Effect of sporadic destratification, seasonal overturn and artificial mixing on CH_4 emissions at the surface of a subtropical hydroelectric reservoir (Nam Theun 2 Reservoir, Lao PDR)

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Abstract

Inland waters in general and specifically freshwater reservoirs are recognized as source of CH_4 to the atmosphere. Although the diffusion at the air-water interface is the most studied pathway, its spatial and temporal variations are poorly documented.

We measured fortnightly CH_4 concentrations and physico-chemical parameters at nine stations in a subtropical monomictic reservoir which was flooded in 2008 (Nam Theun 2 Reservoir, Lao PDR). Based on these results, we quantified CH_4 storage in the water column and diffusive fluxes from June 2009 to December 2012. We compared diffusive emissions with ebullition from Deshmukh et al. (2014) and aerobic methane oxidation and downstream emissions from Deshmukh et al. (2015).

In this monomictic reservoir, the seasonal variations of CH_4 concentration and storage were highly dependant of the thermal stratification. Hypolimnic CH_4 concentration and CH_4 storage reached their maximum in the warm dry season (WD) when the reservoir was stratified. They decreased during the warm wet (WW) season and reached its minimum after the reservoir overturned in the cool dry season (CD). The sharp decreases of the CH_4 storage were concomitant with sporadic extreme diffusive fluxes (up to 200 mmol m^{-2} d^{-1}). These hot moments of emissions occurred mostly in the inflow region in the WW season and during the overturn in the CD season in the area of the reservoir that has the highest CH_4 storage. Although they corresponded to less than 10% of the observations, these CH_4 extreme emissions (>5 mmol m^{-2} d^{-1}) contributed up to 50% of total annual emissions by diffusion.

During the transition between the WD and WW seasons, a new hotspot of emissions was identified upstream of the water intake where diffusive fluxes peaked at 600 mmol m^{-2} d^{-1} in 2010 down to 200 mmol m^{-2} d^{-1} in 2012. In the CD season, diffusive fluxes from this area were the lowest observed at the reservoir surface. Emissions from this area contributed 15-25% to total annual emissions although they occur on a surface area representative of less than 1% of the total reservoir surface. We highly recommend measurements of diffusive fluxes around water intakes in order to evaluate if such results can be generalized.

1. Introduction

Since the 1990s, hydroelectric reservoirs are known to be source of methane (CH_4) to the atmosphere. Their contribution to total CH_4 emissions still needs refinement since the discrepancies among estimates is large, ranging from 1 to 12% of total CH_4 emissions (St Louis et al., 2000;Barros et al., 2011). These two estimates are mostly based on diffusive

fluxes at the air-water interface and they overlook emissions from the rivers downstream of the dams (Abril et al., 2005;Guerin et al., 2006;Kemenes et al., 2007;Teodoru et al., 2012;Maeck et al., 2013;Deshmukh et al., 2015), CH_4 ebullition (DelSontro et al., 2010;Deshmukh et al., 2014) and emissions from the drawdown area of reservoirs (Chen et al., 2009;Chen et al., 2011) although these pathways could largely dominate diffusion at the surface of the reservoirs.

