Continuous in situ measurements of volcanic gases with a diode-laser-based spectrometer: CO₂ and H₂O concentration and soil degassing at Vulcano (Aeolian islands: Italy)

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Abstract

We report on a continuous-measurement campaign carried out in Vulcano (Aeolian islands, Sicily), devoted to the simultaneous monitoring of CO₂ and H₂O concentrations. The measurements were performed with an absorption spectrometer based on a semiconductor laser source emitting around a 2-μm wavelength. The emitted radiation was selectively absorbed by two molecular ro-vibrational transitions specific of the investigated species. Data for CO₂ and H₂O concentrations, and CO₂ soil diffusive flux using an accumulation chamber configuration, were collected at several interesting sampling points on the island (Porto Levante beach- PLB, Fossa Grande Crater – FOG- and Valley of Palizzi, PAL). CO₂/H₂O values, measured on the ground, are very similar (around 0.019 (± 0.006)) and comparable to the previous discrete detected values of 0.213 (Fumarole F5- La Fossa crater rim) and 0.012 (Fumarole VFS – Baia Levante beach) obtained during the 1977–1993 heating phase of the crater fumaroles.

In this work much more homogeneous values are found in different points of the three sites investigated. The field work, although carried out in a limited time window (25th–28th August 2004), pointed out the new apparatus is suitable for continuous gas monitoring of the two species and their ratios, which are important geochemical indicators of volcanic activity, for which other reliable continuous monitoring systems are not yet available.

Background

Continuous monitoring of gases released in volcanic areas, emitted as fumaroles or diffusive degassing from soils, can give an insight for a deeper understanding of the volcanic dynamics and its impact on global climate and human activities [1,2]. The presence of numerous chemical species released in air and their variation vs. time can be related to chemical reactions of magma with host rocks and circulating fluids, occurring during its uprise. CO₂ and H₂O are molecular species which play a major role in volcanic dynamics. In fact, while water vapour can be related to an increasing heat flow caused by magma uprise movements, the volcanic CO₂ is in general very abundant in magma but is less reactive than other gases during the
interaction with aquifers and rocks [3]. The selection of these two species is based on their common use in volcanology and on their potential in providing useful information regarding volcano degassing dynamics. The CO₂ is hence the best indicator of new deep magma supply, because its increase in concentration and flow at ground reflects increased exsolution from deep sources. In addition, since the CO₂ is by far the most abundant gas component in volcanic fluids, the ratio between CO₂ and H₂O closely represent the gas/water ratio in the aquifers, which is sensitive to temperature increase in the deeper systems, which can result, for instance, by rising of hot fluids or magma [4].

Even though in the past years different techniques have been applied to monitoring of volcanic gases, e.g., mass spectrometry and gas chromatography, they are generally performed in laboratory, after collecting sample gases on field. No reliable techniques for long-term, continuous geochemical monitoring with near-laboratory precision do actually exist. Discrete on-site field measurements are generally restricted to CO₂ soil diffusive flux [5]. Remote sensing techniques were developed in the last years for several gas species, such as Differential Optical Absorption Spectroscopy (DOAS) [6], Correlation Spectroscopy (COSPEC) [7], Light Detection And Ranging (LIDAR) [8], Fourier Transform Infrared (FTIR) Spectroscopy [9,10]. Eddy-covariance [11] (EC) micrometeorological method can be alternative solutions, for CO₂ flux, allowing measurements over large gas emission areas. Some of these methods are able to detect few trace species (e.g. SO₂, HCl, HF) while H₂S, CO₂ and H₂O, main constituents of volcanic gases, are spectroscopically hard to determine, therefore the corresponding volcanogenic emission rates can only be derived indirectly. More recently, continuous gas and temperature measurements have been performed using a set of analytical instruments to monitor volcanic gases over weeks [12-14].

