Solid Phase Micro-Extraction (SPME) Coupled with Selectable 1D/2D GC-MS for the Determination of Food Product Flavors

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KEYWORDS
Selectable 1D/2D GC-MS, SPME, food, flavor, fragrance

ABSTRACT
Identification of important trace components in complex samples like fragrances, natural products, polymers or food products can be challenging. Achieving the mass on column and resolution necessary to locate peaks and identify trace components using a single column chromatographic separation can be difficult, if not impossible.

A selectable 1D/2D GC-MS configuration based on Agilent® Technologies capillary flow technology (CFT) and low thermal mass (LTM) GC column modules with dissimilar column phases was used to perform two-dimensional GC analysis of different foodstuffs. Heartcutting was used to transfer analytes from the first to the second column. Mass spectrometry and olfactory detection were performed in parallel. Solid Phase Micro-Extraction was used as a solventless means to introduce sufficient mass of sample onto the precolumn of the multi-dimensional system. SPME offers the added benefit of enabling “tuning” of the selectivity of the extraction through the choice of coating on the fiber.

Separation and identification of selected flavor compounds from food products were used to demonstrate the effectiveness of the system. The main advantages of this configuration were the simple selection of one or two dimensional operation in combination with the ability to use mass spectrometry and olfactory detection in both dimensions for the analysis of odor active compounds.
INTRODUCTION

Two dimensional gas chromatography using heartcutting is an effective tool for resolving compounds in a complex matrix. Typical systems use a non-selective detector, such as an FID, in the first dimension and a selective detector, such as an MSD, in the second. The system described in this work uses capillary flow technology (CFT) hardware from Agilent Technologies to enable parallel mass spectrometry and olfactory detection for both the first and second dimension separation. The system requires no hardware changes for operation in either one- or two dimensional mode. The effectiveness of the system is demonstrated in this study using potato chips and spices as examples.

EXPERIMENTAL

Instrumentation. Analyses were performed on a 6890 equipped with a 5975C inert XL MSD with triple axis detector and LTM Columns (Agilent Technologies), PTV inlet (CIS 4, GERSTEL), MPS 2 Sampler with SPME Option (GERSTEL), and an Olfactory Detection Port ODP 2 (GERSTEL).

Analysis conditions.

MPS: SPME (DVB-Carboxen-PDMS)
CIS 4: SPME liner
   split, 10:1
   250°C (isothermal)
Pneumatics: constant pressure
Oven: 250°C, held for duration
1st Column: 30 m Rtx-5Sil MS (Restek),
   LTM format
   \[d_1 = 0.25 \text{ mm} \quad d_i = 0.25 \mu \text{m}\]
   40°C (1 min); 10°C/min; 160°C; 140°C/min; 300°C (3 min)
2nd Column: 30 m DB-Wax (Agilent), LTM format
   \[d_1 = 0.25 \text{ mm} \quad d_i = 0.5 \mu \text{m}\]
   40°C (17 min); 10°C/min; 210°C; 170°C/min; 40°C
MSD: full scan, 40-350 amu
MSD/ODP: 1/1 split

Sample preparation. Potato chips and spice samples were purchased at a local grocery store. The potato chip samples were crushed. A 3.0 gram sample was placed in a 20 mL screw capped vial. The spice samples, approximately 0.25 grams, were placed directly in a 20 mL screw cap vial.

RESULTS AND DISCUSSION

A diagram of the selectable 1D/2D GC-MS system configuration is shown in Figure 1. Details of the setup have been described elsewhere. [1,2]

![Figure 1. Diagram of Selectable 1D/2D GC-MS setup.](image)

The first example shows the determination of volatile compounds in the headspace inside a potato chip bag based on a single dimension separation. A bag of chips was placed on the dashboard of a car for four hours. The car was parked in the sun to accelerate the degradation of the product. When the bag was opened, a distinct rancid odor was detected. Figure 2 shows a comparison of the chromatograms obtained from fresh and rancid chips. The compounds responsible for the rancid odor, mostly aldehydes, are easily seen. Figure 3 shows a comparison of two different brands of chips. Brand Y shows a higher level of acetone and hexanal, indicating that these chips are less fresh.
Figure 2. Total ion chromatograms obtained from fresh and rancid chips.

