Identification of Antioxidative Constituents from Polygonum aviculare using LC-MS Coupled with DPPH Assay

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Abstract – A method for simultaneously identifying antioxidative compounds was developed using time-based LC-MS coupled with DPPH assay regardless of the time consuming process. The methanolic extract of Polygonum aviculare (Polygonaceae) showed significant DPPH radical scavenging activity. Time-based DPPH assay for simultaneous identification of active compounds from the extracts of P. aviculare was used. Major peaks of ethyl acetate fraction of P. aviculare showed high DPPH radical scavenging activity. A simple phenolic compound (1) and six flavonoids (2-7) were isolated from the ethyl acetate fraction of P. aviculare by silica gel and sephadex LH-20 column chromatography. The structures of seven compounds were determined to be protocatechuic acid (1), catechin (2), myricitrin (3), epicatechin-3-O-gallate (4), avicularin (5), quercitrin (6), and juglanin (7) based on the analysis of the ¹H-NMR, ¹³C-NMR and ESI-MS data. All compounds exhibited significant antioxidant activity on DPPH assay and active compounds were well correlated with predicted one.

Keywords – Polygonum aviculare, Flavonoids, LC-MS, DPPH assay

Introduction

Natural products which have used for thousands of years have showed different therapeutic effects with few side effects by pharmacological studies and clinical trials. However, to screen and purify the bioactive compounds of nature products is not easy because natural products are complex mixtures containing up to hundreds or even thousands of various constituents.¹ Recently, the introduction of HPLC equipped with mass spectrometry has been applied to screen and identify unknown compounds in natural products. The LC-MS coupled with DPPH assay which is performed DPPH radical scavenging assay after separate extracts by LC-MS determine not only information of the compound but also antioxidant activity of the compound.²,³

Polygonum aviculare L. (common knotgrass, birdweed, and pigweed) belongs to the family Polygonaceae and is an annual plant which is widely distributed through Korea, China, Japan and so on. It can be found in field or roadside easily.⁴ Traditionally, the sprouts of P. aviculare has been used for food. Furthermore, it also has been used important traditional medicine to treat rheumatitis, gout, diarrhea, dysentery, hemoptysis and bleeding.⁵ A previous study have been reported flavonoids (quercetin, kaempferol, myricetin, isorhamnetin, luteolin, avicularin, myricitrin, rutin, quercitrin and astragalin)⁶ and phenolic acids (gallic acid, protocatechuic acid and chlorogenic acid)⁴ from P. aviculare. Its compounds provide many pharmacological effects, such as anti-atherosclerotic,⁷ anti-hypertensive, diuretic, anti-obese, astringent, insecticide,⁸ antioxidant,⁴ anti-tumor⁹ and anti-inflammatory.¹⁰ Therefore, the purpose of the study is to simultaneously identify antioxidative compounds from P. aviculare efficiently using time-based DPPH assay regardless of the time consuming process.

Experimental

General experimental procedure – NMR spectra were recorded on a Bruker SPECTROSPIN 300 MHz spectrometer. Silica gel (70 - 230 mesh, Merck, Germany) and Sephadex LH-20 (25 - 100 μm, Amersham Biosciences, Sweden) were used for open column chromatography (CC). Thin-layer chromatography (TLC) was performed on a precoated silica gel 60 F₂₅₄ (0.25 mm, Merck, Germany). Medium pressure liquid chromatography (MPLC) was performed on Biotage Isolera One system equipped
with a HPFC pump, a variable dual-wavelength detector and a collector. Silica gel column for MPLC was SNAP Ultra 25 g. HPLC grade solvents (acetonitrile, water and methanol) were purchased from Fisher scientific Korea Ltd. Formic acid, DPPH (1,1-diphenyl-2-picrylhydrazyl) and curcumin were purchased from Sigma Aldrich Co.

**Plant material** – The herb of *P. aviculare* was purchased from Kyungdong Oriental Herbal Market, Seoul, Korea. The voucher specimen (KUP-HD051) was deposited in the laboratory of pharmacognosy, College of Pharmacy, Korea University.

**HPLC conditions** – The chromatography analytical procedures were performed on an Agilent 1260 Series (Agilent, Santa Clara, CA, USA). LC system equipped with a binary pump, an online degasser, an auto plate-sampler, and a thermostatically controlled column compartment. The columns were maintained at 30 °C. The separation was carried out on an Shiseido CapCell PAK C18 column (150 × 4.6 mm, 5 μm), preceded by a C18 guard column (4.00 × 3.00 mm; Phenomenex, USA). The binary gradient elution system consisted of 0.1% formic acid in water (solvent A) and 0.1% formic acid in acetonitrile (solvent B) and separation was achieved using the following gradient: 15% B at 0 - 5 min; 15 - 95% B at 5 - 30 min. The flow rate was 0.6 mL/min and the injection volume was 10.0 and 20.0 μL. In case of 20.0 μL, it is used for time-dependent DPPH assay.

