Short-Path Near-Infrared Laser Detection of Environmental Gases by Wavelength-Modulation Spectroscopy

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Abstract—The detection of environmental gases, $^{12}$CO$_2$, $^{13}$CO$_2$, and CH$_4$, using near-infrared semiconductor lasers with a short laser path length is studied by means of wavelength-modulation spectroscopy. The developed system is compact and has high sensitivity enough to detect the absorption peaks of isotopic $^{13}$CO$_2$ of a 3-5% CO$_2$ gas at 2 μm with a path length of 2.4 m, where its peak size is two orders of magnitude smaller than that of the ordinary $^{12}$CO$_2$ peaks. In addition, the detection of $^{12}$CO$_2$ peaks of a 385-ppm (0.0385-5%) CO$_2$ gas in the air is made at 2 μm with a path length of 1.4 m. Furthermore, in pursuing the detection of an ancient environmental CH$_4$ gas confined to a bubble in ice at the polar regions, measurements of the absorption spectrum for a trace gas of CH$_4$ in a small area are attempted. For a 100-% CH$_4$ gas trapped in a ~1 mm$^3$ glass container, the absorption peaks of CH$_4$ are obtained at 1.65 μm with a path length of 3 mm, and also the gas pressure is extrapolated from the measured data.

Keywords—Environmental Gases, Near-Infrared Laser Detection, Wavelength-Modulation Spectroscopy.

I. INTRODUCTION

The detection of environmental gases, such as CO$_2$ and CH$_4$, is now attracting considerable attention in the investigation of greenhouse effects [1], [2]. In particular, the measurement of the CO$_2$ concentration in the atmosphere is frequently made by LIDAR (Light Detection And Ranging) [3], which collects the back-scattering data from the target gas with a distance of kilometers by exposure with laser beams generated from high-power, sizable lasers, e.g., CO$_2$ and YAG lasers [4]–[8]. On the other hand, in the field of investigating ancient environmental gases, much attention has been paid to a gas trapped in the bubbles of ice at the polar regions. A conventional method for studying this uses a special chamber that collects the gas by crushing ice for a CO$_2$ gas [9] and by melting ice for a CH$_4$ gas [10]. But, in both cases, the chamber is sizable and takes considerable time to extract the gas. If the chamber is eliminated, the system will be compact. In addition, if a method to collect the data of the gas directly from the bubble is devised, the data acquisition time will be shortened. Moreover, if the system has a laser-targeting system towards the bubble, it will be a very convenient system. A detection system with those properties will be useful for on-site measurements where ice samples are obtained.

For that purpose, a compact laser-based detection system with a short laser path length is developed with near-infrared semiconductor lasers of 1.65 and 2 μm. The detection sensitivity is enhanced by wavelength-modulation spectroscopy (WMS) [11]–[14]. With this system, the absorption peaks of the isotopic $^{12}$CO$_2$ component of a 3-5% CO$_2$ gas are detected at 2 μm with a path length of 2.4 m, where its peak amplitude is two orders of magnitude smaller than that of the ordinary $^{12}$CO$_2$ component. Also, the absorption spectrum of the $^{12}$CO$_2$ component of a 385-ppm (0.0385-5%) CO$_2$ gas is measured at 2 μm in the air with a path length of 1.4 m. Furthermore, the absorption peaks of a trace gas of pure CH$_4$ in a minuscule area (~1 mm$^3$) are obtained at 1.65 μm with a path length of 3 mm. The details of the constructed system are described in the next section.

II. DEVELOPED SYSTEM

Fig. 1 Experimental setup.

A schematic diagram of the developed system with WMS is depicted in Fig. 1. WMS is a method that can extract derivative signals from faint absorption peaks. To perform WMS, a sawtooth-type voltage change with a sinusoidal voltage modulation generated from a function generator is employed, where the sawtooth-type voltage change has a period of $t_0 = 0.1$ s (or a frequency of $f_0 = 10$ Hz) and the sinusoidal voltage modulation has a period of $t = 1.0 \times 10^{-4}$ s (or a frequency of $f = 10$ kHz). $f$ is set at 10 kHz so that the derivative signals can be obtained clearly. This voltage modulation is then sent to a laser diode (LD) driver to convert it into a modulated current. This modulated current is injected into a 2-μm InGaAs LD [15]–[17] with a maximum power of 2 mW to transfer the same modulation to the laser beam.

The sawtooth-type change with $f_0 = 10$ Hz in the LD injection current works as wavelength-sweeping because it causes an LD refractive-index change and thus a wavelength...
shift. Here, the maximum amount of the injection current determines an attainable wavelength-sweeping range (∼1 nm).

