

## Isolation of Kaempferol from Vietnamese *Ginkgo biloba* Leaves by Preparative Column Chromatography

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The aim of this research was to develop a simple procedure to isolate kaempferol from *Ginkgo biloba* leaves extract. Flavonoids present in *Ginkgo biloba* leaves were extracted by 96 % ethanol. These flavonol glycosides which were hydrolyzed in the presence of HCl to convert to aglycones. The flavonol aglycones were extracted by the hydrolysis of solution with ethyl acetate which was evaporated to dryness under reduced pressure to obtain concentrated flavonol aglycones (C2). Flavonol aglycones were then purified by preparative column chromatography with a mobile phase composed of petroleum ether-ethyl acetate to acquire fraction C3. The recrystallization of fraction C3 in acetone realizes to have an amorphous sediment as fraction C4 that was supposed to be a mixture of kaempferol, quercetin and isorhamnetin. Kaempferol in this sediment was purified by column chromatography with a mobile phase composed of CHCl<sub>3</sub>-MeOH to achieve a fraction denoted as fraction C5. The isolated fraction C5 was assigned by its purity and structure by thin layer chromatography (TLC), high performance liquid chromatography (HPLC) and spectroscopic data analyses.

**Keywords:** *Ginkgo* leaves extract, Kaempferol, Silica gel, Column chromatography.

### INTRODUCTION

*Ginkgo biloba*, commonly known as ginkgo or ginkgo, is among the most oldest plants (a living fossil). *Ginkgo biloba* is the only remaining species of ancient Ginkgoales remaining. *Ginkgo biloba* grows widely in China, spread to Japan, Europe, North America and elsewhere [1]. Crucial constituents contained in this medicinal leaves are the terpene trilactones, flavonoids, alkylphenols, simple phenolic acids, 6-hydroxykynurenic acid, 4-O-methylpyridoxine and polyprenols [2].

Ginkgo flavonoids have been recommended for cerebral-related symptoms such as deterioration functional disturbances, vascular blockage. These flavonoids possess antitumor and anti-inflammatory activities [3]. Crude extracts of ginkgo leaves in concentration of 7.8 µg/mL inhibit activity of both Gram-positive bacteria (*Klebsiella pneumoniae*, *Pseudomonas aeruginosa*) and Gram-negative bacteria (*Staphylococcus aureus*, *Staphylococcus epidermidis* and *Streptococcus pyogenes*). Methanol fractions of leaves significantly restrain tumor forming [4,5]. Ginkgo flavonoids are able to diminish the lipid agglome-

ration in relation to up-regulation of carnitine palmitoyl-transferase 1A (CPT1A), a rate-limiting enzyme, in the β-oxidation of long-chain fatty acids [6]. Totally 13 isolated flavonoids from *Ginkgo biloba* were found to inhibit AChE dose dependently at IC<sub>50</sub> from 57.8 to 133.1 µg/mL.

The flavonoids found in the leaves of *Ginkgo biloba* could be analyzed qualitatively and quantitatively by RP-HPLC. After being hydrolyzed in acid to aglycones (quercetin, kaempferol and isorhamnetin) and separated by HPLC, original total content of these glycosids is estimated from that of aglycones [2,7]. Kaempferol which is among principal yellow colouring material is crystalline solid. It is soluble in dimethyl sulfoxide, ethanol and ether. The extraction solvents of kaempferol mainly are methanol, ethanol, acetone and water with the assistance of ultrasound, microwaves, ultrasound and microwaves [8]. In relation to advanced technology of extraction, soxhlet extraction technique have been employed to extract kaempferol with 80 % methanol as a solvent [9]. Extraction using supercritical fluids (usually CO<sub>2</sub>) is an alternative method whose conditions are at lower temperature and higher pressure. This brings about

high extraction performance since substances are not decomposed but requires expensive equipment [8]. The extract is then hydrolyzed with HCl in methanol and partitioned continuously with chloroform to eliminate non-polar contaminants and ethyl acetate. The ethyl acetate fractions were collected and separated by column chromatography with 50 % methanol as an eluant [10]. Another way to purify ethyl acetate soluble fraction is further partition with *n*-hexane. The *n*-hexane fraction was then subjected respectively to vacuum liquid chromatography and reverse-phase column chromatography using Sephadex column LH-20 to achieved highly pure kaempferol [11].

The one step separation technique to isolate kaempferol from a crude ethanol extract of *Ginkgo biloba* leaves could be high-speed counter-current chromatography method. The separation was carried out by applying a mobile phase constituted of *n*-hexane-butanol-ethyl acetate-methanol-0.5 % acetic acid (1:0.5:3.5:1:4, v/v) [3].

