Application of High-Speed Countercurrent Chromatography to the Separation of Flavanonol, Phenylcoumaran and Flavonolignans in *Silybum marianum*

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Abstract
Separation of flavanonol, phenylcoumaran and flavonolignans in *Silybum marianum* was examined using high-speed countercurrent chromatography (HSCCC). In order to prepare analytical standards, a flavanonol of aromadendrin (87.4% purity, 7.8 mg), three phenylcoumarans of jatrointelignan D (84.5% purity, 13.2 mg), dehydrodiconiferyl alcohol (76.1% purity, 1.4 mg) and dihydrodehydrodiconiferyl alcohol (93.0% purity, 5.8 mg), and three flavonolignans of silybin (88.8% purity, 35.6 mg), silydianin (99.3% purity, 25.1 mg) and silychristin (96.9% purity, 12.6 mg) were separated from the seeds of *S. marianum* using common column chromatography and ODS-HPLC, and identified by 1H and 13C NMR spectra. Then, HSCCC with the hexane/ethyl acetate/methanol/water (3 : 7 : 4 : 6, v/v) system was applied to the separation of aromadendrin, jatrointelignan D, silydianin and silybin. In this separation, it was revealed that silybin and silydianin were successfully separated from each other. The present HSCCC system was directly applied to the ethyl acetate extract and resulted in the separation of silybin. The overall results suggested that HSCCC is useful for the separation of bioactive compounds in *S. marianum*.

Keywords: Separation; High-speed countercurrent chromatography; *Silybum marianum*; Natural products

1. Introduction
The genus *Silybum* is a member of the Asteraceae family and *Silybum* spp. are widely distributed in Mediterranean region, North Africa, Asia and also in Japan. The crude extract of *S. marianum* is used for apezpy and the further purified product is used for chronic hepatitis and cirrhosis of liver in Germany. It is known that *S. marianum* contains many flavonolignans, sterols and flavonoids [1-4]. In particular, silymarin is contained at about 30 – 65% in the seeds of *S. marianum* and it is composed of silybin, isosilybin A, isosilybin B, silychristin, isosilychristin, silydianin and taxifolin [5-7]. Silymarin has antioxidant activity, antifibrotic activity, hepatic protection effect, etc., [5,7,10] where silybin covers approximately 50 – 70% of silymarin with other components including isosilybin (about 5%), silychristin (about 20%) and silydianin (about 10%) as well as silimonin, isosilychristin, isosilibinin, etc [7]. Several biological activities of silymarin as described above may be caused by silybin. From this reason, simple separation of the bioactive compounds in *S. marianum* is required to obtain at the preparative level without loss of samples.

Countercurrent chromatography (CCC) is a support-free chromatography based on the liquid-liquid partitioning of analytes between two immiscible liquid phases in the column. This distinctive feature eliminates the adsorption and the denaturation of the target compound caused by the interaction between the solutes and the solid stationary phase which is sometimes observed in the use of common column chromatography. CCC has been widely used for...
the separation and the purification of various bioactive compounds in the natural products [11-13]. Among many CCC instruments developed in the past, the high-speed CCC (HSCCC) is useful for effective separation of natural compounds within shorter separation times.

This paper describes the application of HSCCC to the separation of flavanol, phenylcoumaran and flavonolignans in *S. marianum*.

2. Experimental

2.1. Apparatus

The HSCCC apparatus employed in the present study was constructed at the Machining Technology Center of Nihon University, Chiba, Japan. The design and the fabrication of the apparatus have been described in elsewhere [14,15]. For each separation, four eccentric coil assemblies were equipped in rotary frame of the HSCCC apparatus. Each eccentric coil assembly was prepared by winding 1 mm I.D. PTFE (polytetrafluoroethylene) tubing (Flon Kogyo Co., Tokyo, Japan) onto 5 cm long, 5 mm O.D. nylon pipes making a series of tight left-handed coils for the forward type-J planetary motion or right-handed coils for the backward type-I planetary motion (20 turns for 1 unit). These coil units were arranged symmetrically around the holder hub of 5 cm O.D. in such a way that the axis of each coil unit is parallel to the holder axis (13 units for the first layer and 20 units for the second layer). The total capacity of four columns connected in series was 55 mL.

The HPLC equipment consists of a reciprocating pump (Type LC-20AD, Shimadzu Corporation, Kyoto, Japan), a UV detector (Type SPD-10AVP, Shimadzu), a separation column (Inertsil ODS-4, 250 × 4.6 mm I.D., 5 μm, GL Sciences, Tokyo, Japan for the analytical scale, Inertsil ODS-4, 250 × 14 mm I.D., 5 μm, GL Sciences and COSMOSIL Cholesterol, 250 × 10 mm I.D., 5 μm, Nacalai tesque, Kyoto, Japan for the preparative scale). 