Laser spectroscopy has been shown to be a powerful technique for field, non-invasive and real-time gas detection, based on the selective absorption of laser radiation by the molecules of the target gas [15,16]. Most molecules of interest have fundamental and overtone absorption bands in the infrared region of the electromagnetic spectrum. However, in the spectral region below 2 μm the outstanding progress in the telecommunication technologies made available laser sources, like distributed feed-back (DFB) semiconductor lasers, which are ideal for the development of high-resolution, fast-response spectrometers. In previous works we applied this technology to monitor CO₂ and H₂O in volcanic fumaroles, using a dual-wavelength approach [17], delivering a beam with 1.57 and 1.39 μm diode-laser radiation to a 20-cm-long cell placed upon the emitting fumarole by means of optical fibers. The dual-wavelength spectrometer enabled us to measure water vapor concentration with a 3% accuracy, whereas it was quite inaccurate for carbon dioxide measurements [18]. Subsequently, we implemented a new absorption spectrometer using a single room-temperature 2-μm DFB diode laser for real-time measurement of both CO₂ and H₂O concentration [19,20]. At this wavelength, a considerably larger fractional absorption can be detected, due to the occurrence of stronger ro-vibrational transitions. The measurement technique was based on recording of direct absorption profiles of a gas sample contained in a multiple-reflection cell with a path-length of 50 m. Remote operation was still feasible using telecom-type optical fibers. Laboratory tests showed that it was possible to achieve a short-term reproducibility over several tens of minutes and an overall accuracy below 1% for measurements on ambient CO₂. Simultaneous measurements of the CO₂ and H₂O concentrations maintain similar accuracy levels [20,21].

After the first field tests [20], the spectrometer has been upgraded and in this paper, we present the results of a measurement campaign carried out in August 2004 in Vulcano (Aeolian islands, Sicily). Vulcano, due to the long period of geochemical anomalies and seismicity [22,23], and the presence of high CO₂ degassing [24-27], represents an ideal test site for the new spectroscopic apparatus, both for concentration and flux measurements. Further, the presence at some sites of fumaroles with very aggressive (hot and acid) components allow to test the apparatus and optics also in extreme conditions of operation. The volcanic CO₂ discharges occur both from fumaroles and as diffuse soil emanations form the volcanic edifices [28].

**Geological setting of Vulcano fumaroles and soil degassing composition**

The Vulcano Island is one of the most active volcanoes of the Mediterranean area; since its last explosive eruption, which occurred in 1888–1890 from the Fossa crater, the volcano has experienced fumarolic activity, mostly located along the northern rim of this crater [22]. Owing to a potentially pre-eruptive crisis begun in late 1977, numerous papers have been focused during the last 25 years on understanding the volcanic system at Vulcano Island and on volcanic surveillance. Chemical composition changes of the fumarolic gases have been monitored for two crater fumaroles (FA and FI) by Capasso et al. from the 1988 to 1991 [29]. The fumaroles studied showed an increase in temperature, ranging from 590°C to a maximum of 650°C for the FA fumarole. Moreover, the He/N₂, He/CO₂ and He/S ratios showed large variations with a general increase of helium, nitrogen, carbon dioxide and total sulphur contents and indicating a general trend like He>N₂>CO₂>total S. The behaviour of HCl...
concentration in the fumarole FA is opposite respect to those of He, N₂, CO₂, SO₂. In average the variability of the gas composition at Vulcano have been explained [23] as an indicator of different degree of mixing processes between a Deeper Magmatic Component (DMC) and a Shallow Hydrothermal Component (SHC). Taking into account both non-reactive gases (e.g., N₂, and He) and main constituents (i.e., H₂O, and CO₂), the maximum contribution of the DMC occurred in the summer of the 1988 while the maximum fraction of the SHC in fumarolic fluids was recorded through 1989. The following progressive increase in CO₂ concentration through 1990 indicates an increasing DMC contribution, likely due to an energetic seismic event, which should have caused intense fracturation and permeability increase; it was followed by its decrease during the 1991, and finally followed, during the 1992, by an increase of the DMC contribution [23]. During the 1993–1995 period [30] monitoring of three types of sampled manifestations at Vulcano (Crater fumaroles, Baia di Levante and SIP, Palizzi and P4MAX) showed the occurrence of mixing between different end-members (crater hottest fumaroles, cooler shallower fluids, atmospheric component).