The polarity of three flavonol aglycones is increased from kaempferol, quercetin and isorhamnetin due to the presence of hydroxyl groups in their structure. Therefore, they could be isolated from each other by preparative column chromatography with normal stationary phase which is suitable for relatively less polar substances. This technique is among the most simple and therefore available for larger scale preparative applications. Solvents for elution is manually or automatically applied. Kaempferol purities were determined by HPLC and structurally identified by IR, MS, <sup>1</sup>H and <sup>13</sup>C NMR.

## EXPERIMENTAL

**Plant materials:** *Ginkgo biloba* leaves were purchased from local market store of Ho Chi Minh City, Vietnam during March 2018. The sample was identified by the Department of Botany, Pharmacy Faculty, Nguyen Tat Thanh University. After collection, the leaves were dried under 60 °C for 72 h in an oven with force ventilation and then ground into raw powder.

Methanol and acetonitrile (Merck Darmstadt, Germany) were of HPLC quality. Acetone, ethyl acetate, *n*-hexane, acetic acid were of synthetic quality (Chemsol, HCM city, Vietnam). All the solvents used for HPLC analysis were filtered (0.45 µm) and ultrasonically degassed before use. All the separations were carried out at room temperature (25 °C). The chromatography column (35 cm × 2 cm) was packed with 25 g of silica gel 60 Merck (0.040-0.063 mm).

**Flavonoid extractions and hydrolysis:** About 1.5 kg of *Ginkgo biloba* leaves powder were extracted by repeated cold maceration in 2 L of 96 % EtOH at room temperature for one time a day. The same extraction process was repeated for 4 times in the same leave sample, using totally 8 L absolute ethanol. The extract was filtered and concentrated in vacuum at 60 °C to dryness with a rotary evaporator and denoted as C1. The extract C1 is then distributed into 1 L hot water then added chloroform, shaken and allowed to stand 30 min in order to eliminate non-polar impurities. Each 100 mL of distributed water was shaken with 30 mL CHCl<sub>3</sub> until the CHCl<sub>3</sub> layer exhibited no colour. The aqueous layer after shaking with CHCl<sub>3</sub> was separated and hydrolyzed with 10 % HCl at pH 3-4. The acidic aqueous solution was shaken with EtOAc. Each time of shaking, used 30 mL EtOAc and acidic water layer

was collected. This process was repeated until EtOAc layer yellow colour disappeared. The acidic aqueous layer was hydrolyzed by reflux method with 20 % HCl at pH 1.3-1.4 for 4 h and then left for for 1 h. The hydrolyzed solution was fractionally extracted with EtOAc (30 mL) and the EtOAc layer was collected. This layer was concentrated in vacuum evaporator at 40 °C to dryness and denoted as C2.

**Column chromatography:** The C2 fraction was collected and then separated using column chromatography (CC) method with silica gel as stationary phase and petroleum ether and EtOAc as mobile phase in the ratio of 1:1. The stationary phase was packed with 25 g of silica gel using a wet method. Subfractions were collected every 15 mL with flow-rate of 2.0 mL per minute. Consequently, total 200 subfractions were obtained (Table-1).

TABLE-1  
SUBFRACTIONS OF THE C2

PE-EtOAc	Volume (mL)	Subfractions (15 mL)
100:0	400	Not collected
95:5	300	1-20
90:10	300	21-40
85:5	3000	41-200

Kaempferol in subtractions was verified by TLC (silica gel 60 F<sub>254</sub>), solvent system CHCl<sub>3</sub>-MeOH-HCOOH (95:5:1, v/v), monitoring under an UV lamp. The subfractions from 1 - 200 were collected and concentrated in vacuum evaporator at 40 °C to dryness and denoted as fraction C3.

**Recrystallization in acetone:** The fraction C3 was dissolved in a flask using a minimal amount of acetone. The crude product was eluted out as a pure crystalline solid and leaving mother liquor. Kaempferol with higher purity (C4) was achieved after several times recrystallization. The fraction C4 was verified by the same chromatographic condition and revealed that it consist of three flavonoids.

**Final purification by column chromatography:** The fraction C4 was purified by column chromatography with CHCl<sub>3</sub> as a mobile phase at a flow-rate of 2.5 mL per minute. Eluant was not collected until chromatographically pure kaempferol begins to occur (monitored by TLC). The mobile phase was then changed from CHCl<sub>3</sub> to MeOH. Elution was performed until kaempferol was completely washed out the column. Evaporation of mobile phase at room temperature obtained a fraction C5. It was preserved from light using a dark membrane.