Even if CH_4 diffusion at the surface of reservoir is the best-documented emission pathway, little information is available on spatial and temporal variability of CH_4 emissions by diffusive fluxes. In tropical amictic reservoirs, the highest diffusive fluxes are usually observed during dry periods and when the stratification weaken at the beginning of the rainy season (Guerin and Abril, 2007). A study of CH_4 emissions from a dimictic reservoir suggests a potential large outgassing of CH_4 during the reservoir overturns (Utsumi et al., 1998b) as it is the case in natural monomictic and dimictic lakes (Kankaala et al., 2007;López Bellido et al., 2009;Schubert et al., 2010;Schubert et al., 2012;Fernández et al., 2014). Such hot moments of emissions (McClain et al., 2003) could contribute 45-80% of CH_4 annual emissions by diffusion (Schubert et al., 2012;Fernández et al., 2014). They are rarely taken into account in carbon budgets since they can only be captured by high frequency monitoring. Spatial heterogeneity of CH_4 emissions at the surface of reservoirs is also very high. It mostly depends on the spatial variations of ebullition that is controlled by sedimentation (DelSontro et al., 2011;Sobek et al., 2012;Maeck et al., 2013). The spatial variation of diffusion appears to be low with emissions being slightly higher (1) in area where dense forest is flooded as compare to the former riverbed (Abril et al., 2005), (2) at shallow sites than at deeper ones (Zheng et al., 2011;Sturm et al., 2014) and (3) in inflow zones of reservoirs compare to the main body (Musenze et al., 2014). However, as it was shown for CO_2 emissions from a tropical hydroelectric reservoir, taking into account both spatial and temporal variability of emissions significantly affect carbon budgets and emission factors (Pacheco et al., 2015).

The spatial and temporal variability of CH_4 ebullitive fluxes were intensively studied at the newly flooded subtropical Nam Theun 2 Reservoir (NT2R) (Deshmukh et al., 2014) and downstream as well as total CH_4 emissions are described in Deshmukh et al. (2015). The objective of the present study is to quantify the CH_4 diffusive fluxes at the surface of NT2R. The CH_4 emissions were quantified fortnightly during three and a half year (May 2010 to December 2012) based on a monitoring of CH_4 concentrations that started in June 2009. This was performed at nine stations flooding different types of ecosystems. On the basis of these

92 results, we discuss the spatial and temporal variations of the CH_4 emissions by diffusive
93 fluxes and the significance of hotspots and hot moments in the total emissions from the
94 surface of the reservoir.

95 ## 2. Material and methods

96 ### 2.1. Study area

97 The NT2 hydroelectric reservoir (17° 59' 49" N, 104° 57' 08" E) was built on the Nam Theun
98 River located in the subtropical region of Lao People's Democratic Republic (Lao PDR) on
99 the Nakai Plateau. A detailed description of the study site is given in Descloux et al. (2014).
100 The filling of the reservoir began in April 2008, the full water level was first reached in
101 October 2009 and the power plant was commissioned in April 2010. Annually, the NT2
102 Reservoir receives around 7527 Mm^3 of water from the Nam Theun watershed, which is more
103 than twice the volume of the reservoir (3908 Mm^3). A continuous flow of 2 m^3 s^{-1} (and
104 occasionally spillway release) is discharged from the Nakai Dam (ND in Fig 1) to the Nam
105 Theun River. The water used for electricity production is delivered from water intake (WI in
106 Fig 1) to the powerhouse (PH in Fig 1). The powerhouse is located in the valley 200 m below
107 the plateau.

108 Typical meteorological years are characterized by three seasons: warm wet (WW) (mid June-
109 mid October), cool dry (CD) (mid October-mid February) and warm dry (WD) (mid
110 February-mid June). Daily air temperature varies between 14°C (CD season) to 30°C (WD
111 season). The mean annual rainfall is about 2400 mm and occurs mainly (80%) in the WW
112 season.

113 During the filling of the reservoir, 489 km^2 of soils and different types of vegetation
114 (Descloux et al., 2011) were flooded by the end of October 2008. The water level in the
115 reservoir was nearly constant from October 2008 to April 2010. After the commissioning,
116 during the studied period (June 2009 to December 2012) the reservoir surface varied
117 seasonally and reached its maxima (489 km^2) and minima (168 to 176 km^2 depending on the
118 years) during the WW and WD seasons, respectively. According to these water level
119 variations, the average depth is 8 m for a maximum depth of 39 m.

