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A compact and stable eddy covariance set-up for methane measurements *using off-axis integrated cavity output spectroscopy*

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Abstract

A Fast Methane Analyzer (FMA) is assessed for its applicability in a closed path eddy covariance field set-up in a peat meadow. The FMA uses off-axis integrated cavity output spectroscopy combined with a highly specific narrow band laser for the detection of CH₄ and strongly reflective mirrors to obtain a laser path length of 2-20×10³ m. Statistical testing and a calibration experiment showed high precision (7.8×10⁻³ ppb) and accuracy (< 0.30%) of the instrument, while no drift was observed. The instrument response time was determined to be 0.10 s. In the field set-up, the FMA is attached to a scroll pump and combined with a 3-axis ultrasonic anemometer and an open path infrared gas analyzer for measurements of carbon dioxide and water vapour. The power-spectra and co-spectra of the instruments were satisfactory for 10Hz sampling rates.

Due to erroneous measurements, spikes and periods of low turbulence the data series consisted for 26% of gaps. Observed CH₄ fluxes consisted mainly of emission, showed a diurnal cycle, but were rather variable over time. The average CH₄ emission was 29.7 nmol m⁻² s⁻¹, while the typical maximum CH₄ emission was approximately 80.0 nmol m⁻² s⁻¹ and the typical minimum flux was approximately 0.0 nmol m⁻² s⁻¹. The correspondence of the measurements with flux chamber measurements in the footprint was good and the observed CH₄ emission rates were comparable with eddy covariance CH₄ measurements in other peat areas.

Additionally, three measurement techniques with lower sampling frequencies were simulated, which might give the possibility to measure CH₄ fluxes without an external pump and save energy. Disjunct eddy covariance appeared to be the most reliable substitute for 10Hz eddy covariance, while relaxed eddy accumulation gave reliable estimates of the fluxes over periods in the order of days or weeks.

4.1 Introduction

Methane is the third most important greenhouse gas globally, after water vapour and carbon dioxide. Its concentration has risen by 150% since the pre-industrial era (Foster et al., 2007) and currently 20% of the enhanced greenhouse effect is considered to be due to methane (Foster et al., 2007). Although methane is less abundant in the atmosphere compared to carbon dioxide, it is a relatively strong greenhouse gas: the Global Warming Potential of methane expressed as CO₂-equivalents is approximately 25 (over 100 years) while that of carbon dioxide is by definition 1 (Foster et al., 2007). For water vapour and carbon dioxide, the eddy covariance technique has been used for several decennia (Aubinet et al., 2000) to measure the exchange of these gases with the earth surface. In eddy covariance, the vertical flux (F_s) of an atmospheric property (s) is directly determined by the covariance of that property and the vertical velocity, as shown in Eq. (4.1). This can be obtained by calculating the time averaged product (over the period t_1 to t_2) of the deviation (s') of the atmospheric property (s), from $s = \bar{s} + s'$, and the deviation (w') of the vertical wind velocity (w) from $w = \bar{w} + w'$:

$$F_s = \overline{w' s'} = \frac{1}{t_2 - t_1} \int_{t_1}^{t_2} w'(t) s'(t) dt \quad (4.1)$$

(Aubinet et al., 2000). The eddy covariance technique requires an instrument with high precision, accuracy and system stability as well as high sampling rates and short instrument response time (τ). Unfortunately, the low abundance of methane in the atmosphere hampers adequate concentration measurements of this gas and the eddy covariance technique has therefore been rarely applied for the assessment of methane emissions (Kroon et al., 2007; Hargreaves et al., 2001; Kormann et al., 2001). The advantages of the eddy covariance technique compared to other techniques for measuring trace gases are nonetheless obvious: integrated continuous measurements over a large footprint area (10^2 to 10^4 m²) and longer periods without disturbance from small scale surface features. These properties enable assessments of spatial and temporal variability at the landscape scale.

Since the 1980s, infrared absorption spectrometry using tunable diode lasers (TDLs) has been widely used for measurements of trace gases in the lab. A field technique for eddy covariance using a multipass absorption cell was introduced in 1995 (Zahniser et al. 1995). Unfortunately, serious problems of drift and low sensitivity effects occurred. Additionally, the method had more practical drawbacks: the large nitrogen Dewar for temperature control that needs refill weekly, an extensive and sensitive optical module, and frequent calibration. Improvements were made with the Quantum Cascade Laser (QCL) spectrometer which was first introduced in 1994 (Faist et al., 1994). The method was more stable and accurate in eddy covariance set-up than the TDL spectrometry (Nelson et al., 2004; Kroon et al., 2007), but the practical drawbacks of the methane instruments (large nitrogen Dewar, optical module, and frequent calibration) were still present in the QCL spectrometer technique. Hitherto, the micrometeorological

measurement techniques for methane have thus implied very expensive, large and labour intensive field set-ups with logistic limitations to unattended field deployment.

The objective of this study was to investigate the applicability and quality of the FMA, a new off-axis integrated cavity output spectroscopy (ICOS) technique using a highly specific narrowband laser, for its applicability for eddy covariance field measurements of methane. The relatively user-friendly and low cost set-up is tested for precision, accuracy and system stability as well as its' performance in eddy covariance field set-up. The first data series are analysed and compared with data obtained by existing measurement techniques. To assess the applicability of the instrument at locations that lack external power supply, alternative measurements techniques of the set-up are simulated which could reduce the power requirements.

4.2 Methods and materials

4.2.1 Instrument design

A measurement cell with highly reflective mirrors was combined with a highly specific narrowband laser system by O'Keefe et al. (1998), creating a path length of at least two km in the measurement cell. In this manner, small absorption rates cause much larger reduction of the total transmitted laser intensity. The ICOS technique could therefore be used for the detection of gases with ultra weak absorption, while making the extensive optical module and the nitrogen Dewar superfluous. The operation of the DLT-100 Fast Methane Analyzer (FMA) from Los Gatos Research Ltd. is based on this improved ICOS technique and uses a distributed feedback (DFB) diode laser with a wavelength of nearly 1.65 μm . The DFB diode laser offers tunability, narrow line width and high output power in a compact and very rugged setup. It features a grating structure within the semiconductor, which narrows the wavelength spectrum and guarantees single-frequency emission. Off-axis ICOS implies that a laser beam directed into the measurement cell at a slight angle, after which it is reflected in the cell numerous times by highly reflective mirrors (reflectivity ~ 0.9999), thus creating a path length of $2\text{-}20 \times 10^3$ m by making $1\text{-}10 \times 10^4$ passes in the cell (Bear et al., 2002; Fig. 4.1). The detector measures fractional absorption of light at the methane resonant wavelength, which is an absolute measurement of the methane concentration in the cell.

The path length of the laser, and therefore the time period during which the laser is being reflected in the cell for each measurement, is dependent on the reflectivity of the mirrors in the measurement cell. This period over which the laser is being reflected in the measurement cell is called the mirror ringdown time (MRT) and is continuously monitored by the FMA. The MRT cannot be allowed to drop below 3.0 to 3.5 μs , since the laser path length then becomes too short to detect the changes in laser intensity properly. The mirrors in the measurement cell are sensitive to dirt accumulating in the measurement cell; a small contamination of the measurement cell causes rapid decrease of the MRT. Cleaning the mirrors is a relatively simple procedure that can be done in a dust-free environment.

The FMA measures in the concentration range from 10 to 25×10^3 ppb and operates autonomously. Technically, measurements can be made at rates up to 20Hz and at ambient temperatures of 5 °C to 45 °C, while humidity should be below 95% to avoid condensation. The pressure in the measurement cell (P_{cell}), which can be adjusted by a valve switch on the instrument, should be kept near 210 hPa. To obtain sampling rates higher than 1Hz, an external pump is needed to maintain the required P_{cell} and τ .