**Mass spectrometry conditions** – Mass spectrometry was performed using an Agilent 6530 Q-TOF mass spectrometer (Agilent, Santa Clara, CA, USA) equipped with an electrospray ionization (ESI) interface, and was operated in negative ion mode with parameters set as follows: capillary voltage, 4000 V; fragmentor, 175 V; skimmer, 65 V; OCT 1 RF Vpp, 750 V; pressure of nebulizer, 40 psi; drying gas temperature, 325 °C; sheath gas temperature, 350 °C. Nitrogen was used as sheath and drying gas at a flow rate of 10.0 and 12.0 L/min, respectively. Data were collected in centroid mode and the detection mass range was set at m/z 50 - 1000 using the extended dynamic range. The accurate-mass capability of the TOF analyzer allowed reliable confirmation of the identity of the detected metabolites, normally with mass errors below 5 ppm in routine analysis. An external calibration solution (Agilent calibration solution A) was continuously sprayed in the ESI source of the Q-TOF system, employing the ions with m/z 112.9855 (TFA anion) and 1033.9881 [HP-0921 (TFA adduct)] to recalibrate the mass axis, ensuring mass accuracy and reproducibility throughout the chromatographic run. The Mass Hunter Workstation software LC/MS Data Acquisition for 6530 series Q-TOF (version B.05.00) was used to control all the acquisition parameters of the HPLC-Q-TOF-MS system and also to process the obtained data.

**DPPH assay** – For the measurement of DPPH radical scavenging activity, 190 μL of an ethanol solution of 15 μM DPPH was mixed with 10 μL of ethanol with or without the different concentrations of the tested samples such as extract, n-hexane fraction, EtOAc fraction, n-BuOH fraction, H2O fraction and compounds were isolated from the EtOAc fraction in a 96-well plate. For time-based DPPH assay, each 30 seconds’ effluent of HPLC was manually collected during 30 minutes on each well of 96-well plate. And then samples of each well were applied to the DPPH assay. The plate was incubated in a dark room at room temperature for 30 min and the reduction of the DPPH free radical was measured by reading the absorbance at 517 nm. DPPH, a purple-colored, stable free radical is reduced to the yellow-colored diphenylpicrylhydrazine when antioxidants are added. Curcumin was used as positive control. The inhibition ratio (percent) was calculated from the following equation: % inhibition = [(1 – (S – So) / (C – Co)) × 100, where S is the absorbance of DPPH 190 μL + sample 10 μL; So is the absorbance of ethanol 190 μL + sample 10 μL; C is the absorbance of DPPH 190 μL + ethanol 10 μL; Co is the absorbance of ethanol 190 μL + ethanol 10 μL.

**Extraction and isolation** – The herb of *P. aviculare* (1.0 kg) were accurately weighed and extracted with 80% methanol (MeOH) for 90 min in an ultrasonic apparatus. After evaporation of most of the solvent, the residue (107.23 g) was suspended in H2O and then partitioned successively with n-hexane, ethyl acetate (EtOAc) and n-BuOH. The EtOAc fraction (7.99 g) was subjected to silica gel column chromatography using gradient elution with n-hexane 100% to MeOH:EtOAc (1:1), in order to divide the fraction into fifteen fractions (E1–E15). Fraction E4 was re-chromatographed on sephadex LH-20 using isocratic elution with MeOH to give four fractions (E4-1–E4-4). Compound 2 (64.9 mg) was obtained from E4-4. Fraction E6 was separated over sephadex LH-20 using isocratic elution with MeOH to yield three fractions (E6-1–E6-3). Compound 4 (2.6 mg) obtained from E6-2. Fraction E11 was subjected to sephadex LH-20 with MeOH to afford five fractions (E11-1–E11-5). Compound 3 (27.1 mg) was obtained from E11-4. Fraction E7 was recrystallized with n-hexane. According to the TLC profiles, the crystal was re-chromatographed on sephadex LH-20 using isocratic elution with MeOH to afford nine fractions (E7A-1–E7A-9). Compounds 7 (7.8 mg) and 5 (10.5 mg) were obtained from E7A-3 and E7A-6, respectively.