The modulated laser beam from LD passes through a multi-path gas cell with a path length of 2.4 m and reaches an InGaAs photodiode (PD) detector. The detected signals at PD are then fed into a lock-in amplifier. This lock-in amplifier extracts the 2f component of the signals with low noise, which is equivalent to the 2nd derivative component of the signals. Finally, the extracted signals are sent to an oscilloscope to display the results every 0.1 s. In this way, faint absorption peaks are obtained as derivative signals. This is the merit of using WMS. Note that in addition to small semiconductor LD and PD, compact measurement instruments were employed here such that they were packed into a box of 450 mm × 160 mm × 380 mm (roughly the size of a traveling suitcase).

III. EXPERIMENTAL RESULTS AND DISCUSSION

A. Absorption-Spectrum Measurement of CO$_2$

As a test gas, a 3-% CO$_2$ gas at 100 Torr and 25 deg C was used. This concentration is almost equal to that of a CO$_2$ gas in our breath exhaled to the atmosphere. Here, the composition ratio between $^{13}$CO$_2$ and $^{12}$CO$_2$ was set to be the same ratio in the air. As the rest (97%) of the gas, it was filled with N$_2$.

The laser beam generated from LD was set at around 2.006 μm. For quick identification of $^{13}$CO$_2$ peaks, the relative peak positions of $^{13}$CO$_2$ to those of $^{12}$CO$_2$ were examined in advance by use of HITRAN [18], as shown in Table I, where $\lambda$ [μm] and $\kappa$ [cm$^{-1}$] are the laser wavelength and wavenumber, respectively. Since obtainable data in experiments are the transmission rates of a laser beam passing through the mixture of $^{12}$CO$_2$ and $^{13}$CO$_2$ with a path length of 2.4 m at 100 Torr and 25 deg C, the transmission rate was calculated at the same condition with MOLSPEC [19], which is illustrated in Fig. 2.

Next, we show experimental results: The slope of Fig. 3(a) depicts measured, bare signals at the InGaAs PD detector before the extraction of the 2f signals. Fig. 3(a) exhibits only a big absorption peak corresponding to No. 3 in Table I, which is the main peak at this wavelength region. The sinusoidal modulation thickens the measured curve. But when it was turned off, the other peaks than No. 3 were still obscure.

The 2f signals were then extracted from the above bare signals with the lock-in amplifier. In this case, all clear No. 1−5 peaks were observed, as displayed at the curve of Fig. 3(b). This No. 4 peak is the isotopic $^{13}$CO$_2$ peak, which is about 1/200 smaller in size than the No. 3 main peak. From the

### Table I

<table>
<thead>
<tr>
<th>No.</th>
<th>Molecule</th>
<th>$\lambda$ [μm]</th>
<th>$\kappa$ [cm$^{-1}$]</th>
<th>Spectral strength [a.u.]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>$^{12}$CO$_2$</td>
<td>2.006683</td>
<td>4984.129845</td>
<td>3.158 × 10$^{-23}$</td>
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<tr>
<td>2</td>
<td>$^{12}$CO$_2$</td>
<td>2.0062970</td>
<td>4984.306792</td>
<td>2.962 × 10$^{-23}$</td>
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<tr>
<td>3</td>
<td>$^{12}$CO$_2$</td>
<td>2.0062085</td>
<td>4984.526883</td>
<td>1.033 × 10$^{-21}$</td>
</tr>
<tr>
<td>4</td>
<td>$^{13}$CO$_2$</td>
<td>2.0060544</td>
<td>4984.909710</td>
<td>5.203 × 10$^{-24}$</td>
</tr>
<tr>
<td>5</td>
<td>$^{12}$CO$_2$</td>
<td>2.0058833</td>
<td>4985.334836</td>
<td>2.752 × 10$^{-23}$</td>
</tr>
</tbody>
</table>

Fig. 2 Calculated transmission rate for the 3-% mixture gas of $^{12}$CO$_2$ and $^{13}$CO$_2$ with a path length of 2.4 m at 100 Torr and 25 deg C.

Fig. 3 (a) Measured bare signals for the mixture gas of $^{12}$CO$_2$ and $^{13}$CO$_2$, (b) 2f signals extracted from the measured bare signals (a).

Fig. 4 (a) Measured bare signals for $^{12}$CO$_2$ with a concentration of 385 ppm (0.0385%), (b) 2f signals extracted from the measured bare signals (a), (c) Calculated transmission rate for the 385-ppm $^{12}$CO$_2$ gas.
viewpoint of transmission rates, the developed system has an ability to measure absorption peaks for a transmission rate of 99.5% (i.e., an absorption rate of 0.5%), as seen for the No.4 peak in Fig. 2 and Fig. 3(b).

Next, shown at the slope of Fig. 4(a) are measured, bare signals at around 2.011 μm for a 385-ppm (0.0385-%) 12CO2 gas in the air at 760 Torr and 25 deg C with a path length of 1.4 m, where the sinusoidal modulation was turned off. Although Fig. 4(a) displayed no absorption peaks, after the extraction of the 2f signals, a large peak was observed, as displayed at the curve of Fig. 4(b). We can confirm that this peak matches that at 2.0115 μm with a transmission rate of 99.5% (i.e., an absorption rate of 0.5%), as shown in Fig. 4(c) calculated by MOLSPEC.