## RESULTS AND DISCUSSION

**Purity assessment by TLC:** Fraction C5 was analyzed with three different mobile phases with increasing polarity as (1) CHCl<sub>3</sub>-MeOH-HCOOH (8:2:1, v/v), (2) PE-EtOAc-HCOOH (5:5:1) and (3) CHCl<sub>3</sub>-EtOAc-HCOOH (5:4:1). The chromatogram of fraction C5 showed only one spot on a developed TLC plate. This indicated that the sample was a single component.

**Purity assessment by HPLC:** The purity of fraction C5 was also assessed by HPLC using the analytical conditions as detector: PDA2998, λ = 370 nm, column: Luna® 5 µm C18, (250 × 4.6 mm, 5 µm); mobile phase: MeOH-CH<sub>3</sub>CN-H<sub>3</sub>PO<sub>4</sub> 0.05 % (15:25: 60, v/v); volume of injection: 20 µL; Flow-rate: 1 mL/min and column temperature: 25 °C.

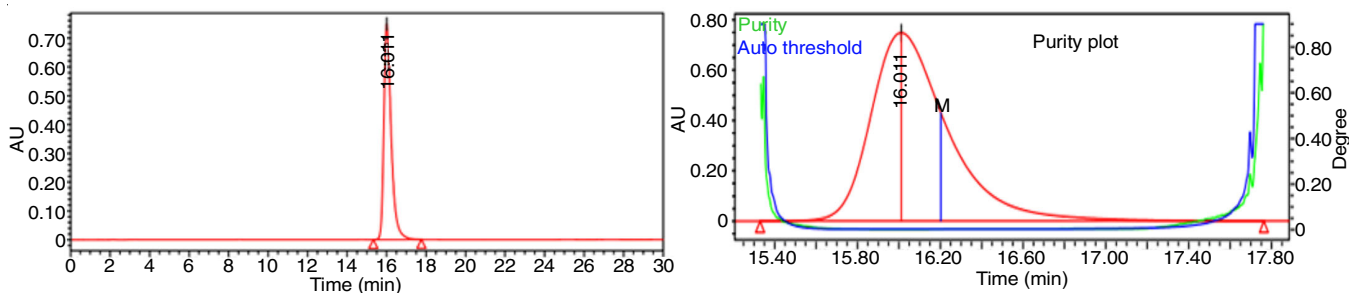


Fig. 1. HPLC chromatograph of fraction C5

Chromatographic purity of fraction C5 was 98 % as according to percentage of peak area over total areas of all peaks in the chromatogram (Fig. 1). In addition, only peak in chromatogram was proved to be not overlapped by any other peaks assessed by threshold value.

**FTIR analysis:** The infrared spectrum of fraction C5 showed broadened band in the region  $3211\text{ cm}^{-1}$  which indicates a hydroxyl group (-OH) and supported by the appearance of absorption at  $1090\text{ cm}^{-1}$  is a prime characteristic of C-OH. A strong absorption band formed at  $1655\text{ cm}^{-1}$  is due to C=O. Another strong bands at  $1613$  and  $1557\text{ cm}^{-1}$  were assigned to C=C aromatic. Similarly, a band at  $818\text{ cm}^{-1}$  was due to -CH bending of an aromatic ring while a weak peak at  $1166\text{ cm}^{-1}$  indicates the presence of C-O bond (Fig. 2) [10].

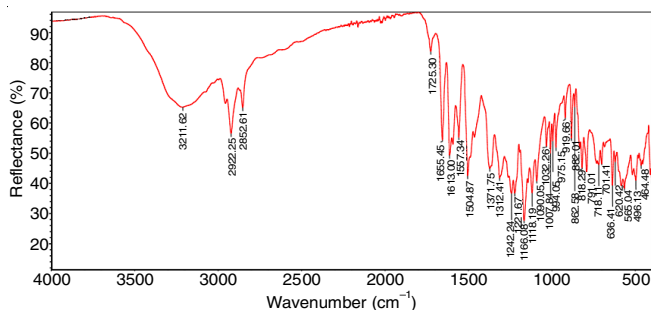


Fig. 2. Infrared spectra of fraction C5

**Mass analysis:** The mass spectrum in ESI mode of fraction C5 (Fig. 3) showed a quasi-molecular ion with  $m/z$  284.8. It indicates that molecular weight of fraction C5 is 286 as same as kaempferol [12].