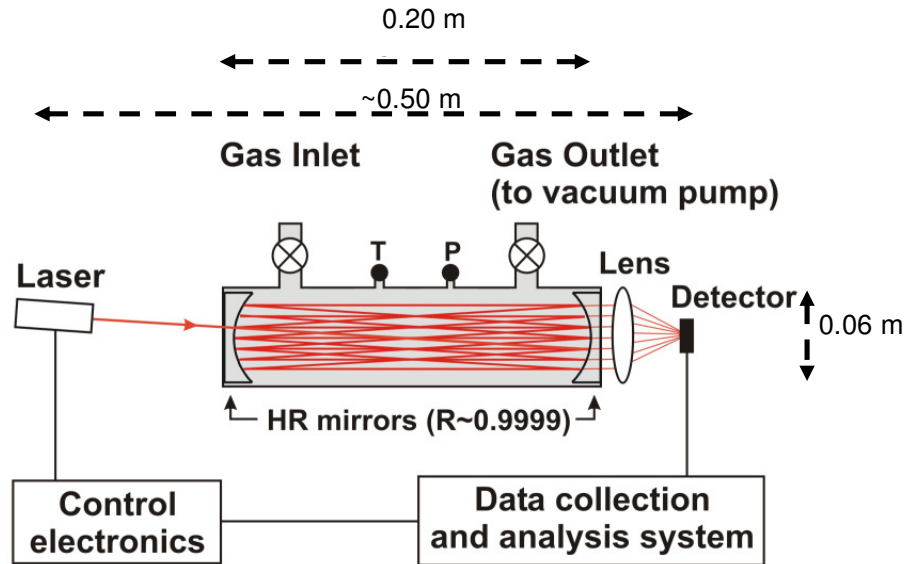


Figure 4.1: Schematic overview of the integrated cavity output analysis (ICOS) technique used in the FMA. (Source: Los Gatos Research Ltd.) “R” is the reflectivity of the mirrors and “T” and “P” are the sensors of T_{cell} and P_{cell} .

The measurement cell has a volume of $0.55 \times 10^{-3} \text{ m}^3$ and a length of 0.20 m (Fig. 4.1). Data output is provided in analogue as well as digital format (RS232&TCP/IP) and the device can store data up to 10 gigabytes. Warm-up time is approximately one minute and measurements as well as performance can be observed on a colour TFT LCD flat panel display. The dimensions of the FMA are 0.25 m height, 0.97 m width and 0.36 m depth and it has a weight of 22 kg. Power requirements are 115/230 VAC, 50/60Hz and 180W and inlet/outlet fittings are of the Swagelok type ($3/8''$, $1/4''$).

4.2.2 Site description

Besides testing in the laboratory, the FMA was tested in an eddy covariance set-up at the Horstermeer measurement site. This site is located in a eutrophic peat meadow area in the central part of the Netherlands and was described extensively by Hendriks et al. (2007). The area has a flat topography and vegetation consists of grasses, small shrubs and reeds. Before, CH_4 fluxes in the area were measured with the flux chamber technique and variation between three land elements was observed: emissions from the saturated land

and water surfaces were high compared to the relatively dry land. The annual weighted CH₄ emission at the site over 2005 and 2006 was estimated at $83.95 \pm 54.81 \text{ nmol m}^{-2} \text{ s}^{-1}$ (Hendriks et al., 2007).

4.2.3 Assessment of instrument stability, precision and accuracy

System stability is a major factor influencing high-sensitivity measurements. Theoretically, the signal from a perfectly stable system could be averaged infinitely. However, real systems are stable only for a limited time period. The length of time over which a laser signal can be averaged to achieve optimum sensitivity, and thus high precision, largely determines the quality of the spectrometer. The precision of concentration measurements should be at least a few parts per thousand of the ambient mixing ratio, which is approximately 4 ppb for a mixing ratio of approximately 1800 ppb (outside air) in the case of CH₄ (Kroon et al., 2007). Both maximum system stability and precision can be determined using the Allan variance (Allan, 1966; Werle et al., 1993; Nelson et al., 2004; Kroon et al., 2007). The Allan variance (σ_A^2), as a function of integration time T , is the average of the sample variance of two adjacent averages of time series of data and is described by Eq. (4.2):

$$\left\langle \sigma_A^2(k) \right\rangle_t = \frac{1}{2m} \sum_{s=1}^m [A_{s+1}(k) - A_s(k)]^2 \quad (4.2)$$

$$\text{with: } A_s(k) = \frac{1}{k} \sum_{l=1}^k x_{(s+1)k+l}, \text{ with: } s = 1, \dots, m \text{ and } m = m' - 1$$

In this equation A is an average of the CH₄ concentration, k is the number of elements in subgroup x , s is the subgroup number, l is the sample-number in the subgroup, and m' the number of independent measurements. It is assumed that data are collected over a constant data interval Δt , therefore the integration time $T = k\Delta t$ (Allan, 1966; Werle et al., 1993; Nelson et al., 2004). The Allan variance decreases when random noise dominates over drift effects. However, when noise caused by instrumental drift of the system starts to dominate, the Allan variance starts to increase, indicating a decrease of system stability and hence precision. CH₄ concentration measurements over a ten minute period of relative constant CH₄ concentrations with a mean value of 1905 ppb and a standard deviation of 4.74 ppb with 10Hz sampling rate were used to determine the Allan variance. Subsequently, the Allan variance of this period was plotted over the integration time T (Fig. 4.2), showing a decreasing Allan variance over integration times larger than 2.4 s. No increase of the Allan variance was observed at larger integration times, indicating an absence of instrumental drift, and thus high system stability and precision, for integration times of a few seconds up to ten minutes. Additionally, an indication for the short term precision (σ) was obtained by the y-axis intercept at the minimum Allan variance ($\sigma_{As}^2 = 6.1 \times 10^{-3} \text{ ppb}^2$). Using Eq. (4.3):

$$\sigma = \sigma_{As} \times f_s^{-1/2} \quad (4.3)$$

in which σ_{As} is the square root of the minimum Allan variance and f_s is the sampling frequency of the system (10Hz), σ was determined as 7.8×10^{-3} ppb Hz^{-1/2}. Precisions of 0.3 ppb Hz^{-1/2} (Nelson et al., 2004) and 2.9 ppb Hz^{-1/2} (Kroon et al., 2007) were found in previous studies of QCL instruments.

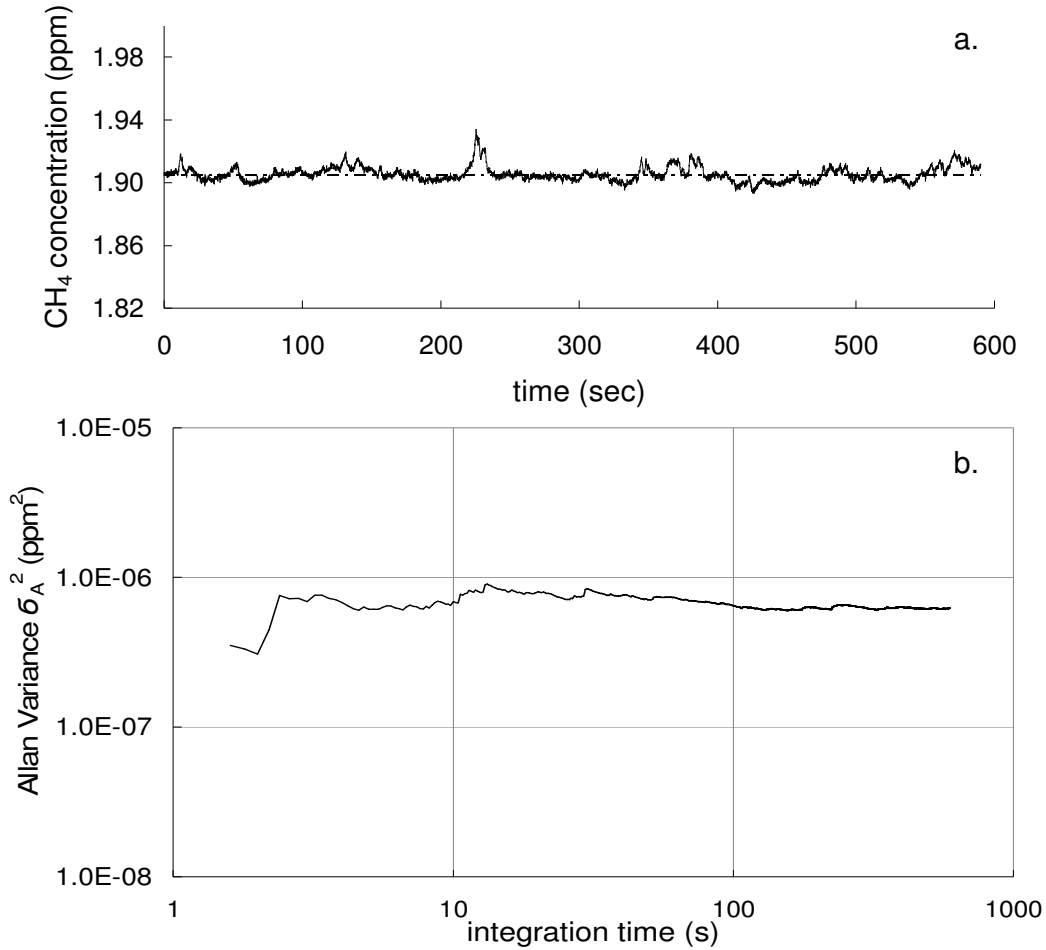


Figure 4.2: Time series of CH₄ concentration measurements with 10Hz sampling rate with a mean of 1905 ppb and a standard deviation of 4.74 ppb (a.) and the Allan variance plot for these data (b.).

