investigation.

**Stereomicroscopy**

Whole fruits were first observed using stereomicroscope for morphology, shape, size, and color parameters. Free hand longitudinal sections were obtained from the whole fruits and observed using Nikon's SMZ25 Fluorescence Stereomicroscope with EGFP Bandpass filter having excitation at 450-490 nm and emission at 500-550 nm (Nikon Instruments Inc., Melville, NY, USA) equipped with NIS Elements High Content Analysis Microscope Imaging Software.

**Scanning Electron Microscopy (SEM)**

Sections of dried fruits were hand-made using a razor blade, mounted on aluminum stubs using double-sided adhesive carbon tape
and were sputtered with a thin layer (20 Å) of gold using Desk V HP Dantôn Vacuum system (Dantôn Vacuum LLC, Mooresestown, NJ, USA) under argon gas. Coated fruit sections were examined and imaged by the scanning electron microscopy Evex Global Nano SEM at an accelerated voltage of 20 KV.

**Infrared Spectroscopy**

FT Raman and FT infrared spectra were acquired using Nicolet IS50 (Thermo Scientific). IR was acquired using IS50 ATR for solid samples.

**High Performance Thin Layer Chromatography (HPTLC)**

Methanolic extracts of the fruit shell and the embryo cotyledons surrounded with the endosperm layer were examined by HPTLC along with standard silymarin solution on silica gel plates in 10 mm bands at 40 μg/band (HPTLC Silica Gel 60F254 20 × 10 Merck K GaA 6427 Darmstadt, Germany). Plate was developed with chloroform, acetone and formic acid 75:16.5:8.5 v/v, dried and observed at a UV of 254 and 366 nm. Plates were also derivatives in 1% vanillin reagent, (1 g of Vanillin, 100 mL of ethanol mixed with 5 mL of conc. Sulphuric acid and 95 mL of ethanol), and the plate was dried in the oven at 100°C for 5 min, after which spots were visualized under white light and 366 nm wavelengths. All applications including spotting, development, derivatization and imaging were performed using the Camag automated instruments (Automatic TLC Sampler 4, Rep Pro Star 3 and ADC 2 Automatic Developer) and win Cats-Planar Chromatography manager software (CAMAG USA, Denver, CO).

**Accurate Mass Spectrometry**

Accurate Mass were measured using LC-QTOF Agilent 6530 system equipped with 1290 binary pump, Phenomenex Kinetex C18 100 mm × 3.0 mm 2.6 μm column. Solvents A was made of methanol with 0.1% formic acid; B was 0.1% formic acid in water run in a gradient protocol starting at 30% A to 60% A over 15 min with a flow 0.2 mL/min. Ion source ESI+Agilent Jet Stream Technology in positive ionization mode. The Jet Stream ESI source was operated in positive/negative mode, and instrument parameters were set as follows: sheath gas temperature, 350°C; sheath gas flow, 8 L/min; nebulizer, 20 psi; dry gas temperature, 300°C; dry gas flow, 5 L/min; and capillary entrance Voltage, 3500 V. Fragmentor and Skimmer were operated at 190 and 65 V, respectively. The MS data were collected with Mass Hunter Data Acquisition B.06.00 (Agilent Technologies), and Mass Hunter Qualitative Analysis B.06.00 (Agilent Technologies) was applied to identify silymarin components. All EICs were obtained with ±10 ppm m/z expansion.

**Results and Discussion**

Fruits of *Silybum marianum* are small elongated ovate and slightly compressed dorsoventrally. Fruits vary in color from light pale yellow to brown and dark reddish brown with shiny and shiny surface, the exocarp (pericarp epidermis) consisting of elongated polygonal cells. Average dimensions of the fruits are 6.8 mm in length and 3.0 mm in width with an average weight of 2.8 g/100 fruit (Figure 1).

Figure 1: Fruits of *Silybum marianum*.

Scanning Electron Microscopy (SEM) examination of the fruit sections revealed that *S. marianum* consist of the pericarp, testa epidermis, endosperm and the embryo. The pericarp is subdivided into exocarp, mesocarp and endocarp. Exocarp is the outermost layer (pericarp epidermis), mesocarp which makes the pericarp parenchyma, and these cells contain dark tannins responsible for the spotted appearance of the fruits. The sclerenchyma cells of the endocarp contain large calcium oxalate prismatic crystals [9]. Testa epidermis layer consisted of elongated macroscleleids and followed by the subepidermal tissue. The endosperm forms a single layer of cells that surrounds the embryo’s two cotyledons with many elongated cells filled with fat droplets (Figure 2).