1H, 13C and 2D NMR spectra were measured using a spectrometer (Type ECA-500, JEOL, Tokyo, Japan) and the chemical shifts were referenced to the residual solvent peaks of methanol-d4 (CD3OD) or acetone-d6 (((CD3)2)CO) as an internal standard (δH 3.31 and δC 49.0 for CD3OD, or δH 2.05 and δC 29.84 for ((CD3)2)CO).

2.2. Reagents and plant material

Hexane, ethyl acetate (EtOAc), methanol (MeOH) and 1-butanol were of the first grade purchased from FUJIFILM Wako Pure Chemical Corporation (Osaka, Japan). All other reagents were of reagent grade. The seeds of *S. marianum* were obtained from a market at Hebei in China.

2.3. Separation and identification of flavanol, phenylcoumarans and flavonolignans in *S. marianum*

The seeds of *S. marianum* (1.5 kg) were extracted with MeOH, and a part of the extract (36.8 g) was dissolved with water and then partitioned with hexane. The aqueous layer was further partitioned with EtOAc to give the extract (25.6 g). A part of this extract (3.6 g) was subjected to a silica gel column (chloroform (CHCl3)/MeOH, 1 : 0 → 50 : 1 → 19 : 1 → 9 : 1 → 8 : 2 → 1 : 1 → 0 : 1) to give 13 fractions. The fraction eluted with CHCl3/MeOH (50 : 1) was subjected to passage over an ODS column (water/MeOH, 6 : 4 → 1 : 1 → 4 : 6 → 3 : 7 → 2 : 8 → 0 : 1) to give jatrointelignan D [16] and silybin [2]. The fraction eluted with CHCl3/MeOH (9 : 1) was subjected to separate over an ODS column (water/MeOH, 8 : 2 → 6 : 4 → 1 : 1 → 4 : 6 → 2 : 8 → 0 : 1) followed by ODS-HPLC (acetonitrile/0.1% formic acid, 3 : 7) using COSMOSIL Cholesterol column to give dehydrodiconiferyl alcohol [17,18], dihydrodehydrodiconiferyl alcohol [18] and aromadendrin [19]. Other fractions eluted with water/MeOH (1 : 1) were further subjected to an ODS HPLC using Inertsil ODS-4 column and a silica gel column (benzene/EtOAc, 1 : 0 → 8 : 2 → 7 : 3 → 6 : 4 → 1 : 1) to give silydianin [2]. The fraction eluted with CHCl3/MeOH (1 : 1) was identified as silychristin [2].

2.4. Measurement of the partition coefficient of analytical standards in the hexane/EtOAc/MeOH/water (HEMWat) system

In order to select a suitable two-phase solvent system for the HSCCC separation, the partition coefficient (KD) values of jatrointelignan D, silybin, aromadendrin and silydianin were measured using the simple test tube method using the hexane/EtOAc/MeOH/water (HEMWat) system established by Friessen et al. [20] as follows: Two milliliters of each phase were put into a test tube, and then about 1 mg of test sample was added and mixed vigorously. After two layers were formed, 1 mL of each phase was put into another test tube and diluted with 2 mL of MeOH. The absorbance of this solution was measured at 288 nm using a spectrophotometer (Type UV-1800, Shimadzu). The wavelength at 288 nm selected in the present study can commonly detect flavanols, phenylcoumarans and flavonolignans. The KD value was calculated from the absorbance of the upper phase divided by that of the lower phase.

2.5. Preparation of the two-phase solvent system and the sample solution

Each HEMWat system examined in the present study was prepared at a desired volume ratio as summarized in Table 3. The solvent mixture was stirred vigorously and then settled to form two layers in a separatory funnel at room temperature. The sample solution was prepared by dissolving an aliquot of the sample in equal volume of each phase for HSCCC separation.
2.6. HSCCC separation and HPLC analysis of the fraction
The coiled column was filled with the stationary phase and followed the injection of the sample solution through the column inlet by the syringe. When the rotation speed of the column reached 1000 rpm, the mobile phase was pumped into the column at a flow rate of 0.5 mL/min. The effluent was collected into test tubes at 2 min/tube using the fraction collector (Advantec. Co., Tokyo, Japan) and measured the absorbance at 288 nm, and then applied to the HPLC analysis. The HPLC conditions are as follows: column: Inertsil ODS-4 (250 × 4.6 mm I.D.), eluent: 0.1% acetic acid (A), acetonitrile (B), A/B = 75 : 25 (initial) – 75 : 25 (15 min) – 60 : 40 (10 min) – 60 : 40 (10 min), flow rate: 1.0 mL/min, column temperature: 40 °C, detection: 288 nm.