Figure 3. Total ion chromatograms obtained from two potato chip brands.

Figure 4 shows a comparison of plain and flavored chips from the same manufacturer. The compounds associated with the individual flavors are easily seen. The sour cream and green onion flavor chromatogram shows some methoxy-propoxypropanols which may stem from the packaging material. The salt and vinegar flavor chips exhibit a large acetic acid peak, and the barbecue chips show a very complex chromatogram with a wide range of terpenes. Olfactory detection combined with automated voice recognition and text conversion was performed on this sample. The result was a rather confusing list of descriptors from overlapping peaks in the chromatogram. The section of the chromatogram from 6.5-10.5 minutes, was then heart-cut and transferred to
the second column for further separation. The first dimension column was backflushed starting at 10.6 minutes. The resulting chromatogram is shown in Figure 5.

As can be seen, additional compounds are separated in the second dimension enabling their identification along with a description of their olfactory qualities. Olfactory detection of the compounds separated in the 2nd dimension was used to identify important pyrazines, which are formed during the cooking process and which contribute to the fried potato flavor [3]. Figure 6 shows an enlarged view of the region from 15.5-19.5 minutes. Table 1 list the identities, descriptors and literature aroma profile for the compounds labeled in Figure 6.
The second example shows the SPME headspace analysis of a poultry seasoning. The seasoning declaration of ingredients listed thyme, sage, marjoram, rosemary, black pepper, and nutmeg. Figure 7 shows the 1D chromatogram from the headspace SPME extraction of the poultry seasoning. Overlapping peaks on the front of the cubebene peak were cut over to the 2nd dimension for further separation and identification. Figure 8, shows the resulting 2D chromatogram.

Table 1. Peak identities for Figure 6.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Descriptor</th>
<th>Aroma profile</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Dimethylpyrazine isomer</td>
<td>Burnt</td>
<td>Nutty, roasted</td>
</tr>
<tr>
<td>2 Dimethylpyrazine isomer</td>
<td>Burnt</td>
<td>Nutty, roasted</td>
</tr>
<tr>
<td>3 Ethyl methylpyrazine</td>
<td>Barbecue</td>
<td>Potato, hazelnut, coffee</td>
</tr>
<tr>
<td>4 Trimethylpyrazine</td>
<td>Burnt Potato</td>
<td>Roasted, baked potato</td>
</tr>
<tr>
<td>5 Dimethyl ethylpyrazine</td>
<td>Potato</td>
<td>None found</td>
</tr>
</tbody>
</table>

Figure 6. Enlarged view of a section of the chromatogram obtained from barbecue flavored chips.

Figure 7. Total ion chromatogram obtained from poultry seasoning.
Figure 9 shows an enlarged view of the area from 20-22 minutes. Two peaks with floral notes, isoledene and ylangene, are well resolved from the cubebene (woody note). The figure also shows the overlay of three separate runs demonstrating the good reproducibility of the system. Only a small portion of the cubebene peak from the first dimension was cut and transferred to the 2D column, so one would not expect good precision for this peak in the second dimension.

Figure 9. Enlarged stacked view of three chromatograms obtained from poultry seasoning.
Stereochemistry can also be an important factor in the odor profile of a compound. The last example shows the use of a chiral column in the second dimension to separate isomers. A standard of 3,3,5-trimethyl-2-cyclohexen-1-ol was introduced into the GC using SPME. Figure 10 shows the 1D and 2D chromatograms for this sample. In the first dimension, a single peak is seen with a flavor descriptor of “spicy”. When the peak is cut over to the 2nd dimension, four peaks are seen. The first major peak is the isomer responsible for the flavor (spicy, tobacco). None of the other isomers resulted in a perceptible olfactory impression at the ODP.

**Figure 10.** Comparison of 1D and 2D chiral separation for 3,3,5-trimethyl-2-cyclohexen-1-ol.

**CONCLUSIONS**

The selectable 1D/2D GC-MS system with a valve-less flow switching device is a powerful system for the analysis of trace components in complex samples. The main features of this configuration are the simple selection of one- or two dimensional operation. The setup of the system is facilitated through the use of GERSTEL 1D/2D Sync software, which calculates critical method parameters for the selected column set.

**REFERENCES**

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