Application News

Gas Chromatograph Nexis[™] GC-2030

Assessment of JetanizerTM and Quantitative Analysis of CO_2 and CH_4 in the Atmosphere

H. Miyabayashi, S. Uchiyama, M. Ishimoto

User Benefits

- ◆ Simplification of the analytical system for CO and CO₂ utilizing the Jetanizer coupling with GC-FID.
- ◆ Low-cost system achieved with adding the Jetanizer to FID-2030 that is already working.
- ◆ N₂ as carrier gas available.

■ Introduction

More and more research and development projects are focusing on being "Carbon Neutral", which is the pursuit of zero Greenhouse gas production from human industrial activity. Greenhouse gas includes Carbon Dioxide (CO2) and Methane (CH₄). In addition, high-purity hydrogen requires high-sensitive analytical system for the impurities such as CO, CO₂, and CH₄. Therefore, easy quantification methods for these gases are required in various fields for becoming "Carbon Neutral". While TCD and BID are excellent for the analysis of inorganic gases, they both have difficulties with these applications. Highly sensitive analysis is not possible in TCD, and carrier gas is limited to Helium in BID. A methanizer can be installed to utilize FID for CO and CO₂ analysis. A traditional methanizer uses a Nickel (Ni) catalyst in a hydrogen rich environment to reduce CO and CO2 to CH4 for analysis by FID. This setup requires an additional heater and gas line for installation. The Ni catalyst is also highly sensitive to oxidation by oxygen. The Jetanizer uses a novel, oxygen durable catalyst in the FID nozzle. The hydrogen for the FID flame provides the hydrogen rich environment for reduction and the FID heater is utilized for heating the catalyst. Thus, the whole system is durable for oxygen, simple to install, and can expand its usefulness to wider fields. This article introduces how the Jetanizer on the GC-2030 is useful for analysis of CO, CO₂ and CH₄.

■ Jetanizer

The Jetanizer is a compact Methanizer for FID developed from Activated Research Company. Highly efficient conversion of $\mathrm{CO/CO_2}$ into $\mathrm{CH_4}$ can be achieved with a catalyst inside FID nozzle. No extra line or heating device must be equipped. Exchanging FID nozzle is the only hardware change for the analysis. Fig. 1 shows an image of the Jetanizer. The insertion depth of the capillary column is shortened to 45 mm to avoid interfering with the catalyst inside the nozzle (usually the tip length is 72 mm).



Fig. 1 JetanizerTM

■ Analytical Condition

Analytical evaluation was performed with our GC-2030(FID) that employs a gas sampling valve. The Jetanizer is installed in the same way as the FID-2030 nozzle . A metal guard column was connected to the end of the capillary column because the Jetanizer operating temperature is 400°C. Schematic illustration of the whole analytical system is shown in Fig. 2. A packed GC column was also evaluated for more loading capability. Table 1 shows the concentration of the standard gas samples (base gas was N₃). Table 2 shows the analytical condition

Table 1 Concentration of the standard gas samples

	1	25	100	0.1	1	25
	(ppm)	(ppm)	(ppm)	(%)	(%)	(%)
СО	1.00	25.2	100	0.101	1.02	24.8
CO ₂	0.99	25.4	100	0.100	0.999	25.1
CH_4	0.98	24.6	98.2	0.0996	0.994	25.1

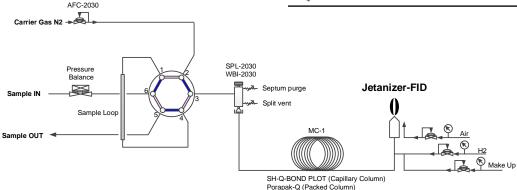


Fig. 2 Schematic illustration of the analytical system

Table 2 Analytical Condition

Model	: Nexis GC-2030		
Gas Sampler	: MGS-2030 + 1 mL Loop		
Inj. Temp.	: 250 °C		

Inj. Mode : ① Split 1:5 ② Direct

Carrier Gas : N₂ Column Flow : 1) 40 cm/sec ② 40 mL/min

: $\hat{\mathbb{I}}$ SH-Q-BOND PLOT Column (30 m imes 0.53 mm l.D., 20 Column

 \odot Porapak-Q 50/80 (2 m imes 2 mm l.D.)