3. Results and discussion
3.1. Separation and identification of flavanonol, three phylcoumarans and three flavonolignans
In order to prepare analytical standards, flavanonol, three phylcoumarans and three flavonolignans were separated from the seeds of S. marianum and identified by 1H and 13C NMR spectra. As a result, aromadendrin (87.4% purity, 7.8 mg), jatrointelignan D (84.5% purity, 13.2 mg), dehydrodiconiferyl alcohol (76.1% purity, 1.4 mg), dihydrodehydrodiconiferyl alcohol (93.0% purity, 5.8 mg), silybin (88.8% purity, 35.6 mg), silydianin (99.3% purity, 7.8 mg), jatrointelignan D (84.5% purity, 13.2 mg), aromadendrin and silybin obtained using a simple test tube method for jatrointelignan D, silybin, aromadendrin and silydianin, which were obtained at relatively large amounts. Table 1 summarizes the K$_D$ values of these four compounds were increased according to the increase of the polarity of the two-phase solvent system (the increase of the number of the HEMWat system). In the most hydrophilic composition ratio of the present solvent system (HEMWat −8), all four compounds were almost partitioned into the lower phase, while in the most hydrophilic composition ratio of the present solvent system (HEMWat +8), these compounds were almost partitioned into the upper phase. As the result, the HEMWat +4 system, hexane/EtOAc/MeOH/water (3 : 7 : 4 : 6, v/v) was selected for the HSCCC separation because the K$_D$ values of these compounds were relatively different from each other.

3.3. HSCCC separation of components obtained from the seeds of S. marianum
Figure 2 illustrates the HSCCC separation of silydianin, jatrointelignan D, aromadendrin and silybin obtained using the HEMWat system (3 : 7 : 4 : 6, v/v) with the lower mobile phase. The stationary phase retention was 14.5%, which was extremely low but almost similar elution pattern was obtained using another type of the HSCCC, the small-scale cross-axis CCC, with the stationary phase retention at 41.2%.

As illustrated in Fig. 2, the eluate obtained in this HSCCC separation was divided into four fractions (Fr. a – d). The purity of the component containing each fraction was calculated from the HPLC data. Figure 3 shows the HPLC chromatograms of each fraction. In the HSCCC, each compound was separated insufficiently except for silydianin in Fr. b. Fraction c contained both jatrointelignan D and aromadendrin as illustrated in Fig. 3C. However, in Fr. d, silybin was detected as highly purified single peak in the HPLC chromatogram. From these results, it was suggested that silydianin and silybin could be separated in the present HSCCC conditions. When using the HEMWat +5 system, hexane/EtOAc/MeOH/water (3 : 7 : 3 : 7, v/v), silydianin and jatrointelignan D may be separated with the lower mobile phase, and silybin and aromadendrin may be separated with the upper mobile phase.
phase. HSCCC is useful for the preparation of bioactive compounds, because the single HSCCC peak with UV detection is sometimes composed of several constituents at high purities, which is different from HPLC.

Table 1. Composition ratios of the HEMWat system and the partition coefficient ($K_D$) values of jatrointelignan D (1), silybin (2), aromadendrin (5) and silydianin (6).

| HEMWat hexane | EtOAc | MeOH | water | $K_D$ 1 | $K_D$ 2 | $K_D$ 5 | $K_D$ 6 |
|---------------|-------|------|-------|---------|---------|---------|---------|
| -8            | 10    | 0    | 10    | 0.54    | 0.11    | 0.04    | 0.22    |
| -7            | 9     | 1    | 9     | 0.49    | 0.10    | 0.08    | 0.04    |
| -6            | 8     | 2    | 8     | 0.44    | 0.03    | 0.01    | 0.01    |
| -5            | 7     | 3    | 7     | 0.43    | 0.25    | 0.02    | 0.06    |
| -4            | 7     | 3    | 6     | 0.40    | 0.01    | 0.09    | 0.04    |
| -3            | 6     | 4    | 6     | 0.42    | 0.28    | 0.05    | 0.05    |
| -2            | 7     | 3    | 5     | 0.27    | 0.11    | 0.02    | 0.08    |
| -1            | 6     | 4    | 5     | 0.37    | 0.08    | 0.09    | 0.15    |
| 0             | 5     | 5    | 5     | 0.49    | 0.11    | 0.10    | 0.07    |
| +1            | 4     | 6    | 5     | 0.57    | 0.27    | 0.29    | 0.09    |
| +2            | 3     | 7    | 5     | 0.66    | 0.56    | 0.59    | 0.15    |
| +3            | 4     | 6    | 4     | 0.79    | 0.90    | 0.66    | 0.20    |
| +4            | 3     | 7    | 4     | 1.07    | 2.50    | 1.59    | 0.36    |
| +5            | 3     | 7    | 3     | 1.76    | 6.63    | 3.47    | 0.48    |
| +6            | 2     | 8    | 2     | 3.40    | 13.14   | 11.13   | 0.96    |
| +7            | 1     | 9    | 1     | 6.00    | 14.97   | 16.53   | 1.50    |
| +8            | 0     | 10   | 0     | 24.19   | 43.49   | 24.17   | 2.01    |