2.2. Sampling strategy

A total of nine stations (RES1-9, Figure 1) located in the reservoir were monitored fortnightly in order to determine the vertical profiles of physico-chemical parameters of the water column and the CH_4 concentrations. The characteristics of the stations are given in the Table 1. Basically, three stations are located on the thalweg of the former Nam Theun River (RES2, RES4, RES6) whereas four other stations are located in a small embayment in the flooded dense forest (RES3), flooded degraded forest (RES5), flooded swamp area (RES7) and flooded agricultural land (RES8). The RES1 station is located 100 m upstream of the Nakai Dam, and RES9 station is located ~1 km upstream of the water intake delivering the water to the powerhouse. All samples and in situ measurements were taken in the morning or early afternoon from an anchored boat. Most of the time, the boat was attached to a buoy at the sampling station. When no buoy was present, an anchor was used with care in order not to re-suspend surface sediments. As the sampling started from the surface, the bottom water was sampled almost an hour later and should not be influenced by the perturbation generated by the anchor.

2.3. Experimental methods

2.3.1. Vertical profiles of oxygen and temperature

Vertical profiles of O_2 and temperature were measured in situ at all sampling stations with a multi-parameter probe Quanta® (Hydrolab, Austin, Texas) since January 2009. In the reservoir, the vertical resolution was 0.5 m above the oxic–anoxic limit and 1 to 5 m in the hypolimnion.

2.3.2. Methane concentration in water

The evolution of CH_4 concentrations has been monitored from May 2009 to December 2012 on a fortnightly basis. Surface samples were taken with a surface water sampler (Abril et al., 2007) and other samples from the water column were taken with an Uwitec water sampler. (Abril et al., 2007). Water samples were stored in serum glass vials, capped with butyl stoppers, sealed with aluminium crimps and poisoned (Guerin and Abril, 2007). Before gas chromatography analysis for CH_4 concentration, a N_2 headspace was created and the vials were vigorously shaken to ensure an equilibration between the liquid and gas phases. The

concentration in the water was calculated using the solubility coefficient of Yamamoto et al. (1976).

2.3.3. Gas chromatography

Analysis of CH_4 concentrations were performed by gas chromatography (SRI 8610C gas chromatograph, Torrance, CA, USA) equipped with a flame ionization detector. A subsample of 0.5 ml from the headspace of water sample vials was injected. Commercial gas standards (10, 100 and 1010 ppmv, Air Liquid "crystal" standards) were injected after analysis of every 10 samples for calibration. Duplicate injection of samples showed reproducibility better than 5%.

2.4. Water column CH_4 storage

Between two sampling depth of the vertical profiles of CH_4 concentrations, the CH_4 concentrations were assumed to change linearly in order to calculate the concentration in each 1-m layer of water. The volume of water in each layer was calculated using the volume-capacity curve (NTPC, 2005). The CH_4 storage was calculated by multiplying the average CH_4 concentrations of each layer by the volume of the layer and summing-up the amount of CH_4 for all depth intervals.

2.5. Aerobic CH_4 oxidation

The depth-integrated CH_4 oxidation rates at each station were calculated on the basis of the specific oxidation rates (d^{-1}) determined at NT2 (Deshmukh et al., 2015) and the vertical profiles of CH_4 and O_2 concentrations in the water column as already described in (Guerin and Abril, 2007). The depth-integrated CH_4 oxidation rates at each station were estimated only from January 2010 since the vertical resolution of the vertical profiles of O_2 and CH_4 was not high enough in 2009.

2.6. Estimation of diffusive fluxes from surface concentrations

The diffusive CH_4 fluxes were calculated from the fortnightly monitoring of surface concentrations with the thin boundary layer (TBL) equation at all stations in the reservoir (RES1-9). The CH_4 surface concentrations in water and the average CH_4 concentration in air (1.9 ppmv) obtained during eddy covariance deployments (Deshmukh et al., 2014) were applied in equation (1) to calculate diffusive flux:

$$F = k_T \times \Delta C \quad (1)$$

where F, the diffusive flux at water-air interface; k_T, the gas transfer velocity at a given temperature (T); $\Delta C = C_w - C_a$, the concentration gradient between the water (C_w) and the concentration at equilibrium with the overlying atmosphere (C_a). Afterward, the k_T was computed from k_{600} with the following equation:

$$k_T = k_{600} \times (600/Sc_T)^n \quad (2)$$

with Sc_T, the Schmidt number of CH_4 at a given temperature (T) (Wanninkhof, 1992); n, a number that is either 2/3 for low wind speed (< 3.7 m s^{-1}) or 1/2 for higher wind speed and turbulent water (Jahne et al., 1987).

