



Fast gas chromatography–surface acoustic wave sensor: An effective tool for discrimination and quality control of Lavandula species

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ABSTRACT

Fast gas chromatography combined with surface acoustic wave sensor (GC/SAW) has been applied for the detection of the volatile aroma compounds emanated from various Lavandula medicinal plants such as *Lavandula angustifolia* (Norfolk, England; Tasmania, Australia; Sault, France; Sequim, USA), *Lavandula hybrida* (Valensole, Sault, France), *L. angustifolia* “Royal Velvet”, *L. hybrida* “Grosso”, and *L. hybrida* “Provence” (Olympic, USA). The GC/SAW involving the fragrance pattern analysis provides a novel analytical method with a very fast separation and the characterization of aromas caused by the difference of chemical composition according to botanical origin. The aim of this study is to show the capability of GC/SAW for the characterization and pharmacologically active aroma quality control to the analysis of Lavandula species. Finally, GC/SAW provided sufficiently reliable results for species identification and quality control.

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1. Introduction

Lavender is one of the most used important aromatic and medicinal plants. Lavandula essential oils are widely used in aromatherapy, fragrance, and pharmaceutical industries. Lavandula is a genus of about 25–30 species of aromatic annual herbs in the family Lamiaceae (mint family), and native to Mediterranean region, tropical Africa and east to India. Its essential oil has many clinical benefits such as antibacterial, antifungal, carminative, sedative, antidepressive, effective for burns and insect bites [1,2]. Thus, the composition analysis of essential oil of aromatic medicinal plants is of great interest in medicinal, broad fragrance industries, and clinical therapy.

For many years, GC and gas chromatography–mass spectrometry (GC–MS) have been widely used because of high resolution capacity for the characterization of the volatile aroma components in Lavandula species. Before volatile aroma compounds can be analyzed, they have to be extracted from the plant material. A wide variety of extraction methods including solvent extraction [3], Soxhlet extraction [4], steam distillation [5], and simultaneous distillation extraction [6] can be used for that purpose. These methods involve excessive manipulation of the sample, a very costly, time-

consuming procedure, low extraction efficiency, losses of some volatile compounds, degradation through thermal or hydrolytic effects which may lead to incorrect results. Especially, volatile aroma analysis demands rapid and simple procedure, because aroma components are thermally sensitive and decomposed to the different new odors. Since 1989, headspace solid-phase microextraction (HS-SPME) as a sampling technique based on sorption of analytes onto an absorbent coated fused silica fiber and then thermally desorption directly into a GC injection port has been introduced for purpose of volatile and semi-volatile compounds analysis in plants, food, biological, and environmental fields. Headspace sampling has many advantages, the most important of which is the elimination of much of the interferences arising from the sample matrix. The merits of the HS-SPME method include solvent-free, smaller sample volumes, and shorter analysis time [7–10]. Also, aromas are usually composed of complex mixtures of many volatiles, human sensory evaluation by trained panelists is important in aroma analysis. However, it has many limitations which involve a very expensive, time consuming procedure, and subjectiveness of expert. Finally, there is enormous request for the development of analytical method which provides rapid, simple, on-line measurement, non-destructive, low-cost procedure and rapid sensory information.

Consequently, one of such methods seems probably the electronic nose (eNose) which is rapid and high sensitivity for the analysis of volatile components [11,12]. But, aroma analysis with the E-nose has not always been successful. It is very sensitive to drift and lacks the possibility for identification of the different

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aroma components causing the signal change [13]. Interestingly, eNose method combined with a principal component analysis (PCA) has successfully applied for the discrimination of olive oils of different qualities and different geographic origins [14,15].

Since 1998, a new technique, fast gas chromatography with surface acoustic wave sensor (GC/SAW, zNose), called electronic nose based on the headspace sampling method provides second unit analysis, simple, low-cost procedure, real time detection, and rapid sensory information [16–27]. Especially, the characteristic of simplicity of fast GC/SAW method does not demand extraction procedures and provides on-line measurements. The headspace vapor is swept into the inlet via a pump, and then the vapor passes through the valve where the compounds are adsorbed onto the trap inside the system. Switching the valve to the injection process causes helium gas to flow backwards through the trap and onto the column. Therefore, GC/SAW has diminished decomposition of plant materials, minimized activity of enzyme. Especially fast GC/SAW permits a high reproducibility and good sensitivity at the high picogram to nanogram level, makes it possible to detect sensitive living volatile materials quantitatively [25]. Also, GC/SAW allows pattern recognition by a visual fragrance pattern, called a Vapor-Print or chromatographic fingerprint derived from the frequency of a SAW sensor. Fragrance pattern analysis also enables to detect environmental pollutant [16,17], detect off-flavors in wine [18], monitor the change of pharmacologically active volatiles according to real-time for living-organisms [21,25], discriminate the botanical and geographical origin for organism [22,23,26,27]. The advantages and the method validation of GC/SAW, adaptability to a variety of applications were reported in my previous publications [25–27].