B. Absorption-Spectrum Measurement of CH4

Next, the absorption spectrum of a CH4 gas in a minuscule area was measured, which could not be obtained by the ordinary wavelength-sweeping method.

As a sample to measure, having in mind investigating an ancient environmental CH4 gas trapped in a bubble of ice at the polar regions, we prepared a small glass container enclosing the gas with an interior volume of only ∼1 mm3. This was made in such a way that both sides of a 1-mm-long hollow glass tube was heat-sealed at 1000 deg C with two 1-mm-thick glass caps in a pure CH4 atmosphere at 760 Torr (Even at this high temperature, pure CH4 was not ignited because of no oxygen). Due to this heat-sealing, as the sample was cooled down to room temperature (25 deg C), the pressure of the enclosed gas would be decreased. The value of this decreased pressure will be estimated later in this section.

To measure the absorption spectrum of the CH4 gas effectively, the 2-μm LD was changed to a 1.65-μm LD [17], and the sample was set between the laser and the detector with a separation of 3 mm.

In addition, to check the peak positions of the CH4 gas for easier measurements, the data of them were obtained in advance by using a much longer glass cell with the ordinary wavelength-sweeping method, where the cell contained a pure CH4 gas with 90 Torr at 25 deg C and was set in place of the multi-path gas cell in Fig. 1. Fig. 5(a) is the target absorption peak in this case, and Fig. 5(b) is the calculated transmission rate for the 20-cm-long cell. Here, we can see that the target peak actually has two peaks at 90 Torr. Fig. 5(c) shows the measured data. From the shape of the observed peaks that fits that of the calculation, wavelength data can be precisely given to the observed peaks.

The slope of Fig. 6(a) shows measured, bare signals through the 1-mm3 CH4 gas in the sample after the replacement of the 20-cm-long cell, while pointing the laser beam towards the small hole of the sample filled with CH4. But as long as the ordinary wavelength-sweeping method was used, no absorption peaks were observed, as depicted at the slope of Fig. 6(a). By the extraction of the 2f signals, two peaks were clearly observed, as shown at the curve of Fig. 6(b), whose peak positions fit those of Fig. 6(c) (the same one in Fig. 5(c)).

Finally, from those two observed peaks, we estimated the aforementioned decreased pressure, P, of CH4 in the sample that was cooled down to room temperature (25 deg C). To do this, the transmission rate through the 1-mm3 CH4 gas was computed in advance by changing P, as illustrated in Fig. 7. This indicates that a broad spectral line at 760 Torr becomes narrow as P decreases and that two peaks start to appear at around 200 Torr. From Fig. 7, P is estimated to be 100 < P < 200 Torr. A more precise estimation of P can be made by Boyle-Charles’ law, PV = nRT, where V is the gas volume, n is the number of moles of the gas, R is the gas constant (or the product of Boltzmann’s constant and Avogadro’s constant), and T is the absolute temperature. Using
Fig. 7 Calculated transmission rate for the 1-mm$^3$ CH$_4$ gas at 25 deg C when the pressure is changed.

the heat-sealing temperature, 1000 deg C, at 760 Torr, we can obtain $P$ at the room temperature, 25 deg C, for the same volume and the same number of moles as $P = 760 \text{Torr} \times \left[\frac{(25 + 273.15) \text{ K}}{1000 + 273.15 \text{ K}}\right] \approx 178 \text{ Torr}$, which is an in-between value of 100 and 200 Torr.

IV. SUMMARY

Constructing a compact laser-based detection system with WMS, we detected the absorption peaks of environmental gases, $^{12}$CO$_2$, $^{13}$CO$_2$, and CH$_4$, using near-infrared semiconductor lasers with a short path length. Here, WMS helped to find faint absorption peaks by extracting their 2$^s$ signals (i.e., 2nd derivative signals) with low noise. The detection sensitivity reached that for a laser transmission rate of 99.5 % (i.e., a laser absorption rate of 0.5 %). With this sensitivity, isotopic $^{13}$CO$_2$ of a 3-% CO$_2$ gas was measured at 2 $\mu$m with a path length of 2.4 m. Also, the absorption peaks of $^{12}$CO$_2$ in the air with a concentration of 385 ppm (0.0385 %) were obtained at 2 $\mu$m with a path length of 1.4 m. In addition, measuring the absorption spectrum of a CH$_4$ gas captured in a small area of ice-bubble size was attempted. For a pure CH$_4$ gas enclosed in a $\sim$1-mm$^3$ glass container, its absorption peaks were detected while pointting a 1.65-$\mu$m laser beam towards the small hole of the sample filled with CH$_4$ with a path length of 3 mm, and also its pressure was obtained from the measured data.

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REFERENCES