For the determination of k_{600} at the stations RES1-8, we used both the formulations from Guerin et al. (2007) which includes the cumulative effect of wind (U_{10}) and rain (R) on k_{600} ($k_{600} = 1{:}66e^{0{:}26U10} + 0{:}66R$), and the average formulation of MacIntyre et al. (2010) (($k_{600} = 2.25\ U_{10} + 0.16$) whatever the buoyancy fluxes. As shown by (Deshmukh et al., 2014), the average of the fluxes obtained from these two relationships compared well with fluxes measured by floating chambers at the reservoir surface and no enhancement of the CH_4 fluxes could have been attributed to the variations of buoyancy fluxes when the eddy covariance system was deployed. Since the water current velocities were lower than 1 cm s^{-1} in most of the reservoir (Chanudet et al., 2012), the effect of water current on k_{600} was not included. For calculation purpose, wind speed (at 10 m height) and rainfall from two adjacent meteorological stations located at Nakai Village (close to RES9 station) and at the Ban Thalang Bridge (close to RES4 station, Figure 1) were used. At these stations, the average k_{600} was 6.5 cm h^{-1} over the course of the year.

At the water intake (RES9) where the hydrology and hydrodynamics is different from the other stations, it was impossible to quantify the k_{600} since the boat drifted quickly to the shoreline because of water currents in the narrow channel. According to Chanudet et al. (2012), water current velocity in this area of the reservoir is about 0.2 m s^{-1}. After Borges et al. (2004), the contribution of such water currents in a water body with depth ranging from 9 to 20 m is 2.0±0.5 cm h^{-1} which should be summed up with the contribution of wind and rainfall from Guerin et al. (2007) and MacIntyre et al. (2010). It gives an average of 9 cm h^{-1}. The k_{600} was determined in the regulating dam (Deshmukh et al., 2014) located downstream of the turbine where we visually observed vortexes similar to those observed at RES9. In the

209 regulating dam, the k_{600} was 19 cm h^{-1} on average for 4 measurements (not show). In order to
210 be conservative for the estimation of emissions from the water intake, we considered a
211 constant value of k_{600} (10 cm h^{-1}) which is in the lower range of (1) the k_{600} calculated from
212 (Guerin et al., 2007), MacIntyre et al. (2010) and Borges et al. (2004), and (2) k_{600} values
213 determined in area with comparable hydrology/hydrodynamics.

214 **2.7. Extrapolation of fluxes for the estimation of the NT2 total emissions**

215 Based on physical modelling (Chanudet et al., 2012), it has been showed that the station
216 RES9 located at the water intake is representative of an area of ~3 km^2 (i.e. 0.6% of reservoir
217 water surface), whatever the season. This 3-km² area was used to extrapolate specific
218 diffusive fluxes from RES9. The embayment where RES3 is located represents a surface area
219 of 5-6% of the total surface area of the reservoir whatever the season (maximum 28 km^2), to
220 which were attributed the specific diffusive fluxes from RES3. The diffusive fluxes calculated
221 for RES1, RES2, RES4, RES5, RES6, RES7 and RES8 stations were attributed to the water
222 surface area representative for each station, taking into account the seasonal variation of the
223 reservoir water surface from the surface-capacity curve (NTPC, 2005).

224 **2.8. Statistical and correlation analysis**

225 Statistical tests were performed to assess the spatial and temporal variations in the surface
226 CH_4 concentrations and diffusive fluxes at all stations in the reservoir. Normality of the
227 concentration and diffusive datasets was tested with R software (R Development Core Team,
228 2008) and the Nortest package (Gross and Ligges, 2015). The data distribution was tested
229 with the Fitdistrplus package (Delignette-Muller et al., 2015).