Next, a calibration experiment was carried out by administering standard gas mixtures with concentrations of 125 ppb and 2002 ppb CH₄, both with an accuracy of 0.2 ppb, to the FMA at ten instances within a 10-day period. During the whole experiment the FMA was never turned off in order to imitate longer measurement periods as will be the case in the field set-up. The deviation from the standard gas values of CH₄ (MD) was calculated for each measurement by Eq. (4.4):

$$MD = \frac{M - S}{S} \quad (4.4)$$

where M is the measured CH_4 concentration in ppb and S is the CH_4 concentration of the standard gas in ppb. A two-point calibration factor (F_{cal}) was calculated for each set of measurements by Eq. (4.5):

$$F_{\text{cal}} = \frac{S_H - S_L}{M_H - M_L} \quad (4.5)$$

where S_H and S_L are the high and low standard values of CH_4 in ppb and M_H and M_L are the high and low measured values of CH_4 in ppb. Although the measured concentration sometimes varied one or two ppb from the standard values over the experimental period for both gases, no actual drift was observed in the instrument (Table 4.1). F_{cal} was 1.000 on average with fluctuations $< 0.30\%$, indicating high accuracy of the FMA.

Calibration experiment FMA							
measurement date	time	time elapsed (days)	low CH_4 gas (125 ppb) M_L (ppb)	MD_L	high CH_4 gas (2002 ppb) M_H (ppb)	MD_H	F_{cal}
02/04/2007	11:50:00	0.000	125	-0.10%	2005	0.20%	0.998
03/04/2007	09:00:00	0.882	127	1.50%	1999	-0.10%	1.003
03/04/2007	15:30:00	1.153	125	-0.10%	1999	-0.10%	1.002
04/04/2007	09:20:00	2.896	125	-0.10%	2000	-0.10%	1.001
04/04/2007	16:45:00	2.205	125	-0.10%	1999	-0.10%	1.002
05/04/2007	09:30:00	2.903	125	-0.10%	1999	-0.10%	1.002
05/04/2007	17:30:00	3.236	124	-0.90%	2001	0.00%	1.000
10/04/2007	09:45:00	7.913	125	-0.10%	2004	0.10%	0.999
11/04/2007	09:40:00	8.910	125	-0.10%	2002	0.00%	1.000
11/04/2007	16:30:00	9.194	125	-0.10%	2002	0.00%	1.000
12/04/2007	09:15:00	9.892	125	-0.10%	2004	0.10%	0.999

Table 4.1: Results of the 10-day calibration experiment with two standard gases of 125 and 2002 ppb respectively. Time elapsed since start of experiment, CH_4 concentrations measured by FMA (M_L and M_H), MD values and F_{cal} values are shown.

Additionally, over a period of a week, CH_4 concentration measurements at the Horstermeer site were compared with CH_4 concentrations measured at 20 and 60 m height at a measurement site in Cabauw (51°58'N and 4° 55'E)¹, approximately 30 km from the Horstermeer site. The CH_4 concentrations at Cabauw were measured with a Carlo Erba

¹ The CH_4 concentration measurements in Cabauw are measured with a gas chromatograph by the Energy Centre of the Netherlands (ECN).

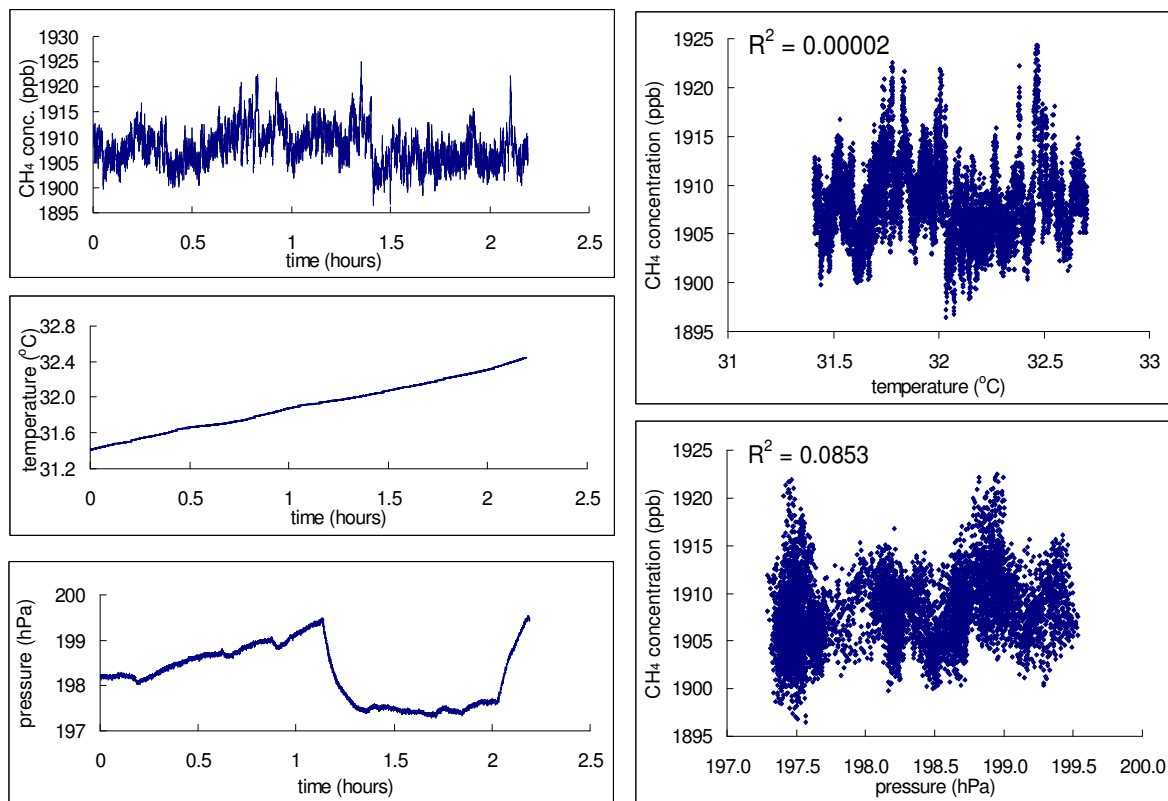


Figure 4.3: Results of the laboratory experiment on the effect of T_{cell} and P_{cell} on CH_4 concentration measurements. a.: time series of CH_4 concentration measurements; b.: time series of T_{cell} measurements; c.: time series of P_{cell} measurements; d.: scatter plot of CH_4 concentration against T_{cell} with correlation coefficient (R^2); e.: scatter plot of CH_4 concentration against P_{cell} with R^2 .

gas chromatograph system and have a precision of 2 ppb. The CH_4 concentration at the Horstermeer site was 15 ppb higher on average, which might be the result of the relatively high CH_4 emissions from the peat meadow area in which the measurements were taken. Nonetheless, the increasing trend of CH_4 concentration at the Horstermeer site was similar to the trend at the Cabauw site. Generally, slow variation in concentrations is caused by the difference between continental and marine background concentration, the continental concentration being approximately 50 ppb higher (Eisma et al., 1994). During the measurement period, prevailing winds were from the east (continental). This accounted for the slow rise in CH_4 concentrations at both measurement locations.

The influence of changing temperature and pressure conditions in the measurement cell on CH_4 concentration measurements was assessed in the laboratory. During a time series of continuous measurements, a step change in P_{cell} was induced, while the temperature in the measurement cell (T_{cell}) increased steadily. An effect of the increase of T_{cell} on CH_4 concentration was observed neither from the time series, nor from the correlation plots (Fig. 4.3). A decrease in the CH_4 concentration data was observed 990 sec after the sharp decrease in P_{cell} . However, this feature did not result in a clear correlation between CH_4

concentration and P_{cell} . Also, this type of P_{cell} changes did not occur under normal circumstances when P_{cell} had a stable value near 210 hPa.