Examination of the fruit sections using fluorescent stereomicroscope using Nikon’s SMZ25 Fluorescence Stereomicroscope revealed that silymarin is located in the pericarp layer of the fruit. Certified silymarin standard was found to give a strong green fluorescent at 500-550 nm when exited at 450-490 nm using the EGFP Bandpass filter; fats and proteins did not show any emission using that filter. Fluorescent mapping of the fruit longitudinal section is shown in Figure 3. Fluorescence emission under the stereomicroscope produced a diffused image which is difficult to locate the exact locations of the silymarin as appeared as continues green...
band in the pericarp, the apical region of the fruit, and to lesser extent at the testa epidermal layer.

**Figure 3:** Stereomicroscope micrographs of *Silybum marianum* longitudinal sections: A and B no fluorescent applied; C and D epifluorescence showing the distribution of silymarin in the pericarp.

Raman spectra mapping showed precisely that silymarin is only located in the layer of the pericarp (endocarp) that included the crystals, and also in the apical region. It appears as none continues band that reflects the distribution of the crystals within the endocarp tissues. The red color indicated the highest concentrations of the silymarin and the blue color indicates the lowest as shown in Figure 4.

**Figure 4:** Raman spectra mapping of silymarin in *S. marianum* section: A) outside the fruit surface; B) pericarp; C) embryo.

To confirm our results, the fruits were mechanically separated as shell layer and endosperm/embryo layers and each layer was extracted with methanol to examine their contents of silymarin. The two extracts were examined first by HPTLC and revealed that the endosperm/embryo layer did not have any silymarin and was mostly consisted of proteins and lipids. On the other hand, the shell layer was very rich in silymarin as shown in Figure 5. Yield of silymarin in the shell was 6% compared to 2% in the whole fruits.

**Figure 5:** HPTLC of *Silybum marianum* extracts: 1) endosperm/embryo; 2) fruit shell; 3) standard silymarin.

FTIR of the solid materials of the shell and the other inside layers of the fruits (endosperm and embryo) showed that IR spectra confirmed that silymarin is located only in the shell of the fruits. The chromone carbonyl group (1660 cm\(^{-1}\)), the aromatic ether linkage group (1160 cm\(^{-1}\)), the phenolic hydroxyl group (3500 cm\(^{-1}\)), the keto group (1745 cm\(^{-1}\)) and the lactone ring group (1752 cm\(^{-1}\)) characteristic absorption for silymarin were only detected in the shell of the fruits. Raman bands found in the shell of the fruits but not in the endosperm layers were the symmetric aromatic ring stretching at 1605 cm\(^{-1}\), the C=C ring stretching at 1649 cm\(^{-1}\), and the asymmetric ring vibration at 1500 cm\(^{-1}\).

Further confirmation was carried out by examining the methanol extracts of the shell and the inner layers of the fruits using accurate mass QTOF analysis showing again that silymarin components exist only in the shell as presented in Table 1.

<table>
<thead>
<tr>
<th>Accurate Mass</th>
<th>Taxifolin</th>
<th>Silychristin</th>
<th>Silydianin</th>
<th>Silybin A</th>
<th>Silybin B</th>
<th>Isosilybin A</th>
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**Table 1:** Accurate mass data of silymarin individual compounds.
Conclusion

Micro spectroscopy using fluorescence microscopy and Raman microscopy of the fruit sections showed that silymarin is located exclusively in the pericarp layer or the fruit shell. Extraction of silymarin from the mechanical separated fruit sections into its inner part (endosperm and embryo) and the seed shell (pericarp) with methanol using microwave assisted extraction revealed that silymarin is mostly located in the pericarp with a recovered yield of 6%. No silymarin was recovered from the endosperm/embryo layer. The obtained results suggested that using the pericarp is more efficient in extracting silymarin than using the whole fruit. It also reduces consumption of solvents necessary for defatting of the fruit while the endosperm/embryo part, which is rich in fatty oil, can be used for production of biodiesel fuel.

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References