230 Since all tests indicated that the distribution of the data were neither normal nor lognormal,
231 Kruskal-Wallis and Mann-Whitney tests were performed with GraphPad Prism (GraphPad
232 Software, Inc., v5.04). No significant differences were found between the seasons and/or the
233 stations. These test results were attributed to the very large range of surface concentrations
234 due to the sporadic occurrence of extreme values (over 4 orders of magnitude). In order to
235 reduce this range, the log of the concentrations was used. For each station, the time series of
236 the log of the CH_4 surface concentrations were linearly interpolated and re-sampled every 15
237 days in order to compare time series with the same number of observations. The log of the
238 concentrations was used to determine the frequency distribution, the skewness of the dataset

(third order moment), the auto-correlation of each time series and the correlation between the different stations. All analyses were performed using Matlab.

3. Results

3.1. Temperature and O_2 dynamics in the reservoir water column

During the three and half year of monitoring at the stations RES1-8, the NT2R was thermally stratified with a thermocline at 4.5 ± 2.6 m depth in the WD (Feb-Jun) season as revealed by the vertical profiles of temperature (Figure 2). In the WW season, the temperature vertical profiles at the stations RES1-8 either showed a thermocline (RES7 and RES8 in 2010 and 2011, Figure 2) whereas in some occasions, the temperature decreased regularly from the surface to the bottom during sporadic destratification (RES1-3, Figure 2). On average during the WW season, a thermocline was located at 5.8 ± 4.8 m depth. During the CD season, the reservoir overturned as already mentioned by Chanudet et al. (2012) and the temperature was constant from the surface to the bottom (Figure 2) in the different years. In order to illustrate the destratification, a stratification index (ΔT) which corresponds to the difference between the surface and bottom water temperature was defined. During the periods of stratification in the WD seasons, ΔT was up to 10°C higher than during reservoir overturn in the CD season with ΔT close to zero (Figure 3a). During the WW season, the ΔT decreased gradually.

During the WD season at the stations RES1-8, an oxicline was most of the time located at a depth concomitant with the depth of the thermocline whereas oxygen penetrated deeper in the WW season (Figure 2). During these two seasons, the epilimnion was always well oxygenated with O_2 concentrations higher than 200 μmol L^{-1}. In the WD season, the hypolimnion was completely anoxic whereas O_2 reached occasionally the hypolimnion during the sporadic destratification events in the WW season (29±54 μmol L^{-1}, Figure 2 and 3b). During the CD season (reservoir overturn), the water column was often oxygenated from the top to the bottom of the reservoir (Figure 2). On average over the whole reservoir, the lowest hypolimnic oxygen concentration was observed in 2010 before the reservoir was commissioned (Figure 3b).

After the commissioning of the reservoir (April 2010), the water column located near the water intake (RES9) got totally mixed as revealed by the homogeneous temperature and oxygen profiles from the surface to the bottom whatever the season (Figure 2). The water column at RES9 was always well oxygenated (163 ± 62 μmol L^{-1}, Figure 2).