4.2.4 Incorporation of FMA in eddy covariance system

After testing in the laboratory, the FMA was installed in a closed-path eddy covariance field set-up (Fig. 4.4). In order to obtain 10Hz measurements, a dry vacuum scroll pump (XDS35i, BOC Edwards, Crawly, UK) was used with a maximum pumping speed of $9.72 \times 10^{-3} \text{ m}^3 \text{ s}^{-1}$. However, at the required pressure of 210 hPa the actual pumping speed was $5.50 \times 10^{-3} \text{ m}^3 \text{ s}^{-1}$. The scroll pump has relatively high power requirements: 100/200V to 120/230V, 50/60Hz and 600W. It was placed at the end of the set-up, connected to the FMA by a wire-reinforced tube with an internal diameter of $1.9 \times 10^{-3} \text{ m}$, sucking air through the system. The FMA was placed in a heated, water resistant box, while the scroll pump was placed in an aerated box that prevented it from getting wet and from overheating. In addition to the internal filter with a pore size of $2 \mu\text{m}$ (Swagelok part no. SS-4FW4-2), a filter with a pore size of $60 \mu\text{m}$ was placed at the inlet in order to prevent dust, aerosols, insects and droplets from entering the tubing. The inlet was shielded from the rain by a stainless steel cap. To prevent any water that accidentally passed the first filter from moving down toward the analyzer the air was first led up through a stainless steel tube (diameter of $6.4 \times 10^{-3} \text{ m}$) that bends sharply at 0.5 m after which the air moves

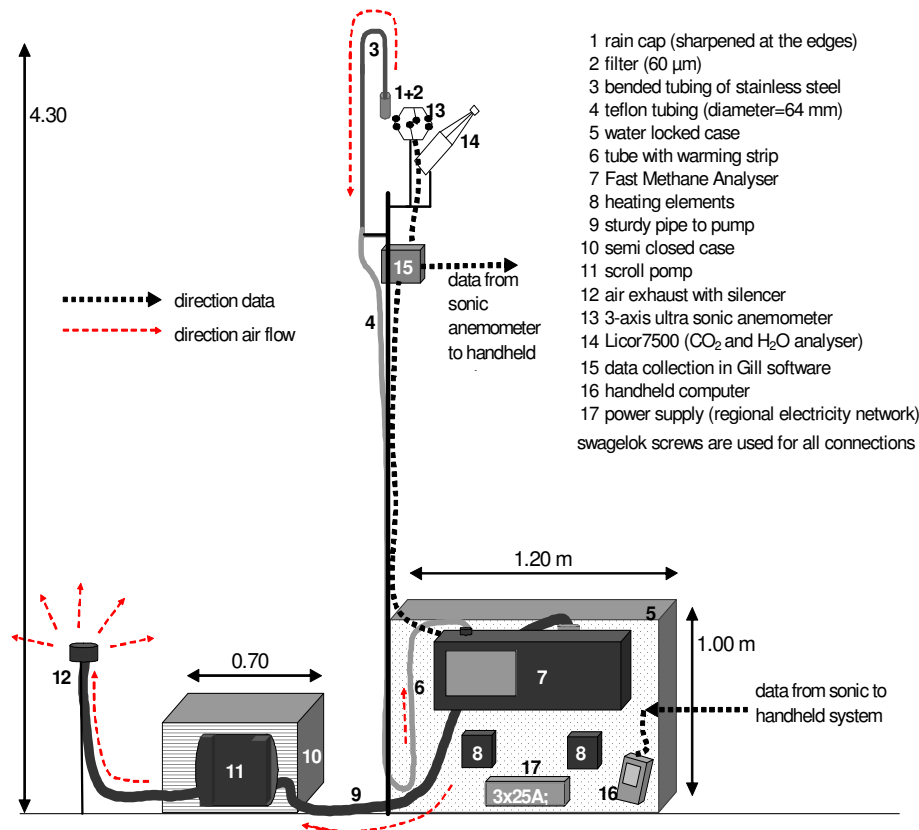


Figure 4.4: Schematic overview of the combined field set-up of the closed path eddy covariance system for CH_4 using the FMA and the open path eddy covariance system for CO_2 and water vapour.

down toward the analyzer through a Teflon tube (diameter of 6.4×10^{-3} m). After the air has passed through the FMA measurement cell it flows through the scroll pump and was exhausted through a silencer. The gas-inlet filter was positioned 0.2 m away from the LI-7500 open path infrared gas analyzer (LI-COR Lincoln, NE, USA) and a Windmaster Pro 3-axis ultrasonic anemometer (GILL Instruments Limited, Hampshire, UK) directed into the prevailing wind. Both instruments were installed at 4.3 m above the surface at the Horstermeer measurement site (Hendriks et al., 2007). Data were logged digitally on a handheld computer at a rate of 10 Hz (Van der Molen et al., 2006).

4.2.5 Assessment of FMA in measuring CH_4 fluxes

In general a sampling rate of 10Hz, with a Nyquist frequency of 5Hz, is used for eddy covariance techniques (Aubinet et al., 2000; Kroon et al. 2007). Therefore an instrumental time response of 10Hz or greater is required in order to correlate with the wind measurements made with the 3-axis ultrasonic anemometer. The measurement rate of an instrument is determined by both electronic signal processing and by the τ of the measurement cell (Nelson et al., 2004). Electronic signal processing is dependent on the spectral complexity of the measurement technique as well as the technical design of the ICOS technique. In the case of the FMA, this was defined by the designers (Los Gatos Research Ltd.) as 20Hz. The limiting factor of the maximum sampling rate is often τ , which could be determined by the volume of the measurement cell ($0.55 \times 10^{-3} \text{ m}^3$) divided by the actual pumping speed ($5.50 \times 10^{-3} \text{ m}^3 \text{ s}^{-1}$) giving a flow response of 0.10 s. Additionally, τ was determined by applying a step change in concentrations at 20Hz sampling rate (Fig. 4.5) (Moore, 1986; Zahniser et al., 1995; Nelson et al., 2004). Each data point was the average mixing ratio of multiple step change events at a certain t (time elapsed since step change in concentration). The τ was defined by the exponential fit to the decay of the CH_4 mixing ratio and was calculated $\tau = 0.11$ s by Eq. (4.6):

$$[\text{CH}_4]_t = [\text{CH}_4]_{t=0} e^{(-t/\tau)} \quad (4.6)$$

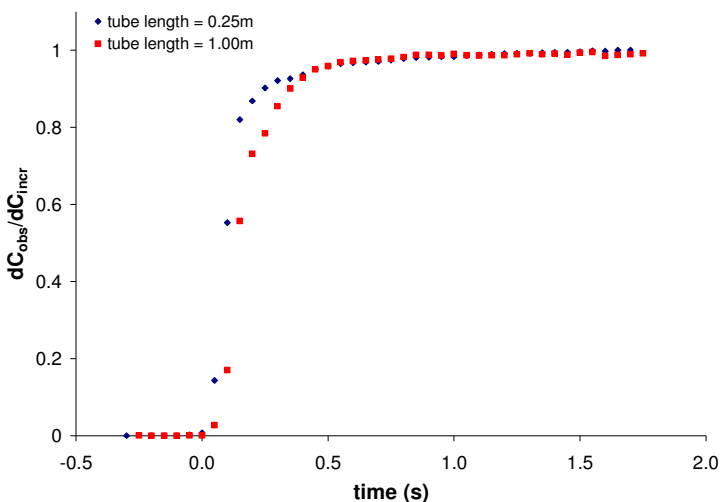


Figure 4.5: Averaged and normalised time series of CH_4 concentration data showing the instrument response with 20Hz sampling rate changing from ambient air to a gas sample with a high CH_4 concentration for tube lengths of 0.25 m and 1.0 m.