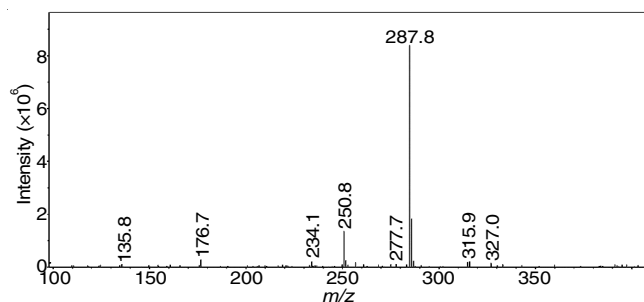


Fig. 3. Mass spectrum of fraction C5

**NMR analysis:**  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ ) spectrum was used to identify the number of proton and its multiple in the molecular structure of fraction C5. In these data, there are several groups of signals consisting of six protons aromatic

(Fig. 4). The emergence of doublet signals at 6.490 ppm (1H, d,  $J = 1.5$  Hz) and 6.229 ppm (1H, d,  $J = 1.5$  Hz) revealed an occurrence of two protons in coupling position of *m*-aromatic ring. The existence of two doublet signals at 8.021 ppm (2H, d,  $J = 9$  Hz) and at 6.943 ppm (2H, d,  $J = 8.5$  Hz) proved that *ortho*-coupling position on C-6',2' and C-5',3' of B-ring. The emergence of singlet signals at  $\delta$  10.251 and 12.438 ppm show the position of hydroxyl group. All these data (Table-2) revealed a model of flavonol skeleton.

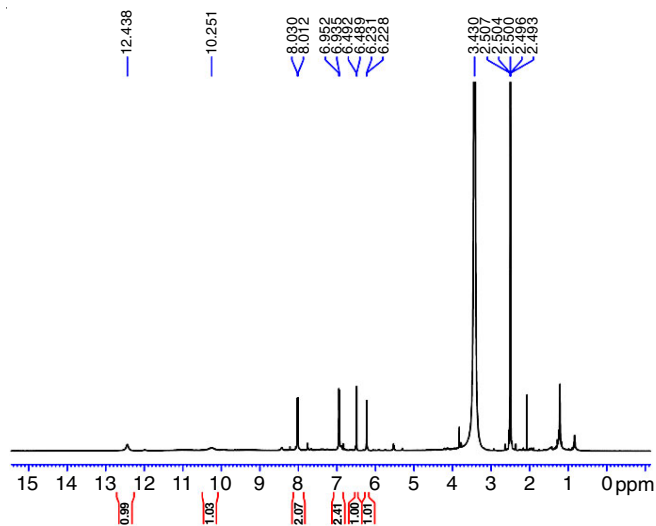


TABLE-3  
INTERPRETATION <sup>13</sup>C NMR SPECTRUM OF C5 IN  
COMPARISON WITH KAEMPFEROL (125 MHz, DMSO)

Position of C	C5 (ppm)	Kaempferol [13] (ppm)	Position of C	C5 (ppm)	Kaempferol [13] (ppm)
C-2	146.88	146.1 (s)	C-10	103.01	102.9 (s)
C-3	135.72	135.5 (s)	C-1'	121.66	121.6 (s)
C-4	175.97	175.7 (s)	C-2'	129.46	129.3 (s)
C-5	156.19	156.0 (s)	C-3'	115.53	115.3 (d)
C-6	98.34	98.2 (d)	C-4'	159.33	159.0 (s)
C-7	164.14	163.8 (s)	C-5'	115.53	115.3 (d)
C-8	93.58	93.4 (d)	C-6'	129.46	129.3 (d)
C-9	160.66	160.5 (s)	–	–	–

Based on spectroscopic data analysis, it can be concluded that the fraction C5 was found to be kaempferol (Fig. 5) having molecular formula C<sub>15</sub>H<sub>10</sub>O<sub>6</sub>.

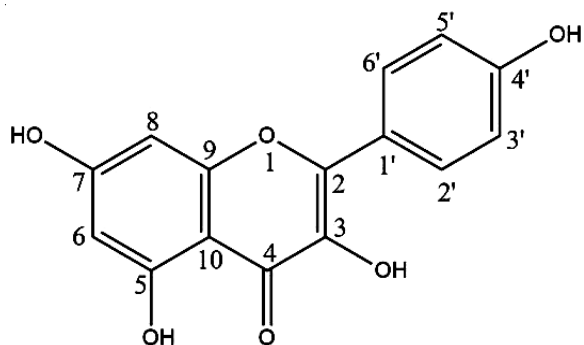


Fig. 5. Structure of kaempferol

## Conclusion

The isolation of flavonol glycoside (kaempferol) from *Ginkgo biloba* leaves extract using simple column chromatography was reported [14]. Flavonoids were first hydrolyzed from *Ginkgo biloba* leaves extract and then by converting them into aglycones. The resulting three aglycones were purified by column chromatography with normal phase silica gel. Isolated kaempferol confirmed by spectral (IR and NMR) and mass analysis. Isolated kaempferol showed a molecular ion peak at *m/z* 286 having molecular formula C<sub>15</sub>H<sub>10</sub>O<sub>6</sub>.

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## CONFLICT OF INTEREST

The authors declare that there is no conflict of interests regarding the publication of this article.

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