Column Temp. : 40 °C

: FID + Jetanizer (column insertion depth : 45 mm) Detector

FID Temp. : 400 °C

: ① N_2 , 24 mL/min ② N_2 , 4 mL/min Makeup Gas

H₂ Flow : 32 mL/min Air Flow : 250 mL/min

①:with capillary column ②:with packed column

■ Fundamental Performance of Jetanizer

Repeatability of the peak area, linearity of the calibration curve, and the converting efficiency (C.E.) into Methane were evaluated with the standard gas samples shown in Table 1. Fig, 3-1 shows a chromatogram of 0.1% standard gas sample with capillary column and Fig. 3-2 shows the one with packed column. Each Limit of Detection (LOD) value was calculated with S/N from 1 ppm standard gas sample. Table 3 shows the result.

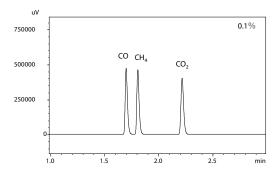


Fig. 3-1 chromatogram of 0.1% sample with capillary column

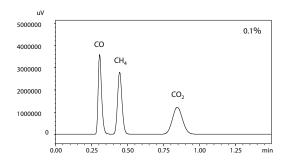


Fig. 3-2 chromatogram of 0.1%sample with packed column

Table 3 LOD of CO, CH₄, CO₂

	со	CO ₂	CH₄
Capillary column	0.34 ppm	0.37 ppm	0.27 ppm
Packed column	0.08 ppm	0.10 ppm	0.17 ppm

■ Repeatability of the Peak Area and **Efficiency of Methane Conversion**

CO and CO₂ cannot be directly detected with FID. Converting them into $\bar{\text{CH}}_4$ with the Jetanizer is crucial for the analysis and its efficiency is highly influential to the analysis.

C.E. can be calculated with peak area (A) and concentration (C) of CO/ $\rm CO_2$ and $\rm CH_4$. The relation between them can be described as formula (1). The area repeatability %RSD and C.E. of each sample are shown in Table 4-1 and Table 4-2.

C. E. (%) =
$$\frac{A(CO \text{ or } CO_2)}{A(CH_4)} * \frac{C(CH_4)}{C(CO \text{ or } CO_2)} * 100$$
 (1)

Table 4-1 Repeatability of peak area (n=10) and the C.E when using capillary column

	со		c	CO ₂	
	%RSD	C.E.	%RSD	C.E.	%RSD
1 ppm	2.11	97.6	2.23	100.3	2.48
25 ppm	0.39	98.4	1.03	97.8	0.74
100 ppm	0.29	100.2	0.35	100.2	0.28
0.1 %	0.08	99.0	0.09	99.2	0.09
1 %	0.17	99.8	0.21	100.2	0.16
25 %	0.33	100.9	0.41	100.0	0.33

Table 4-1 Repeatability of peak area (n=10) and the C.E when using capillary column

	со		c	CO ₂	
	%RSD	C.E.	%RSD	C.E.	%RSD
1 ppm	1.18	104.9	1.58	124.5	0.93
25 ppm	0.27	98.2	0.31	95.5	0.28
100 ppm	0.05	99.3	0.09	97.6	0.05
0.1 %	0.05	98.5	0.05	98.4	0.05
1 %	0.15	98.7	0.16	101.0	0.16
25 %	0.13	68.7	0.12	94.6	0.11

Sufficient repeatability and C.E. were obtained when using capillary column for both CO and CO2. In packed column analysis, the repeatability and C.E was sufficient except for CO in 25% standard gas sample. Here, the C.E. limited to 68.7% which is likely due to overloading the catalyst. In such case, reducing the volume of the sample injection should be an effective and direct solution. Slowing down the flow rate could also solve the low C.E. because the sample volume touching the catalyst per a unit of time should be decreased.