Figure 2. The elution curve of selected compounds isolated from *S. marianum* using HSCCC. Experimental conditions: Apparatus: the universal HSCCC equipped with eccentric coil assemblies, 1.5 mm I.D. × 2.5 mm O.D., 55 mL capacity; Sample: jatrointelignan D (1 mg), silybin (1 mg), aromadendrin (1 mg) and silydianin (1 mg) for lower mobile phase; Solvent system: HEMWat (3:7:4:6); Rotation speed: 1000 rpm; Flow rate: 0.5 mL/min; Fractionation: 1.0 mL/tube; Detection: 288 nm; SF: solvent front.

Figure 3 illustrates the HSCCC separation of the EtOAc extract obtained from the seeds of *S. marianum* using the HEMWat (3 : 7 : 4 : 6, v/v) system with the lower mobile phase. The eluate was divided into 10 fractions. As illustrated in Fig. 5, silybin was successfully separated in Fr. 9 at high purity while the elution time was slightly shorter than that in Fig. 3. This may be caused by lower stationary phase retention at 12.7% due to the matrix effect of the sample. However, this result suggests that silybin can be directly separated from the EtOAc extract without any purification process.

Fig. 3. HPLC chromatograms of standards (red or green) and each fraction (blue) after HSCCC separation. (A) Fraction a, (B) Fraction b and silydianin (6), (C) Fraction c, jatrointelignan D (1) and aromadendrin (5), (D) Fraction d and silybin (2). Experimental conditions: Column: Inertsil ODS-4 (4.6 × 250 mm, 5 µm); Mobile phase: A/B 0.1% formic acid/acetonitrile = 75 : 25 (initial) – 75 : 25 (15 min) – 60 : 40 (10 min) – 60 : 40 (10 min); Flow rate: 1.0 mL/min; Column temperature: 40 °C; Detection: 288 nm.
Fig. 4. The elution curve of EtOAc layer extracted from the seeds of S. marianum using established HSCCC separation method, and fraction number and elution time of each fraction after HSCCC separation. Experimental conditions: Apparatus: the universal HSCCC equipped with eccentric coil assemblies, 1.5 mm I.D. × 2.5 mm O.D., 55 mL capacity; Sample: EtOAc layer extracted from the seeds of S. marianum for lower mobile phase; Solvent system: HEMWat (3 : 7 : 4 : 6); Rotation speed: 1000 rpm; Flow rate: 0.5 mL/min; Fractionation: 1.0 mL/tube, Detection: 288 nm; SF: solvent front.

Fig. 5. HPLC chromatograms of standards and fraction 9 after HSCCC separation of the EtOAc extract from the seeds of S. marianum. Experimental conditions: Column: Inertsil ODS-4 (4.6 × 250 mm, 5 µm); Mobile phase: A/B 0.1% formic acid/acetonitrile = 75 : 25 (initial) – 75 : 25 (15 min) – 60 : 40 (10 min) – 60 : 40 (10 min); Flow rate: 1.0 mL/min; Column temperature: 40 °C; Detection: 288 nm.

4. Conclusions
The HSCCC was applied to the separation of flavanonol, phenylcoumaran and flavonolignans obtained from the seeds of S. marianum. As the result, it was revealed that silydianin and silybin was separated using the HEMWat system (3 : 7 : 4 : 6, v/v) with the lower mobile phase. Silybin was also directly separated from the EtOAc extract at high purity. The present HSCCC is useful for the simple separation of silybin because it is known as a bioactive compound.

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Conflicts of Interest
The authors declare no conflict of interest.

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