Gurrieri and Valenza have reported a diffusive CO₂ gas monitoring at Vulcano island inserting in the soil a pipe opened at the base and measuring the dynamic concentration of the CO₂ which is proportional to the ϕCO₂, unless a conversion factor, depending on the experimental device, working conditions, physical characteristics of the soil [31]. Since the end of 1984, several campaigns have been carried out using the latter method in the area of Vulcano Porto and in the area surrounding the cone of La Fossa [32,33], reporting values of flux as high as 8·10⁻⁴ cm/s. Carapezza and Diliberto [34] have measured helium and CO₂ soil degassing respectively using a mass spectrometer in the lab and using an IR spectrometer, directly in field, with an accuracy for CO₂ concentration of 100 ppm. The average concentration of CO₂ measured in the 1989–1991 in fifty-one different points shows a general increasing trend with fluctuations, from 1989 to the end of 1991, with a maximum value at the end of 1990. Moreover during this campaign it was evidenced that some points, located on the southern base of the cone (Grotta Palizzi) and on the W and NW cone slope (Forgia Vecchia) show higher values than those found in Vulcano Porto area. CO₂ fluxes measured since the 1989 [33] suggest an increasing trend since the end of 1989 (~9·10⁻⁴ cm/s per area unit) to the end of 1991 (~4·9·10⁻⁴ cm/s per area unit), decreasing during the 1992–1993 period (~1.2·10⁻⁴ cm/s per area unit). It is worth of note that the observed temporal variations are not influenced by seasonal variation of atmospheric parameters. This means that measurements of CO₂ soil degassing can constitute an useful indicator of deep dynamics in case of further evolution of the activity in the island whereas the crater fumaroles get inaccessible. Moreover, the collected data at specific points (Point 4) during the 1990–1992 years interval seem to suggest that a feeding system was active and enriched in He and CO₂, probably connected to La Fossa volcano [35], whereas elsewhere in the surveyed area He concentration was controlled by the extent of mixing between air and exhaling CO₂. Continuous gas monitoring of CO₂ in soils [36] carried out at different areas of the Vulcano island shows that the mean values of CO₂ degassing in the most representative areas were lower than those of the previous year, in agreement with the observation carried out in the area of Vulcano Porto during the monthly surveys [33].

Accumulation chamber methodology used during a campaign in the period April–July 1996 in the upper part of the Fossa cone [25] measured a total output of 200 td⁻¹ of CO₂ which corresponds to approximately 1000 Mgd⁻¹ of steam. In order to obtain the ϕCO₂ in the plains adjacent to the western and southern slopes of the Fossa volcanic edifice and in the isthmus separating Porto di Levante from Porto di Ponente, Chiodini et al. [26] measured emissions from 0.2 to 2900 gm⁻² d⁻¹ for a total amount of diffuse CO₂ output of 75.3 Mgd⁻¹. A survey on fifteen measurements of ϕCH₄, ϕCO₂ and temperature was conducted at Porto di Levante [24] obtaining respectively: 0.007 to 3.9 g m⁻² d⁻¹, 6.4 to 12000 gm⁻² d⁻¹, and 20 to 100°C. Carapezza and Granieri [37] have proposed a comparison between active Dynamic Concentration and passive Accumulation Chamber methods to measure the carbon dioxide soil flux at the La Fossa areas of Vulcano Island. This work showed that the active methods overestimate the CO₂ flux, and the results are proportional to it only in high-flux zones. Moreover, the carbon dioxide soil flux values were controlled in proximity of active gas-releasing fractures and in presence of changes in barometric pressure and soil permeability.

**Experimental setup**

The Portable Diode Laser Spectrometer (PDLS) is based on a diode laser emitting infrared radiation around 2 μm, where different ro-vibrational transition of CO₂ and H₂O occur. The PDLS has been previously described and char-
acterized in details in [19-21]. Here we recall the principle of operation and the major features, mainly discussing the changes made in order to improve both the robustness of the system as well as its compactness, as required for a field instrument.

The spectrometer is made by three parts: the laser breadboard, the detection bench and the electronics. A 30 × 40 cm² ultra-light breadboard (10-kg-weight) hosts a DFB semiconductor laser (Sensor Unlimited) emitting at 1997 nm. The emitted radiation is collimated and coupled to a 30-m-long single-mode optical fiber by means of two steering mirrors and a input lens which focus the light on one fiber end. In this way we are able to steady couple about 10% of the incoming power.