■ Linearity of the Calibration Curve

Fig. 4-1, 4-2, and 4-3 shows the calibration curves CO, CO $_2$ and CH $_4$, respectively. Calibration point of 25% standard gas sample was excluded in packed column. The coefficients of determination (R2) were more than 0.9999 for all CO, CO $_2$ and CH $_4$ when using capillary column. The R2 for CO, CO $_2$ and CH $_4$ were 0.9988, 0.9994 and 0.9996, respectively. Fig. 5 shows relative response factors (RRFs) from 1ppm to 25% standard gas samples when using capillary column. Contamination from the air slightly influenced the RRFs when the sample was 1ppm. The RRFs for CO, CO $_2$ and CH $_4$ were consistent for all other standard concentrations.

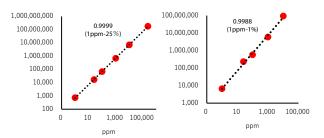


Fig. 4-1 Calibration curve of CO (Left: capillary column Right: packed column)

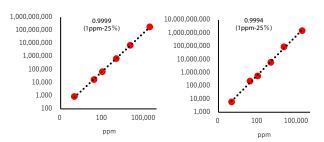


Fig. 4-2 Calibration curve of CO₂ (Left: capillary column Right: packed column)

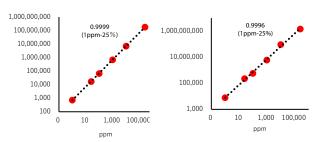


Fig. 4-3 Calibration curve of CH₄ (Left: capillary column Right: packed column)

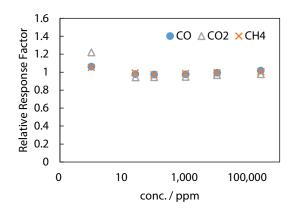


Fig. 5 Relative reference factors when using capillary column

■ Evaluation of Durability to Oxygen

Ni catalyst is vulnerable to oxygen damage; therefore, the analytical system must strictly exclude contamination of oxygen. The catalyst employed in the Jetanizer is durable for oxygen. For evaluating durability to oxygen, more than 3500 injections of air samples, containing approximately 20% oxygen, were completed. Periodically during the air sample injections, 0.1% standard gas sample was injected. Fig. 6 shows the $\rm CO_2$ /CH₄ value of each injection. Sensitivity fluctuation was not observed even after more than a total volume of 3.5 L of air was injected. The Jetanizer catalyst should be stable for exposure to oxygen.

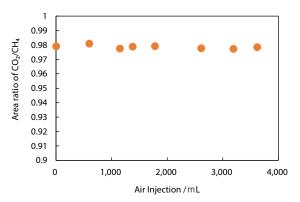


Fig. 6 CO_2 /CH₄ value of 0.1% standard gas sample

■ Quantification of CO₂ and CH₄ in the Air

Oxygen may produce a baseline deflection when eluting to the Jetanizer in air samples. Column selection for avoiding coelution is important in such cases. MICROPACKED-ST is an option for obtaining full separation of O2, CH₄ and CO₂.

The air sample was injected five times sequentially to the GC. Table 5 and Fig. 7 show the analytical condition and the obtained chromatogram, respectively. CH₄ and CO₂ were clearly separated from O2 . The CH₄ and CO₂ in the air were quantified using a single point calibration of a gas standard containing 100 ppm of CH₄ and 1000 ppm of CO₂ diluted with N2. The result is shown in Table 6. CH₄ was 2.1 ppm and CO₂ was 419 ppm, respectively. The repeatability of the area (%RSD) was sufficient. The concentrations and repeatability are shown in Table 6.