3.2. Seasonal dynamics of the CH_4 concentration and storage in the reservoir water column

At the station RES1-8, when the water column is thermically stratified with a steep oxicline in the WD and often in the WW seasons, CH_4 concentrations are in average ~150 times higher in the reservoir hypolimnion (246 ± 234 µmol L^{-1}) than in the epilimnion (1.6 ± 7.7 µmol L^{-1}) (Figure 2). The gradient of CH_4 concentration at the thermocline/oxicline was steeper during the WD season than during the WW season (Figure 2). During the CD season, the average CH_4 concentration in the reservoir bottom water lowered by a factor of three compare to the WD and the WW seasons. However, the reservoir overturn increased the average CH_4 concentrations in the epilimnion by a factor of two (3.4 ± 14.8 µmol L^{-1}) in comparison with the WD and WW seasons. After the commissioning, the CH_4 vertical profiles of concentration before turbine intake (RES9) were homogeneous from the surface to the bottom. The average CH_4 concentration from the surface to the bottom peaked up to 215 µmol L^{-1} with averages of 39.8 ± 48.8, 29.9 ± 55.4 and 1.9 ± 4.3 µmol L^{-1} during the WD, WW and CD seasons, respectively (Figure 2). The concentrations at RES9 were up to 10 times lower than the maximum bottom concentrations at the other stations for a given season. Since the station RES9 behaved differently from the other stations, results from this station will be treated separately.

The overall bottom CH_4 concentration (Figure 3c) and dissolved CH_4 stock in the reservoir (Figure 3d) increased at the beginning of the WD season. The higher bottom CH_4 concentration and storage in the reservoir are concomitant with both the establishment of anoxia in the hypolimnion and the reservoir thermal stratification (Figure 3). Hypolimnic CH_4 concentration and storage reached their maxima (up to 508 ± 254 µmol L^{-1} and 4.7 ± 0.5 Gg(CH_4), Figure 3c,d) at the end of the WD-beginning of the WW season when the residence time of water in the reservoir was the lowest (40 days, Figure 3d). Along the WW season, the thermal stratification weakened (Figure 3a) and the CH_4 concentration and dissolved CH_4 stock decreased (Figure 3c,d) while the residence time of water increased (Figure 3d). In the CD season, the reservoir overturns as evidenced by the low ΔT and the penetration of O_2 to the hypolimnion (Figure 3a,b). During CD season, the bottom CH_4 concentration and the storage reached their minima (down to 1.3 ± 4.5 µmol L^{-1} and 0.01 ± 0.001 Gg(CH_4), Figure 3c,d) when the residence time of water was the longest (Figure 3d). The sharp decrease of CH_4 storage and concentration in the transition from the WW to the CD seasons is

concomitant with a sharp increase of O_2 concentration at the bottom (up to 160 ± 89 μmol L^{-1}, Figure 3).

3.3. Interannual variations of the CH_4 concentrations and storage in the reservoir water column

During the three and a half years of monitoring, the same seasonal pattern is observed although the annual CH_4 bottom concentration and storage was threefold higher in 2009 and 2010 than in the year 2011 (Figure 3c,d). In the dry year 2012, the reservoir bottom CH_4 concentration and storage was almost twice higher than in wet year 2011.

3.4. Aerobic CH_4 oxidation in the reservoir

Between 2010 and 2012, the depth integrated aerobic CH_4 oxidation rates ranged between 0.05 and 380 mmol m^{-2} d^{-1} at the stations RES1-RES8 (Figure 4). On average, aerobic oxidation was higher in the WW season (55±63 mmol m^{-2} d^{-1}) than in the CD (30±46 mmol m^{-2} d^{-1}) and WD (36±32 mmol m^{-2} d^{-1}) seasons and it was not statistically different for the three years. In the WD season, aerobic CH_4 oxidation was on average twice higher in 2010 than for the two following years whereas in the CD season, the highest aerobic oxidation rate was observed in 2012.

3.5. Spatial and seasonal variability of surface CH_4 concentration and diffusive fluxes at the reservoir surface (RES1-RES8)

The surface concentrations at the stations RES1-8 ranged from 0.02 to 150 μmol L^{-1} and were 2.0±10.5 μmol L^{-1} (median = 0.9), 1.5±5.5 μmol L^{-1} (median = 0.4) and 3.4±14.7 μmol L^{-1} (median = 0.2) on average for the CD, WD and WW season, respectively. The surface concentration followed a loglogistic distribution, which indicates the existence of extremely high values. This is confirmed by the fact that the skewness of the time series of the log of the CH_4 concentrations for all stations is positive (Figure S3), especially at the stations RES1, RES3 and RES7 for which the skewness is >1. Over the course of the three and a half year of survey, the surface concentrations were not statistically different between all stations and no statistically significant seasonal variations were observed because of the occurrence of sporadic events at all season (Figure S2a). The normalized distribution of concentrations (in log) according to seasons (Figure 5) indicates that these high concentrations were observed without any clear seasonal trend at the station RES1, RES5 and RES6 (<1 up to 150 μmol L^{-}