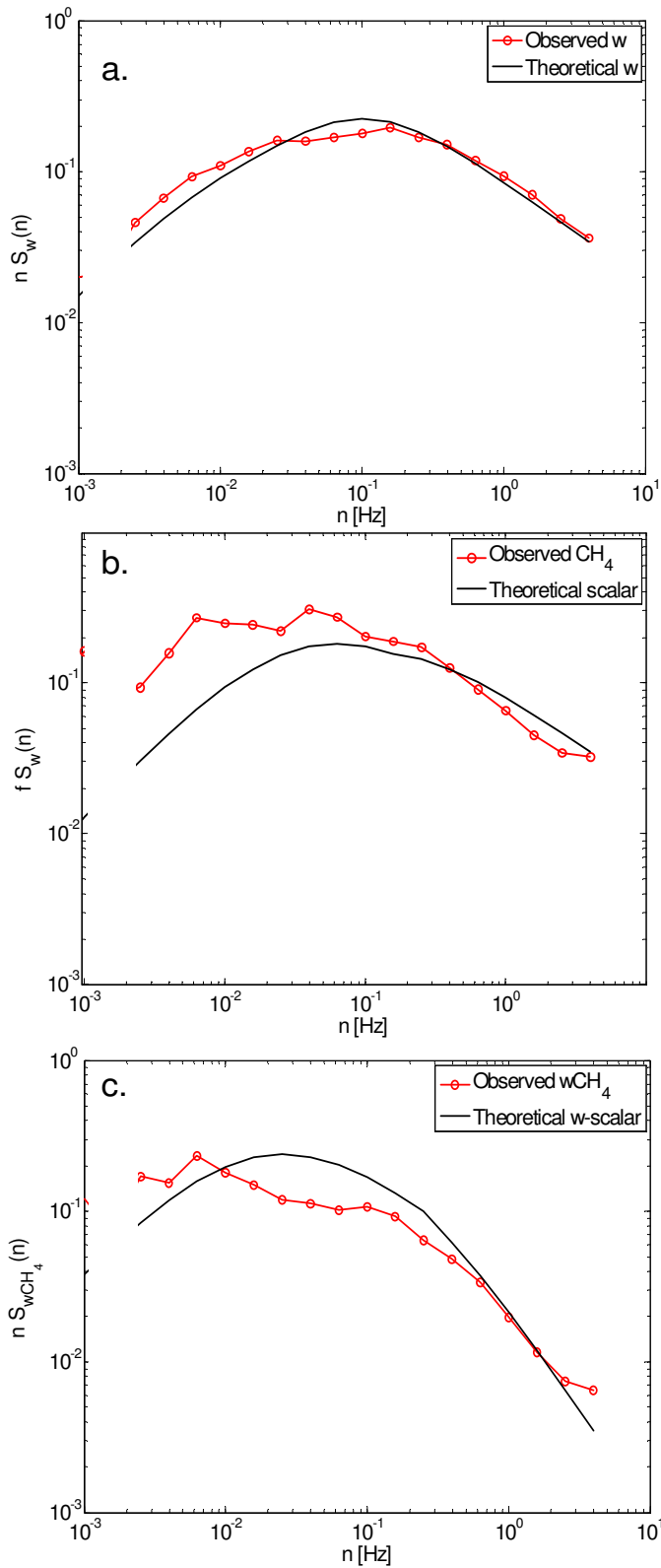


Figure 4.6: Averaged and binned results of the spectral analyses of the FMA eddy covariance set-up for two moderately turbulent days (six half hour periods per day). a.: Observed and theoretical power-spectrum w ; b.: Observed and theoretical power-spectrum of $[CH_4]$; c.: Observed and theoretical co-spectrum of $w'[CH_4]$.

where $[CH_4]_t$ is the CH_4 concentration at instance t . In this, sensitivity to tube length could be observed (Fig. 4.5). The data in the graph are normalized by the transformation dC_{obs}/dC_{incr} , where dC_{obs} is the increase of $[CH_4]$ between the starting time and time t and dC_{incr} is difference between the final $[CH_4]$ and $[CH_4]$ at the starting time. Since the effect of tube damping was corrected separately, and τ only refers to the response of the instrument itself, the actual τ was determined as 0.10 s.

The suitability of the FMA eddy covariance set-up was further evaluated by examining the power spectra of w and $[CH_4]$ and co-spectrum of the covariance $w'[CH_4]'$ (Stull et al., 1988; Kaimal and Finnigan, 1994). For this purpose twelve half hours of data from two days with moderately unstable conditions were analysed. The results of the spectral analysis were averaged and binned and the logarithmic spectral and co-spectral densities were plotted against frequency (Fig. 4.6). Additionally, the theoretical spectra defined by Kaimal et al. (1972) and Højstrup (1981) were plotted in the graphs (Smeets et al., 1998). The power spectrum of w very closely followed the whole theoretical spectrum, while both the $[CH_4]$ power spectrum and the $w'[CH_4]'$ co-spectrum showed slight deviations from the theoretical spectrum. Most important is that the inertial subranges of both the power spectra and the co-spectrum were in general agreement with the theoretical curve. The $[CH_4]$ power spectrum showed a slight tendency to extend upward in the low frequency range. This was also observed in the $[CO_2]$ power spectrum from the LI-7500 analyzer at the site (Hendriks et al., 2007), indicating that the upward tendency in the low range was not instrument related, but was rather an effect of environmental factors. Kaimal et al. (1976) found high densities of low frequencies in the temperature power spectrum in response to a diurnal cycle. Since a diurnal cycle was observed for both $[CO_2]$ and $[CH_4]$ at the Horstermeer site, this might have been the cause of the upward tendency observed in the power spectra. However, the overestimation of low frequencies will automatically cause an underestimation of the high frequencies, since the total area under the power spectrum is a fixed surface (Kaimal et al., 1972; Højstrup, 1981). Considering the $w'[CH_4]'$ co-spectrum, the observed peak near $n=10^{-1}$ Hz generally matched the peak of the theoretical spectrum and was in agreement with spectral analysis of CH_4 flux measurements by Verma et al. (1992), Kormann et al. (2001) and Werle and Kormann (2001). The spectral performance of the eddy covariance set-up for temperature and CO_2 fluxes were discussed previously in a paper by Hendriks et al. (2007) and were in agreement with the spectral analyses of the CH_4 flux measurements.

Additionally, in previous research by Hendriks et al. (2007) the energy balance consisting of the eddy covariance measurements of latent heat and sensible heat, the micro-meteorological measurements of incoming and reflected radiation components and the ground heat flux showed a closure of 82%. This indicated that the data from the eddy covariance set-up were acceptable (Lloyd et al., 1997).

The travel time of the air in the closed path set-up from the inlet filter to the FMA, caused a time lag with respect to the in situ measured wind data. For 20 half hour periods, the covariance $w'[CH_4]'$ of w' at instance $t = 0.0$ s and $[CH_4]'$ at $t = 0.0$ s was determined, as well as the covariance of w' at $t = 0.0$ s and $[CH_4]'$ at the $t = 0.1$ s, $t = 0.2$ s, $t = 0.3$ s, ..., $t = 2.0$ s. For all half hour periods the highest value of $w'[CH_4]'$ occurred with $[CH_4]'$ at $t =$

0.6 s. For the calculation of the actual covariance the time lag of the CH₄ measurements compared to the wind measurements was therefore taken as 0.6 s.

4.2.6 Data processing

The EUROFLUX methodology (Aubinet et al., 2000) was applied to the eddy covariance data to calculate the CH₄ fluxes from the closed path system and the CO₂ fluxes from the open path system (Hendriks et al., 2007) on a thirty minute basis. Since system instrumental drift was not observed, an overestimation of the fluxes as a result of this averaging period was not expected. The damping effect of the τ on the CH₄ signal was corrected for in the flux calculation procedure as well as for the time lag of 0.6 s between closed path CH₄ and open path wind, temperature and CO₂ measurements (Moore et al., 1986). The tube length of the set-up was over 1000 times the inner diameter of the tube and therefore the air temperature in the measurement cell could be considered stable. As a result, the Webb correction for density fluctuations arising from variations in temperature that was applied to the open path CO₂ measurements, was not required for the closed path measurements of CH₄ (Leuning and Moncrieff, 1990). The Webb correction for density fluctuations arising from variations in water vapour (measured with the LI-7500) was applied according to Leuning and Moncrieff (1990). Frequency loss corrections for closed-path systems were applied according to the theory of Leuning and King (1992). Since 3-axis ultrasonic anemometers were found to under measure wind speed at large angles, the method of Nakai et al. (2006) was used to apply the angle of attack dependent calibration (Gash and Dolman, 2003, Van der Molen et al., 2004).