Table 5 Analytical Condition

Model	: Nexis GC-2030
Gas Sampler	: MGS-2030 + 1 mL Loop
Inj. Temp.	: 150 °C
Inj. Mode	: Split 1:3
Carrier Gas	: N ₂ , constant flow mode (10 mL/min)
Column	: MICROPACKED-ST (1.0 m $ imes$ 1.0 mm I.D.)
	(Input value to the software: 125 m \times 0.50 mm I.D., df = 15 μ m due to flow re-calculation)
Column Temp.	: 40 °C (1 min) \rightarrow 40 °C/min \rightarrow 200 °C (3 min)
Detector	: FID + Jetanizer (column insertion depth : 45 mm)
FID Temp.	: 400 °C
Makeup Gas	: N ₂ , 14 mL/min
H ₂ Flow	: 32 mL/min
Air Flow	: 250 mL/min

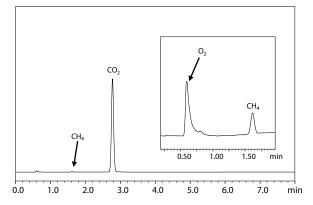


Fig. 7 Obtained chromatogram of the air sample

Table 6 Quantification(ppm) and repeatability (%RSD) result (n=5)

	CH₄	CO ₂
Quantification (ppm)	2.1	419
Repeatability (%RSD)	0.49	0.51

■ Summary

Analytical performance of the Jetanizer was evaluated on GC-2030 (FID) coupling with gas sampling valve. Converting efficiency from CO/ CO₂ to CH₄ was evaluated and obtained successful result. Fundamental analytical performance such as linearity and repeatability also shows sufficient result. Endurance for oxygen was also assessed with sequential injections of the air. The sensitivity remained stable even after 3500 air injections. CO₂ and CH₄ were quantified in the air

The Jetanizer will be useful in the situation such that "the concentration is too small for TCD" and "BID is not available due to Helium shortage".

■ Notice

Observe the following precautions when using the Jetanizer.

- Column insertion depth must be 45 mm. Inserting over 45mm of capillary column may damage the catalyst inside or the column itself.
- * Flow rate of the air should be around 250 mL/min.
- * Too much column flow may extinguish the flame of FID.
- * Baseline or noise level may be higher than usual FID analysis.
- * Hydrocarbons with triple bond structure (such as acetylene), substances with sulfur (such as hydrogen sulfide), and substances with halogens may influence the Jetanizer mechanism.
- * Sample matrix gas of O₂ or H₂ may influence the shape of the flame after elution. Therefore, quantification may be interfered immediately after elution of $\rm O_2$ and $\rm H_2$. The target compounds should be distant from $\rm O_2$ and $\rm H_2$ in chromatogram.
- *The Jetanizer is not a product of Shimadzu. The product is out of our support coverage.
- * Contact to our salesperson for more information.

Nexis is a trademark of Shimadzu Corporation or its affiliated companies in Japan and other countries. Jetanizer is a trademark or registered trademark of Activated Research Company in the United States and other countries.



Shimadzu Corporation

www.shimadzu.com/an/

For Research Use Only. Not for use in diagnostic procedures.

01-00599-EN First Edition: Oct. 2023

This publication may contain references to products that are not available in your country. Please contact us to check the availability of these

products in your country.
The content of this publication shall not be reproduced, altered or sold for any commercial purpose without the written approval of Shimadzu.
See http://www.shimadzu.com/about/trademarks/index.html for details.
Third party trademarks and trade names may be used in this publication to refer to either the entities or their products/services, whether or not

they are used with trademark symbol "TM" or "®". Shimadzu disclaims any proprietary interest in trademarks and trade names other than its own

The information contained herein is provided to you "as is" without warranty of any kind including without limitation warranties as to its accuracy or completeness. Shimadzu does not assume any responsibility or liability for any damage, whether direct or indirect, relating to the use of this publication. This publication is based upon the information available to Shimadzu on or before the date of publication, and subject to change without notice.