With respect to the previous version of the spectrometer a factor 3 in the weight of the laser breadboard has been gained. The use of the fiber allows to keep the laser and the controlling electronics in a safe place, far from the sampling point, where environmental conditions can often be rather hostile and compromise a normal operation. The radiation is guided by the optical fiber to the detection bench, where it is collimated and injected by lenses and tilting mirrors into an open-path multipass Herriott cell. This cell is made by two gold-coated spherical mirrors, 50-mm-diameter and ~400-mm curvature radius, rigidly aligned and bolt 400 mm apart in the central part of the detection bench. All the lenses were coated with anti-reflective layer for reducing the occurrence of fringes which could deteriorate the absorption profile. A rectangular aperture (30 × 30 cm²) at the bottom of the bench allows the sampled gas flowing between the mirrors. The total optical path of the cell is 20 m long. The radiation transmitted by the cell is finally detected by a preamplified extended InGaAs detector. Also the detection bench has been modified, as a consequence of the experience for the previous campaigns (2002–2003) [20]: all the external surfaces are made of a titanium alloy, but also more resistant to acid compounds emitted in volcanic areas. As a result of experience of a previous field campaign (2002–2003) [20], the system as well as its compactness, as required for a field instrument.

The electronics and a laptop computer (LC) operate the laser, acquire the absorption signals and process them in real time. The laser temperature is actively stabilized providing a coarse tuning of its emission wavelength, then the frequency of the laser is continuously scanned by sweeping its injection current with a triangular current ramp (2-Hz scan rate) around an offset constant value. The scanned frequency range is chosen in such a way to probe simultaneously the two absorption molecular transitions of CO₂ and H₂O. The laser frequency is swept over a range of ~20 GHz which includes the two absorption features of CO₂ (line R(34) of the ν₁ + 2ν₂ + ν₃ combination band) and H₂O (line 15ₐ₀₁₅ → 1₄₀₁₄ of the ν₂ + ν₃ band). The absorption spectra are acquired by means of a multifunction PCMCIA digital acquisition board (National Instruments NI DAQCard-6036E for PCMCIA), with a 16-bit analog-to-digital converter (ADC), at a maximal acquisition rate of 200 kS/s, interfaced to the LC. The temperature of the gas is also monitored in real time, with accuracy of 0.1 °C, by acquiring the signal of a thermocouple via one of the ADC channels. A single absorption signal is acquired at the frequency scan of 2 Hz, so concentration values can be obtained every 0.5 s. Otherwise, it is possible to average several signal before processing the spectra and retrieve the concentration values.

The process is made in real time, even though all the spectra are stored for further off-line analysis. The acquisition program, written in Lab View® language, controls the instrument settings and analyzes the acquired spectra. The fitting procedure described below is made by running a Matlab routine called by the main program.

**Signal analysis and preliminary tests**

If \( P_0(ν) \) is the power emitted by the laser as a function of the frequency \( ν \), then the power \( P(ν) \) detected after passing through the absorbing cell is given, according to the Beer-Lambert law, by

\[
P(ν) = P_0(ν) \exp[-A_{CO₂}g(ν - ν_{CO₂}) + A_{H₂O}g(ν - ν_{H₂O})].
\]

(1)

Here \( g(ν - ν₅₃) \) is the unit-area normalized lineshape function, centred at corresponding line center frequency \( ν₅₃ \) for the transition \( X \), which is assumed to be Lorentzian, as the gas is usually at atmospheric pressure:
where $g_X$ is the Half Width at Half Maximum (HWHM). $A_X$ is the integral area of the absorption feature which is the observable related to the gas concentration $N_X$, according to