4.2.7 Simulation of alternative flux measurement approaches

In eddy covariance, the sampling rate determines the number of samples that are taken out of an infinitely large number of samples. The higher the sampling rate, the higher the statistical reliability, which results in higher accuracy of the observed means and covariances (Van der Molen, 2002). In order to obtain reliable estimates of fluxes, a sampling rate of at least 10Hz is generally used for eddy covariance techniques. However, measurements performed at lower rates than 10Hz can generate reliable results too (Rinne et al., 2000 and 2001; Graus et al., 2006; Businger and Oncley, 1990). 1Hz eddy covariance, disjunct eddy covariance, 1Hz eddy covariance and relaxed eddy accumulation (REA) are possible alternatives for the preferred 10Hz eddy covariance. These measurement techniques could make the external pump of the FMA eddy covariance set-up superfluous and thus save over half of the energy required for the eddy covariance set-up. This might be necessary for operation in remote places where an external power source is not available. Here, the raw CH₄, CO₂ and water vapour 10Hz field data were manipulated to simulate 1Hz eddy covariance, disjunct eddy covariance and REA.

With 1Hz eddy covariance, F_s is determined in the same manner as 10Hz eddy covariance (Eq. (4.2)), but at a lower frequency. The method was tested by averaging each 10 consecutive data points of CH₄ concentration as well as wind velocity for the 10Hz data set, thereby simulating a slower sampling rate (1Hz) with a longer τ (1.0 s instead of 0.10 s).

Disjunct eddy covariance uses a subset of the whole 10 Hz time series to determine the flux of an atmospheric property F_s according to Eq. (4.7):

$$F_s = \langle w' s' \rangle = \frac{1}{N} \times \sum_{i=1}^N w_i' s_i' \quad (4.7)$$

where N is the subset of the data (Rinne et al., 2000 and 2001). Here, the disjunct eddy covariance method was tested by sampling the first data point of every 10 data points from the 10Hz data set, thereby creating a time interval of 0.9 s between the sampling moments, while the τ remained 0.10 s. To build a field set-up for disjunct 1Hz eddy covariance a ‘snap sampling’ instrument would have to be mounted in front of the inlet of the system to obtain samples that are sampled with a frequency of 1Hz, while maintaining the sampling duration as short as possible (‘snap’). This method would give the FMA an analysis time of 1.0 s instead of 0.10 s.

REA is a conditional sampling method in which air samples are drawn into two separate reservoirs depending on the direction of w . The criterion of valve switching is based on values of the standard deviation of w (σ_w), which is measured by the 3-axis ultrasonic anemometer at 10Hz. The valve is activated according to the threshold condition $|w_0| = 0.6\sigma_w$. In the case of $-w_0 \leq w \leq w_0$ (deadband values), neither “up” nor “down” samples are taken, but samples are discarded from the sampling system (Graus et al., 2006). Here, we simulated REA by dividing all eddy covariance data points of one half hour measurement period into three data matrices based on the direction of the w (upward, downward and deadband values). Next, the measured concentrations are summed and the turbulent flux of the scalar s (F_s) was determined according to Eq. (4.8):

$$F_s \approx b \times \sigma_w \times \Delta s \quad (4.8)$$

(Businger and Oncley, 1990). The b -value is the correction for the deadband and Δs the difference between the concentrations in the accumulation reservoirs (Bowling et al., 1999).

4.3 Results

4.3.1 First data series

From the eddy covariance data series 11% consisted of gaps due to failure of the eddy covariance set-up caused by rain events and instrumental errors. Additionally, 3% of the data series consisted of spikes (CH_4 flux $> 100.0 \text{ nmol m}^{-2} \text{ s}^{-1}$ and CH_4 flux $< -10.0 \text{ nmol m}^{-2} \text{ s}^{-1}$) and were removed from the data set. During nighttime periods with low friction velocity (u_*) the turbulence of the atmosphere can become too low for the performance of eddy covariance measurements (Wohlfahrt et al., 2005; Dolman et al., 2004). In order to determine the critical u_* value for CH_4 eddy covariance measurements at this specific site, the CH_4 flux data and u_* data from periods with incoming shortwave radiation (SWin) $<$

20W m⁻² were selected (Dolman et al., 2004). The nightly CH₄ fluxes showed a significant decrease for periods with $u_* < 0.09 \text{ m s}^{-1}$ (Fig. 4.7). This result is similar to the critical u_* value of 0.10 m s^{-1} found for CO₂ fluxes at the same site (Hendriks et al., 2007). CH₄ flux data measured during periods with $u_* < 0.09 \text{ m s}^{-1}$ occurred at 12% of the data series and were removed too. The total amount of data gaps accounted for 26% of the whole data series.

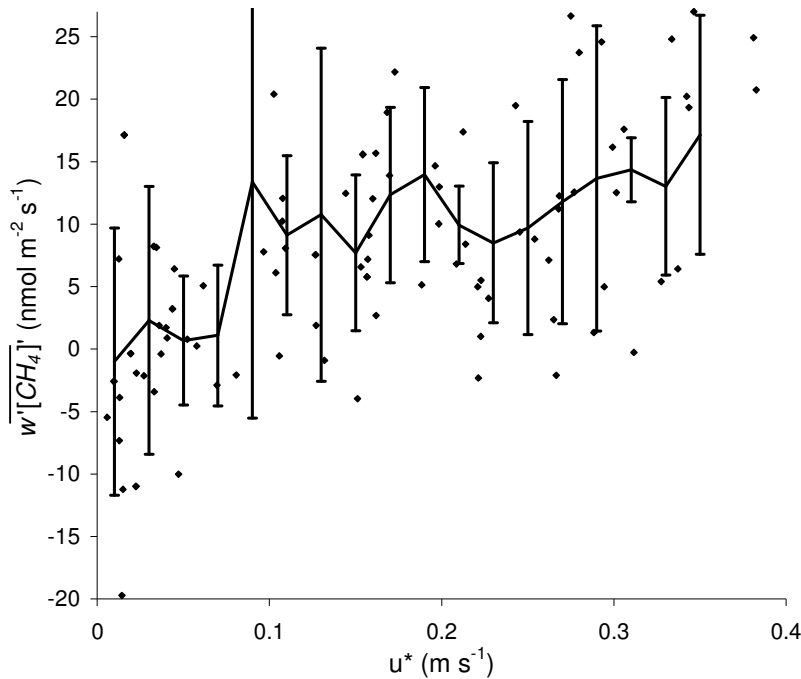


Figure 4.7: Results of the analyses of the effect of low turbulence on nightly CH₄ fluxes, showing a drop in flux magnitude below of u_* of 0.09 m s^{-1} . Data are binned over u_* classes of 0.02 m s^{-1} and error bars show the standard deviation per class.

CH₄ fluxes and CO₂ fluxes (net ecosystem exchange (NEE)) were plotted for a two week period in June 2006 at the Horstermeer site (Fig. 4.8). Although the CH₄ fluxes are rather variable over time, a diurnal cycle can be observed with low emission during the night and high emission during the day. CO₂ fluxes have a similar, but opposite, diurnal cycle. The observed CH₄ fluxes consisted mainly of emission and had an average of $29.7 \text{ nmol m}^{-2} \text{ s}^{-1}$, while the typical maximum CH₄ emission was approximately $80.0 \text{ nmol m}^{-2} \text{ s}^{-1}$. The typical minimum flux was approximately $0.0 \text{ nmol m}^{-2} \text{ s}^{-1}$ and at three occasions a small uptake was observed.

4.3.2 Intercomparison with flux chamber data

For two days during the CH₄ eddy covariance measurement period at the Horstermeer site, flux chamber measurements were made in the footprint of the eddy covariance tower with a Photoacoustic Field Gas-Monitor (type 1312, Innova AirTech Instruments, Ballerup, Denmark) connected with tubes to closed, dark chambers (Hendriks et al., 2007). Flux chamber data were collected at the three land elements occurring in the footprint of the eddy covariance tower: relatively dry peat land, saturated peat land and ditch water surfaces. The fluxes from the various land elements are averaged with respect to their relative surface area (70%, 20% and 10% respectively; Hendriks et al., 2007). On June 10 the

