## Gaseous Aliphatic Aldehydes in Chinese Incense Smoke

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Aliphatic aldehydes were found during the combustion of materials. Tobacco smoke contains aldehydes (Crupm and Gardiner, 1989). Fire fighters were exposed to aldehvdes when thev conducted firefighting (Brandt-Rauf et al, 1988). Aldehydes in ambient air come mainly from the incomplete combustion of hydrocarbons and from photochemical reaction (World Health Organization, 1987). Most aldehydes in ambient air are formaldehyde and acetaldehyde 1988). (Kalabokas et al. Formaldehvde. acetaldehvde. propionaldehyde, butyraldehyde, and benzaldehyde were found in the atmosphere in Los Angeles (Grosjean, 1982).

Burning Chinese incense for whorshipping deities is a Chinese was suspected to be a factor daily routine. It causing Chen et al nasopharvngeal cancer(Mimi et al. 1990: 1987). Epidemiological studies correlated it with the high risk of childhood brain tumor (Martin et al, 1982) and the high risk of childhood leukemia (Lowengard et al, 1987). Ames test identified the mutagenic effect of the smoke from burning Chinese incense The smoke had been proved to contain 1987). (Rasmussen, polycyclic aromatic hydrocarbons and aromatic aldehydes (Schoenotol and Gibbard, 1967). Suspicion about formaldehyde and other alphatic aldehydes was evoked, when a survey of indoor air pollution was conducted in Taipei city (Lin and Yao, 1993). This study determined the presence of aliphatic aldehydes in the smoke from burning Chinese incense under a controlled atmosphere.

## MATERIALS AND METHODS

Four types of Chinese incense containing sandalwood powder were used in this study. They were made in Taiwan, Mainland China and Japan. A chamber  $(68 \times 48 \times 44 \text{ cm}^3)$  was constructed by using polypropylene, for burning the Chinese stick incense. A constant

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flow (5 L/min) of purified air was humidified by applying a Flow-Temperature-Humidity Control System (Model HCS 301, Miller Nelson through the chamber to the exhaust. Research) and flowed Temperature and humidity inside the chamber were monitored by thermometer and Dew Point Meter (model 91HC, YSI). A stick of incense was ignited by a nickel-chrome wire controlled by a The smoke generated in the chamber was voltage regulator. collected by a 2,4-dinitrophenyl-hydrazine coated Sep-Pak DNPHsilica cartridge (Part No.37500, Millipore) in 2 L/min for 2min. Then, the Sep-Pak was eluted by 5 mL acetonitrile (BDH, Hipersolv, England). The high performance liquid chromatograph (Model 590, Waters, USA) equipped with U.V detector at  $\lambda$  max 254nm was utilized to determine the derivatives of aliphatic aldehydes in the eluent. The injection volume was 10  $\mu$  L. The column was Nova-Pak C18 (150mm  $\times$  3.9mm) and the flow rate of the solvent system (acetonitrile/H<sub>2</sub>O; 60/40 v/v) was 1.0 mL/min. A quantified and mixed aldehydes standard solution obtained from the aldehydes standard solution kit (TK-151, Alltech) was spiked onto the Sep-Pak and eluted by 5 mL acetonitrile for forming the stock solution of aldehydes 2,4-dinitrophenyl- hydrazones. From this solution, a series of standard solution was prepared. A quantified 4fluorobenzonitrile (46680, Fluka) in acetonitrile (0.134 g/10mL) serving as an internal standard was added to the standard solutions and to the eluents of the samples. The retention time of each derivative of aldehvdes relative to that of the internal standard was determined. The relative peak area for each aldehyde derivative was also calculated when a sample was quantified. After the analysis by HPLC were completed, the samples were again analyzed by employing GC/MS (gas chromatograph / mass spectrometer, HP 5890 SeriesII/HP 5971) equipped with HPG 1034B MS Chemstation. The column of GC was HP-1 column (methylene silicon  $12m \times 0.2mm \times 3 \mu m$ ). Helium was used as the carrier gas, at a flow rate of 0.72 mL/min. The operating conditions of the mass spectrometer were: ionization energy 70ev, accelerating voltage 1055 volt, and resolution 1 amu.

A field sample was also collected by a Sep-Pak DNPH-silica cartridge and analyzed by HPLC. The cartridge was located 150cm above ground and 10cm away from an incense holder in a courtyard of Lungshan Temple, a Chinese temple in Taipei.

## RESULTS AND DISCUSSION

Figure 1. shows the HPLC chromatograms of the derivatives of aldehydes in the smoke from burning four types of Chinese incense. Formaldehyde, acetaldehyde, acrolein and propionaldehyde were

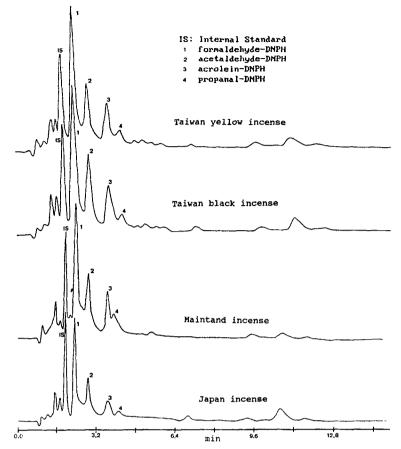


Figure 1. The HPLC chromatograms of the derivatives of aldehydes in the smoke from burning four types of Chinese incense.