$$A_X = -S_X(T) \ln N_X$$  \hspace{1cm} (2)$$

where $S_X(T)$ is the line strength of the transition, which depends on the temperature of the sample through a known relation, and $L$ is the absorption path length. The identification of the chosen spectral lines has been made according the spectroscopical database HITRAN [38], which also excluded the presence of other interfering gas species. The frequency scale has been previously determined by calibrating the time scale of the modulating ramp with a Fabry-Pérot cavity with known free spectral range [19]. Due to the long time stability of the laser source, this calibration has been shown to be stable over several months. Eq. (1) is fitted to the absorption spectrum according to the Levenberg-Marquardt method, and from the knowledge of the parameters resulting from the fit we can calculate the integral area $A_X$ of the absorption and finally retrieving, through Eq. (2), the concentration of the two species, knowing the two experimentally determined transition line strengths, $S_{CO_2} = 4.77(3) \times 10^{-22}$ cm/mol at 301.4 K [19] and $S_{H_2O} = 2.28(4) \times 10^{-24}$ cm/mol at 298 K [21], and the optical path length $L = 20.3$ m. The baseline has been assumed to be a second degree polynomial, as in laboratory condition it has been shown to well reproduce the signal when no absorption is present [20]. This baseline is very stable on a time basis of hours, as it is essentially determined by the increasing of the emitted power as the injection current of the laser increases during frequency scan. No effect of fringes was observed on baseline stability. During the field measurement environmental additional effect (i.e., sudden attenuation of signal from smoke or condensed vapour) could drastically change the baseline, but these cases were uncommon and were easily excluded by monitoring the $\chi^2$ of the fit as a reliability parameter of the fit. Figure (1) shows a typical experimental absorption spectrum recorded during the present campaign, along with the best fit of Eq. (1). In the lower part it is also shown the residual in percentage between experimental signal and fitted curve, normalized with respect to the laser power. The error bar associated to each concentration value in the reported plots are determined by the propagation of the errors given by the fit for each parameters. For each fit the resulting $\chi^2$ is monitored in order to verify the reliability of the results.

A preliminary laboratory survey has been performed on the spectrometer in order to verify its performances as well as the procedure to retrieve the gas concentrations. Several tests were done using certified gas mixtures as reported in [19,21] with more details: the short term reproducibility over several minutes was found to be of 0.3% and 0.5% for $CO_2$ and $H_2O$, respectively. Long term measurements (24 hours) were also performed. The dynamic range of the spectrometer was fixed in the interval 30 – 10000 ppm, as mainly limited from the maximum acceptable relative uncertainty of 10% for the retrieved concentration (lower limit), and from the maximum frequency scan to be imposed on the laser (fixed to 1 cm$^{-1}$) (upper limit) in order to keep the tails of the absorption within.

A comparison between the performances of the laser spectrometer and a commercial infrared gas analyzer (IRGA) was also performed [19]. The robustness of the fitting procedure was verified with different values of the initial parameters values, as well as with Levenberg-Marquardt damping factors. Once the convergence in the fit procedure is achieved, the values of the retrieved concentration resulted not seriously biased
neither by the initial values of the parameter, nor by the damping factors. Conversely the right choice of both these parameters turned out to be crucial in the number of iterations necessary to achieve convergence.

Field measurements
The measurement campaign was held in August 25th-29th, 2004 at Vulcano island. During the campaign we analysed the emissions in different sites: PLB, PAL, and FOG, Fig. 2. These sites were choosen taking into account the previous studies [22-27,29,30,32-37,39-41] of the volcanic activity on Vulcano, identified as the extensive fumarole field in the northern part of La Fossa crater (100°C < T < 600°C), low-temperature fumaroles (T ≈ 100°C) in the Baia di Levante area, and widespreaded manifestations in the soil in the Vulcano Porto area and around the volcanic cone in the Palizzi valley. We performed measurements with two different configurations, with and without the breadboard cover, depending on the type of emissions we found and on the environmental conditions.

The first configuration was without the breadboard top cover, with the possibility of placing it at different heights from the ground. The second configuration was with the breadboard placed at the ground and with the top cover closed, in such a way to form an inverted accumulation chamber allowing the gas to flow from the soil to the sensing cell, through the aperture of the breadboard. The sealing of the base is ensured by the slight burying of the breadboard in the soil. The latter was used when (1) there was no appreciable difference between the CO2 concentration and the average background value, if measured with an open configuration, and (2) there was low water vapour emission. Indeed, when concentration of water vapour in the accumulation chamber is so high that condensation of liquid water occurs on the colder surface of the mirrors and lenses, the final power on the detector shows strong attenuation, decreasing the signal to noise ratio and finally the sensitivity. The time to determine the flux is comparable with that of other accumulation chamber. However the sampled air is measured directly at the surface were the flux occurs, with no need of move the air with pumps to the measurement cell.

Gas concentrations
Gas concentration measurements at Vulcano island have been performed at the three main sites (Fig. 2), namely PLB, FOG crater and PAL, and relevant data are resumed in Table 1. Different configurations were exploited.