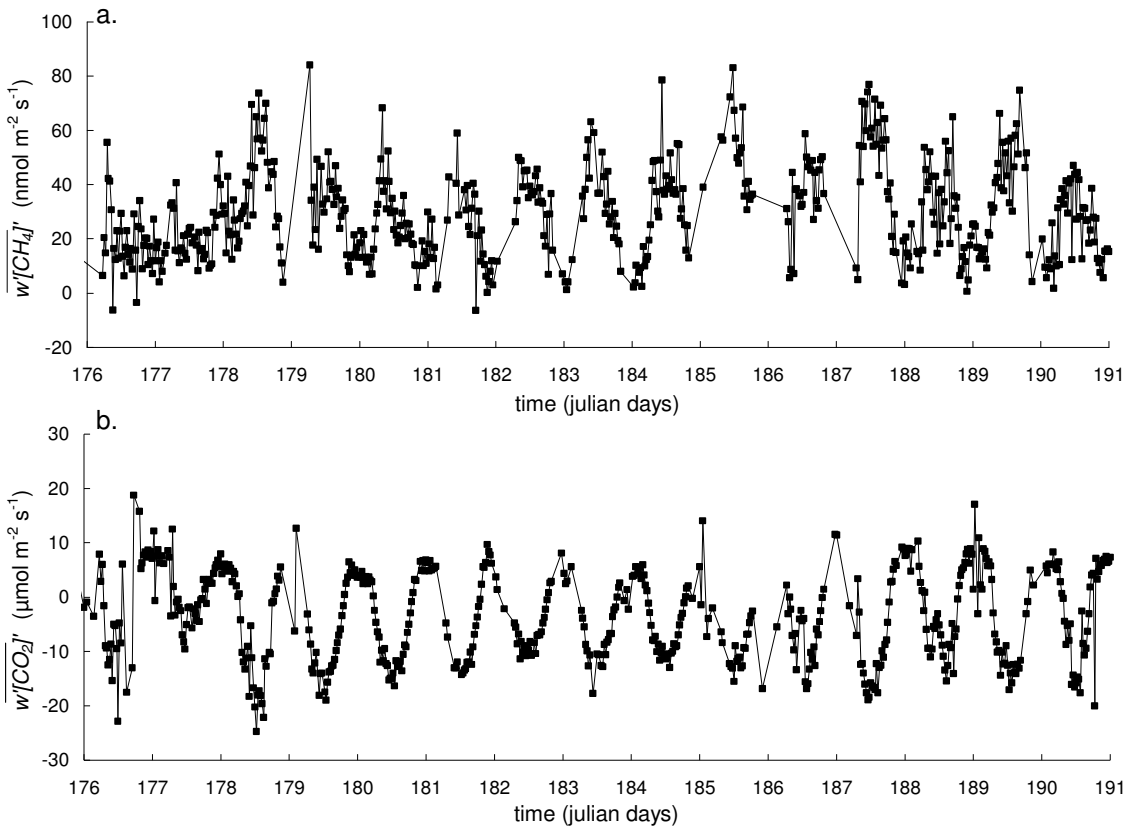


Figure 4.8: CH₄ flux data series over a two week period in the summer of 2006 (a.) and CO₂ flux (NEE) data series over the same period (b.).

flux chamber measurements showed an average CH₄ flux of $114.5 \pm 9.6 \text{ nmol m}^{-2} \text{ s}^{-1}$, while the eddy covariance measurements showed an average CH₄ flux of $83.2 \text{ nmol m}^{-2} \text{ s}^{-1}$ over the same period (Fig. 4.9). On October 3 the average CH₄ flux from the chamber measurements was $53.6 \pm 11.2 \text{ nmol m}^{-2} \text{ s}^{-1}$, while that from eddy covariance was $61.6 \text{ nmol m}^{-2} \text{ s}^{-1}$.

4.3.3 Simulation of alternative flux measurement approaches

The simulated 1Hz and disjunct eddy covariance data as well as the simulated REA data were compared with the original 10Hz eddy covariance data as half hourly averages over a 15 day period (Table 4.2). It can be observed that the various methods show similar but not identical results and that all alternative methods showed variations from the 10Hz eddy covariance data (Fig. 4.10). The 1Hz eddy covariance method showed on average a slight overestimation of the half hourly fluxes for CH₄ measurements (2%) and the standard deviation of the time series was somewhat higher than that of the 10Hz eddy covariance. Average CO₂ and water vapour fluxes determined with 1Hz eddy covariance showed however a large underestimation compared to the 10Hz eddy covariance measure-

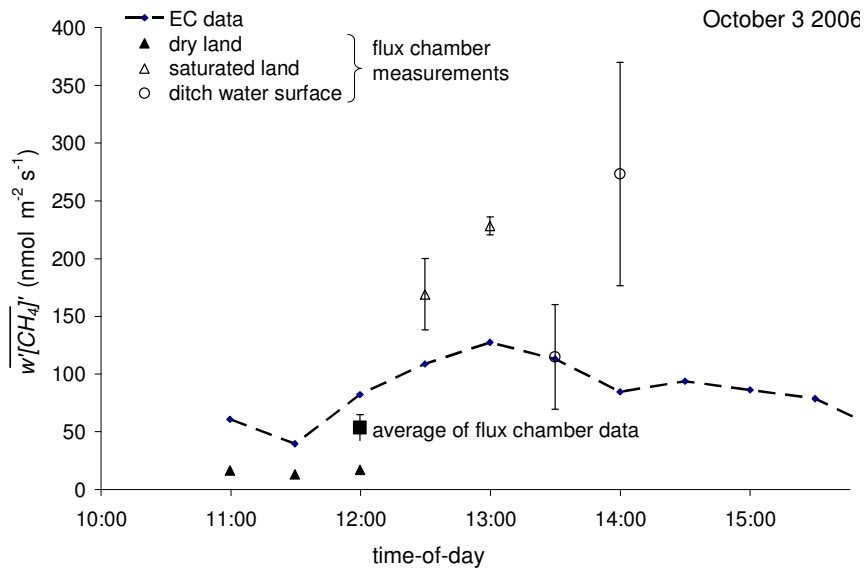
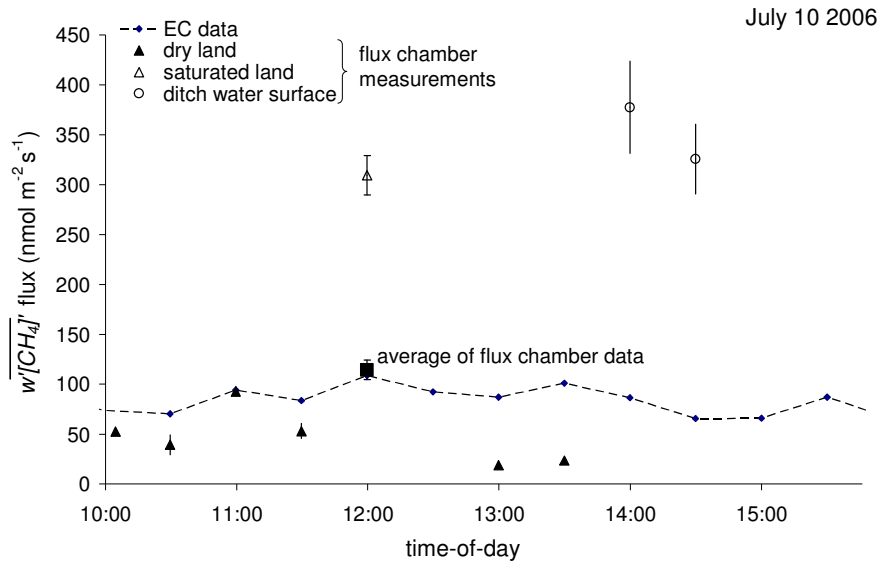


Figure 4.9: Hourly CH₄ flux data at July 10 and October 3 (both 2006) plotted in combination with flux chamber data from various land elements in the footprint of the eddy covariance tower collected at the same day. The square marks show the weighed average of the flux chamber measurements.

ements. The disjunct eddy covariance method showed on average a slight underestimation of the half hourly fluxes for CH₄ as well as for CO₂ and water vapour measurements (-2% and -3%), while the standard deviation was somewhat higher than that of the 10Hz eddy covariance.

The *b*-value used in REA for correction of the deadband area was determined by Pattey et al. (1993) as 0.56. For the measurement set-up investigated here, a *b*-value of 0.20 showed results most similar to the normal 10Hz eddy covariance measurements for CH₄ measurements as well as for CO₂ and water vapour measurements. The results of the REA method showed on average a small underestimation of the half hourly fluxes for CH₄ as well as for CO₂ and water vapour measurements (-4% and -2%). The standard deviation however, was significantly higher than that of the 10Hz eddy covariance, pointing at relatively large deviations from the 10Hz measurements at a half hourly basis.