 Table 1. Retention time of aldehydes-DNPH relative

 to internal standard

	Relative retention time							
Aldehydes-DNPH*	n**	Mean	S.D.	C.V.(%)				
formaldehyde-DNPH	12	1.217	0.007	5.75				
acetaldehyde-DNPH	12	1.495	0.019	1.27				
acrolein-DNPH	12	1.910	0.030	1.57				
propanal-DNPH	12	2.134	0.034	1.59				
methacrolein-DNPH	12	2.817	0.070	2.48				
butanal-DNPH	12	2.981	0.040	1.34				

\* DNPH:dinitrophenylhydrazone

\*\* n:observation number

identified based on retention times relative to the internal standard as shown in Table 1.

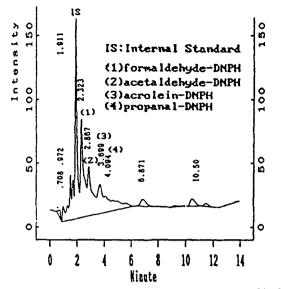


Figure 2. The HPLC chromatogram of the DNPH-derivatives of aldehydes in the smoke collected from a Chinese temple.(DNPH: dinitrophenylhydrazone)

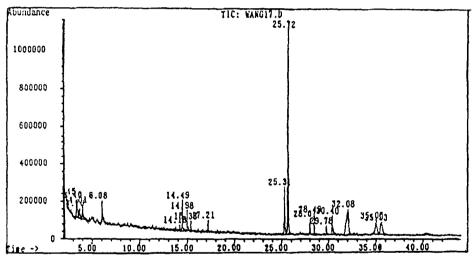
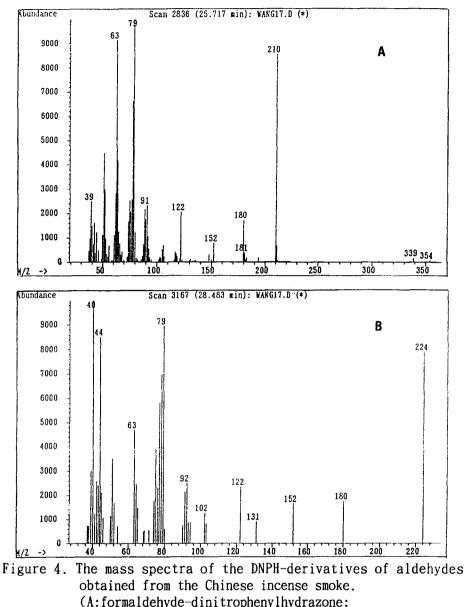


Figure 3. GC chromatogram of the DNPH-derivatives of aldehydes obtained from the Chinese incense smoke.

Figure 2. gives the HPLC chromatogram of the DNPH-derivatives of aldehydes in the smoke collected from a Chinese temple. The above mentioned aldehydes were also found on the chromatogram. The results of GC/MS examination of Chinese incense smoke is shown in Figures 3, 4 and 5. Figure 3 is the GC chromatogram.



B:acetaldehyde-dinitrophenylhydrazone.)

Figure 4-(A), 4-(B), 5-(A) and 5-(B) show the mass spectra from the compounds with retention time of 25.717 min, 28.483 min, 30.406 min and 32.079 min respectively. The respective molecular ion pattern at m/e 39-210, 40-224, 40-236 and 40-276 is typical of formaldehyde-DNPH, acetaldehyde-DNPH, acrolein-DNPH and furnanal-DNPH. Propionaldehyde was not identified by GC/MS, probably because the amount was so small that it could not be detected.

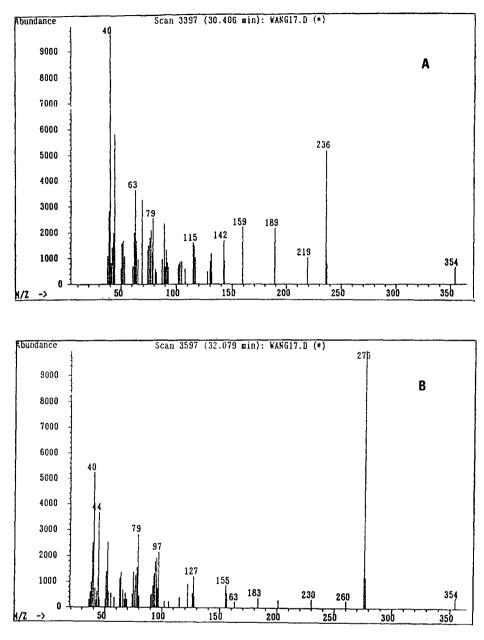


Figure 5. The mass spectra of the DNPH-derivatives of aldehydes obtained from the Chinese incense smoke. (A:acrolein-dinitrophenylhydrazone; B:furnanal-dinitrophenylhydrazone.)

Tables 2. indicates the concentration of each aliphatic aldehyde generated by burning one gram of incense under controlled atmosphere with relative humidity ranging from 46.0% to 83.2%. The

result shows that the ppm/g varies with the type of Chinese incense and the relative humidity. The concentration of aliphatic aldehydes decreased with an increase of humidity. Overall, formaldehyde was predominant in the smoke from burning Chinese incense.

Tables 2. Generation of aliphatic aldehydes from burning one gram of incense at various humidities

	Concentration normalized to gram of incense burned at controlled relative humidity and temperature, ppm/g											
Aldehydes	RH 46.0%; 24.8°C				RH 64.3%; 21.2°C				RH 83.2%; 23.4°C			
	1	2	3	4	1	2	3	4	1	2	3	4
formaldehyde acetaldehyde												
acrolein												

- \* 1 Taiwan yellow incense
  - 2 Taiwan black incense
  - 3 Mainland incense
  - 4 Japan incense
- \*\* Observation numbers in each set were above two times

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