Table 1 resumes the mean values and errors for CO2, H2O and their ratios at the different sampling points. Measurements at PLB (August 25th, 18:30 local time) have been performed setting the apparatus at various heights over the ground (PLB1 at 0 cm; PLB2 at 40 cm; PLB3 at 130 cm), in order to estimate the influence, on CO2 and H2O concentrations, due to the distance of the breadboard from the ground (Fig 3). Values of CO2 concentrations range from 1500 to 450 ppm, with average values measured decreasing from 779 ppm at ground to 450 ppm at 130 cm of height. The average value is rather stable from 44 to 130 cm, indicating that we are measuring, at these heights, almost the gas concentration in the atmosphere. H2O concentration is, on average, of about 4% at ground, slightly decreasing to 3.5% at 130 cm of height. We took also a measurement at 20 cm from the ground which however presented very noisy spectra, resulting in unreliable concentration values greater than those measured at ground level, which we cannot explain. Examples of measurements obtained at FOG (August 27th, 16:00) are shown in Fig. 4. At this site, four different points, at distances of few meters each other, have been sampled, in order to have an idea of the variability of results within small distances. Average values for CO2 concentrations range from 564 ppm to 481 ppm, whereas H2O ones are between 2.9% and 3.5% as shown in Table 1. At PAL site (August 26th, 17:00), three points within distances of some meters where selected for measurements, and examples are shown in Fig. 5 while results are reported in Table

Figure 2
DTM (Digital Terrain Model) of the Vulcano island with the indication of the main measurement sites: PLB: Porto Levante Beach; FOG: Fossa Grande Crater; PAL: Valley of Palizzi.
Average values for CO$_2$ concentrations lie in the range 384 to 564 ppm while for H$_2$O, they are between 2.3% and 2.6%.

Flux measurements with accumulation configuration

In the accumulation chamber configuration, the rate of change of the measured concentration can be modeled by the following differential equation,

$$\frac{dC_c(t)}{dt} = \alpha[C_s - C_c(t)]$$

where $C_c$ and $C_s$ are the concentration of the CO$_2$ in the chamber and in the soil, respectively, and $\alpha$ is the flux rate. Assuming a concentration $C_c(0)$ at the time $t = 0$

| Site                          | T/°C | Mean CO$_2$/ppm (error) | st.dev. | Mean H$_2$O/ppm (error) | st.dev. | Mean CO$_2$/H$_2$O (error) | st.dev. |
|-------------------------------|------|-------------------------|---------|-------------------------|---------|-----------------------------|---------|
| Porto Levante Beach (2/8/04) |      |                         |         |                         |         |                             |         |
| PLB1                          | 26   | 769 (19)                | 240     | 39700 (400)             | 5600    | 0.0195 (5)                  | 0.0058  |
| PLB2                          | 26   | 477 (13)                | 122     | 36200 (400)             | 3500    | 0.0132 (4)                  | 0.0034  |
| PLB3                          | 26   | 450 (12)                | 115     | 34500 (400)             | 3600    | 0.0131 (3)                  | 0.0032  |

| Palizzi (26/8/04)             |      |                         |         |                         |         |                             |         |
| PL1                           | 26   | 384 (3)                 | 40      | 25540 (80)              | 1140    | 0.0150 (1)                  | 0.0016  |
| PL2                           | 26   | 468 (3)                 | 44      | 25830 (75)              | 1050    | 0.0182 (1)                  | 0.0018  |
| PL3                           | 24   | 564 (12)                | 120     | 23130 (75)              | 740     | 0.0244 (5)                  | 0.0053  |

| Fossa Grande (27/8/04)        |      |                         |         |                         |         |                             |         |
| FOGL                          | 25   | 481 (24)                | 170     | 29000 (1000)            | 7500    | 0.0174 (12)                 | 0.0090  |
| FOG2                          | 26   | 564 (15)                | 120     | 34300 (500)             | 3400    | 0.0165 (4)                  | 0.0031  |
| FOG3                          | 29   | 541 (22)                | 200     | 35200 (2000)            | 19600   | 0.0195 (5)                  | 0.0058  |
| FOG4                          | 26   | 505 (20)                | 173     | 30200 (900)             | 8000    | 0.0178 (3)                  | 0.0074  |

1. Average values for CO$_2$ concentrations lie in the range 384 to 564 ppm while for H$_2$O, they are between 2.3% and 2.6%. 