The aim of this study is to show the capability of GC/SAW for the characterization and pharmacologically active aroma quality control to the analysis of *Lavandula* species. In addition, volatile composition profile data for *Lavandula* species by GC/SAW were analyzed by PCA. Also, HS-SPME-GC-MS method was employed to confirm the identification of aroma components of *Lavandula* species and compared to GC/SAW.

2. Experimental

2.1. Materials

Dried buds of *Lavandula* species (*Lavandula angustifolia*, *Lavandula hybrida*, *L. angustifolia* “Royal Velvet”, *L. hybrida* “Grosso”, *L. hybrida* “Provence”) were obtained from lavender farms of Norfolk, England; Tasmania, Australia; Valensole, Sault, France; and Sequim, Olympic, USA. All standard chemicals of analytical grade were purchased from Sigma–Aldrich (St. Louis, Mo, USA) and Tokyo Kasei (Nihonbashi, Tokyo Japan). Organic solvents of a chromatographic grade were obtained from J.T. Baker. The commercially available carboxen-divinylbenzene-polydimethylsiloxane (CAR-DVB-PDMS) SPME fiber (film thickness, 50/30 μm) was purchased from Supelco (Bellefonte, PA, USA) and used.

2.2. GC/SAW (zNose) description

GC/SAW (4100 vapor analyzer, Electronic Sensor Technology, New Bury Park, USA) composed with the fast gas chromatograph, a fast response integrating surface acoustic wave sensor detector, a pre-concentrating trap containing Tenax absorbent, a short GC column is used to detect vapors of the volatile and semi-volatile organic compounds. The SAW sensor consists of an uncoated 500 MHz acoustic resonator, bonded to a thermoelectric heat pump to heat or cool the quartz substrate. The GC/SAW is especially sensitive to low concentrations [25].

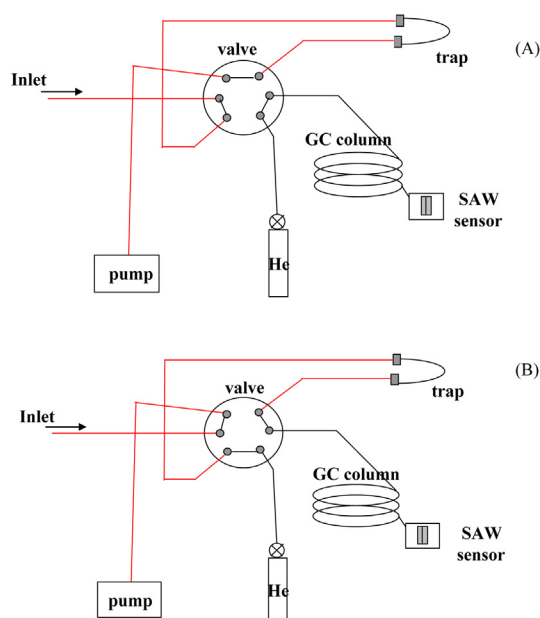


Fig. 1. Schematic diagram of the headspace sampling process (A) and injection process (B) by GC/SAW.

2.3. GC/SAW analytical conditions

About 0.8 g of each dried *Lavandula* sample was weighed into a 40-ml glass vial sealed with a screw cap containing a Teflon/silicone septa. The capped vial was kept to equilibrate with the headspace vapor at 25 °C for 1 h just before analysis.

The headspace vapor of sample was swept into the trap inside the system for 1 s (Fig. 1(A)). The trap was heated rapidly to 200 °C to desorb and transfer the volatiles to the analytical column (Fig. 1(B)). GC column was heated from 30 °C to 90 °C at a rate of 2 °C/s. Helium (99.999%) was used as a carrier gas at 3.0 ml/min (0.05 ml/s). 6% Cyanopropyl phenyl polydimethylsiloxane (DB-624, J&W Scientific, Folsom, CA, USA, 1 m \times 0.25 mm i.d., 0.25 μm film thickness) fused silica capillary column was used. The set-up temperatures were at 20 °C for sensor, 130 °C for inlet port, and 110 °C for valve. Triplicate measurements per sample were carried out. All analytical procedures were completed within 30 s. The shorter total time-to-result per sample allows several replicated analyses of a sample.

2.4. Headspace solid-phase microextraction

About 2.0 g of dried *Lavandula* sample was placed in 25-ml vial sealed with an aluminum cap containing a Teflon/silicone septa. The capped vial was kept to equilibrate at 25 °C for 1 h before HS-SPME sampling. The carboxen-divinylbenzene-polydimethylsiloxane (CAR-DVB-PDMS) SPME fiber (film thickness, 50/30 μm) was used because it was most efficient among the various types of fiber for most volatile organic compounds [7,28]. The SPME fiber was exposed to the headspace above the *Lavandula* sample vial at 25 °C for 1 h. After adsorption, the SPME fiber was retracted from the sample vial and immediately inserted into the injection port of the GC-MS where thermal desorption was performed at 240 °C for 1 min.

2.5. GC-MS analysis

All experiments were performed on a Hewlett-Packard 6890 Series GC system with an Agilent 5973 N Mass Selective Detector (Agilent Technologies, Wilmington, DE, USA) equipped with 5% phenyl polydimethylsiloxane (Ultra 2 column, Agilent,

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