		10Hz EC	1Hz EC	Disjunct 1Hz EC	REA (d=0.6;b=0.2)
average gas flux	CH ₄ flux (nmol m ⁻² s ⁻²)	26.74	27.35	26.29	25.76
	CO ₂ flux (μmol m ⁻² s ⁻²)	-6.53	-5.84	-6.31	-6.41
	H ₂ O flux (mmol m ⁻² s ⁻²)	3.13	2.79	3.05	3.01
average deviation from normal 10Hz eddy covariance	CH ₄ flux (%)	0%	2%	-2%	-4%
	CO ₂ flux (%)	0%	-11%	-3%	-2%
	H ₂ O flux (%)	0%	-11%	-3%	-4%
standard deviation	CH ₄ flux (nmol m ⁻² s ⁻²)	11.65	12.23	12.7	14.34
	CO ₂ flux (μmol m ⁻² s ⁻²)	9.95	9.13	9.86	11.22
	H ₂ O flux (mmol m ⁻² s ⁻²)	2.82	2.59	2.8	2.77

Table 4.2: Summary of results of 1Hz eddy covariance, disjunct eddy covariance and REA compared to 10Hz eddy covariance for CH₄, CO₂ and water vapour (H₂O) flux measurements. Average gas flux over the 15 day period, average deviation from the 10Hz eddy covariance data and standard deviations of half hourly data are shown.

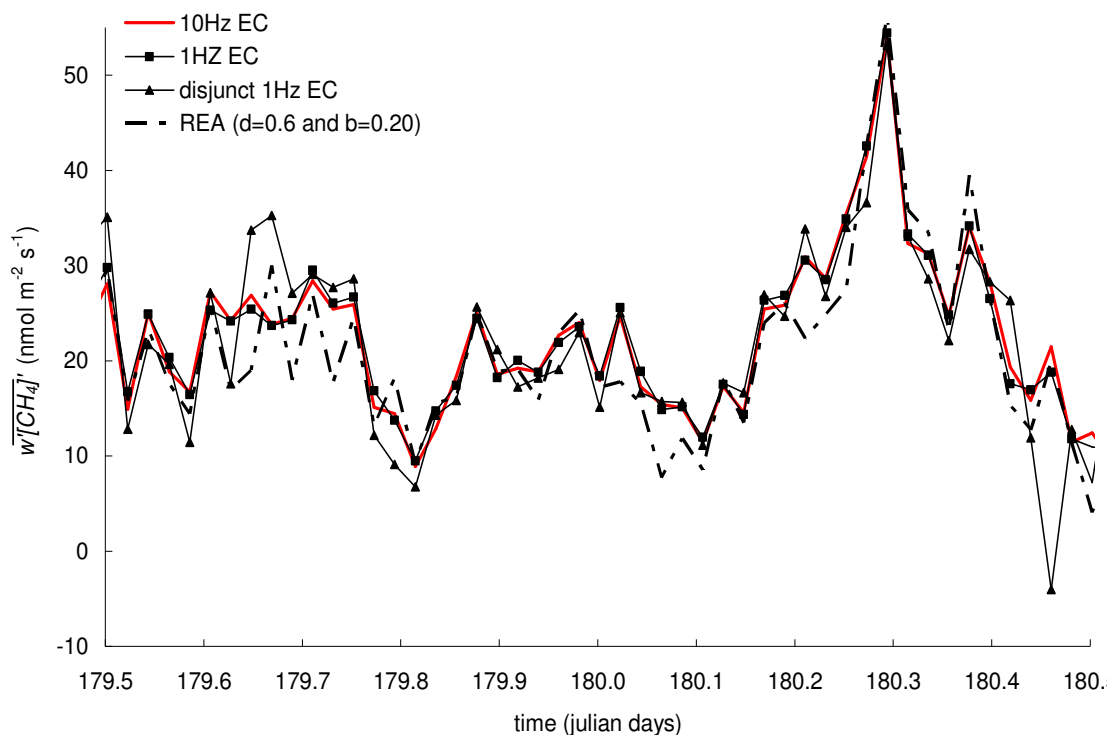


Figure 4.10: Time series of CH₄ fluxes for a one day period: half hourly averages of 10Hz eddy covariance and simulated 1Hz eddy covariance, disjunct eddy covariance, and REA.

4.4 Conclusions and discussion

The FMA, which uses the new off-axis ICOS technique, was found to perform satisfactorily in laboratory experiments and in the eddy covariance field set-up. Compared to other techniques, the absence of a large nitrogen Dewar for cooling that would need weekly refill, the compact, narrow band and stable laser, the relative user friendliness and low costs are considerable advantages. The FMA eddy covariance set-up was found to perform independently in an unattended field situation. However, care should be taken when placing the instrument with a scroll pump in the field. The FMA should be kept dry and in ambient temperatures between 5°C and 45°C, while the scroll pump should be kept dry and cool. Contamination of the measurement cell should be prevented, since the mirrors inside the measurement cell are sensitive and only a small contamination might cause a rapid decrease in reflectivity of the mirrors and performance of the instrument.

The analysis of the Allan variance indicated a high precision and system stability of the FMA. Additionally, the calibration experiment showed sufficiently high accuracy (< 0.30 %). As long as ambient temperatures do not exceed the range of 5 to 45 °C and pressure in the measurement cell is near 210 hPa, the CH₄ measurements are not affected by changes in T_{cell} and P_{cell}. Since no instrumental drift was observed in the Allan variance analyses, in the calibration experiment or in the comparison with CH₄ concentration data at a field site nearby, it was concluded that frequent calibration of the FMA was not necessary.

The observed τ was 0.10 s, implying a maximum sampling rate of 10Hz which is sufficient for eddy covariance measurements. The closed path FMA eddy covariance system performed well, as shown by the power- and co-spectra which corresponded well to the theoretical spectral curves. Additionally, the energy balance showed satisfactory closure. CH₄ fluxes appeared to be underestimated during periods with low turbulence and a u^* correction was applied for CH₄ flux data during periods with $u^* < 0.09 \text{ m s}^{-1}$. Due to erroneous measurements, spikes and periods of low turbulence the data series consisted for 26% of gaps.

Observed CH₄ fluxes consisted mainly of emission, showed a diurnal cycle, but were rather variable over time. The average CH₄ emission was $29.7 \text{ nmol m}^{-2} \text{ s}^{-1}$, while the typical maximum CH₄ emission was approximately $80.0 \text{ nmol m}^{-2} \text{ s}^{-1}$ and the typical minimum flux was approximately $0.0 \text{ nmol m}^{-2} \text{ s}^{-1}$. These CH₄ fluxes were in agreement with QCL flux measurements at a managed peat meadow site in the Netherlands (Reeuwijk), where emissions were $40 \pm 31 \text{ nmol m}^{-2} \text{ s}^{-1}$ on average (Kroon et al., 2007) and average annual CH₄ emission of $30 \text{ nmol m}^{-2} \text{ s}^{-1}$ from peat lands in Germany and the Netherlands (Drösler et al., 2007). From the comparison of the eddy covariance measurements with flux chamber measurements, it was observed that the fluxes from the two techniques showed a discrepancy of approximately 20%. However, considering the fact that the flux chamber measurements are point measurements at the soil surface while the eddy covariance has a footprint of hundreds of square meters, some degree of variation may be expected.

Additionally, raw CH₄, CO₂ and water vapour 10Hz field data were manipulated to simulate 1Hz eddy covariance, disjunct eddy covariance and REA. It was concluded that,

when using a scroll pump is not possible for technical or practical reasons; disjunct eddy covariance was the most reliable substitute for 10Hz eddy covariance. This method showed the highest degree of resemblance with the 10Hz eddy covariance for all three gases. The simulated 1Hz eddy covariance of CO₂ and water vapour fluxes, showed relatively large deviations from the 10Hz eddy covariance data, indicating a reduced reliability of the method and no possibility to test the validity of the method at sites without the eddy covariance set-up for CH₄. Simulation of REA did show similar results for all three gases; however, the standard deviation of the time series was significantly higher than that of the 10Hz eddy covariance, pointing at relatively large deviations from the 10Hz measurements at the half hourly basis. REA was therefore evaluated to generate reliable estimates of fluxes over periods in the order of days or weeks. Importantly, the *b*-value for CH₄ flux measurements was the same as that for the CO₂ and water vapour flux measurements. This indicates that it will be possible to determine the *b*-value for CH₄ measurements with REA at new and remote locations using an eddy covariance set-up with low power requirements for CO₂ or water vapour. Finally, it should be taken into account that the ‘snap sampling’ instrument for disjunct eddy covariance and the valve switching system for REA will introduce additional artefacts, which also require certain amounts of power (Rinne et al., 2000; Kuhn et al., 2005; Graus et al., 2006).

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Chapter 4

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