Figure 3

Two examples of concentration measurement at different height from the ground, 0 cm (left), 40 cm (right), made at the PLB (Aug 25th 2004) and corresponding to acquisition PLB1 and PLB3 of Table 1.
when the chamber is closed, the concentration in the chamber will change according to

\[ C_c(t) = C_s - (C_s - C_c(0))e^{-\alpha t}. \]  

By fitting the last expression to the experimental data we can retrieve the flux rate \( \alpha \) and the concentration \( C_s \). Examples of fitting procedure are shown in Fig. 6. The retrieved fit parameters are listed in Table 2. Knowing the volume of the chamber and the open area at base of the chamber, we can estimate the mass flux at the soil, as reported in the last column of Table 2.

Measurements of \( \text{CO}_2 \) flux at three sites of PAL (August 28th, 18:00) resulted in rates of 0.0031, 0.00090, 0.0018 s\(^{-1}\) (Table 2). At the first site, in a measurement took two days before (August 26th, 17:00), we get a rate of 0.0017 s\(^{-1}\) (Fig. 6). The concentration of water vapour remained constant during each flux measurement and its value is the same in the three different measurements of day August 28th, while for the two measurements made in the same place but in two different days (August 26th and 28th) we observed a change in water vapour concentration. This suggests that the presence of water is merely due to humidity of air. A measurements at Porto Levante Beach (August 29, 11:00) gave a flux rate of 0.0017 s\(^{-1}\). During this measurement we observed an increase of water vapour concentration, however we believe it is likely due to the evaporation of water soil content due to sun at the measuring site after raining. Flux measurements at FOG, at distance of few meters from the main fumarole, showed not detectable flux (Table 2). Similar accumulation measurements in the fumarole failed for the condensation of water vapour on the mirrors which reduced the signal to zero.

**Conclusion**

We have carried out a measurement campaign in Vulcano island with a portable spectrometer based on a semiconductor laser, which can monitor in-situ concentration and flux of \( \text{CO}_2 \) and \( \text{H}_2\text{O} \). The spectrometer has a response time of the order of one second which allows the monitoring of gas dynamics on a very fast time scale. The measurements obtained at Vulcano, together with several tests, performed continuously for several days, indicate the system has no problem to be used for long-time unattended
Figure 6
Flux measurement at PAL – Aug 26th (above) and 28th (below) 2004. Solid lines represent the best fit of Eq. (3) to experimental data. Fluxes corresponding to data PAL Fl and PAL F2 of Table 2.
measurements as well. The only occasional problems for a long-term remote installation could be the condensation of water vapour on the mirrors; such problem could be be reduced by heating the mirrors’ surface. The average values of the CO2/H2O at ground level of the different measurement sites (0.0195–0.0150) is consistent with an average value measured for example at the FO fumarole (see the text for details).

### Table 2: Accumulation measurements. Concentration rate $\alpha$ and saturation values $C_s$ retrieved by fitting Eq. (3) to the experimental data. In the last column we list the respective flux of CO2 (at Standard Temperature and Pressure). The sites were labelled in the first column (see the text for details).

| Site          | $T/^\circ$C | $\alpha/10^{-3}$s$^{-1}$ | $C_s$/ppm | flux/g m$^{-2}$s$^{-1}$ |
|---------------|-------------|--------------------------|-----------|-------------------------|
| Palizzi (26/8/04) |             |                          |           |                         |
| PAL F1        | 24          | 1.65 (4)                 | 8390 (60) | 1.73                    |
| Palizzi (28/8/04) |             |                          |           |                         |
| PAL F2        | 36          | 3.06 (6)                 | 3140 (10) | 3.20                    |
| PAL F3        | 40          | 0.90 (6)                 | 6750 (270)| 0.94                    |
| PAL F4        | 34          | 1.82 (10)                | 12950 (440)| 1.90                   |
| Porto Levante Beach (28/8/04) |             |                          |           |                         |
| PLB F1        | 33          | 1.72 (6)                 | 28330 (510)| 1.80                   |

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