^{1}). At the stations RES2 and RES3, the concentrations up to 128 μmol L^{-1} were mostly observed in the CD season when the reservoir overturns. At the station RES4 located at the Nam Xot and Nam Theun confluence and at the stations RES7 and RES8 both located in the inflow region of the Nam Theun River, the high surface concentrations (up to 64.60 μmol L^{-1}) were mostly observed during the WW season when the reservoir undergoes sporadic destratification. The auto-correlation function of the time series of the log of the surface CH_4 concentrations and diffusive fluxes at the stations RES1-8 indicated that at all stations (except RES1) have a memory effect of 30 to 40 days (Figure S1). This implies that with a sampling frequency of 15 days, we captured most of the changes in the surface CH_4 concentrations. At the station RES1, the changes in CH_4 concentrations are faster than at other stations and would have deserved a monitoring with a frequency higher than 15 days.

During the monitoring at RES1-RES8 stations, the average diffusive flux was 2.8 ± 12.2 mmol m^{-2} d^{-1} ranging from 0.01 to 201.86 mmol m^{-2} d^{-1} without any clear interannual and seasonal trends (Figure S2b). As for the concentrations, flux data followed a loglogistic distribution. The median flux in the WD season is 40 to 80% higher than the median in the WW and CD season, respectively. However, the average fluxes in the WW and CD season are 30% higher than in the WD season (Table 2). This confirms the presence of extremely high values during WD and CD seasons, as expected from the surface concentrations. All seasons together, around 7% of the diffusive fluxes that we observed were higher than 5 mmol m^{-2} d^{-1} which corresponds to extremely high diffusive fluxes in comparison with data from the literature for reservoirs and lakes (Bastviken et al., 2008;Barros et al., 2011). The median and average of these extreme fluxes higher than 5 mmol m^{-2} d^{-1} were 2 times higher in the WW and CD seasons than in the WD season (Table 2).

At NT2, diffusive CH_4 fluxes covered the whole range of fluxes reported for tropical reservoirs, depending on the season. Most of the fluxes at the NT2R Reservoir were around one order of magnitude lower than the ones at Petit Saut Reservoir (French Guiana) just after the impoundment (Galy-Lacaux et al., 1997), and in the same order of magnitude as reported for reservoirs older by 10 to 18 years (Abril et al., 2005;Guerin et al., 2006;Kemenes et al., 2007;Chanudet et al., 2011). However, some diffusive fluxes at the stations RES1-8 in the WW and the CD seasons (up to 202 mmol m^{-2} d^{-1}) are among the highest ever reported at the surface of a hydroelectric reservoir or a lake (Bastviken et al., 2011;Barros et al., 2011) and rivers downstream of dams (Abril et al., 2005;Guerin et al., 2006;Deshmukh et al., 2015).

3.6. Surface methane concentrations and diffusive fluxes at the water intake (RES9)

After the commissioning of the reservoir (Julian day 450), the concentrations at the stations RES9 (Figure 6a) located at the water intake were up to 30 times higher than at any other stations that is 36.6±35.8 μmol L^{-1} (median = 24.3), 37.6±67.0 μmol L^{-1} (median = 0.9) and 1.0±1.7 μmol L^{-1} (median = 0.3) in the WD, WW and CD season, respectively. The surface concentrations at RES9 were significantly higher in the WD and WW seasons than in the WW and CD seasons ($p = 0.0002$ and Figure 6a). The highest concentration was observed each year at the end of the WD season-beginning of the WW season in between June and August. These maxima decreased from 215 μmol L^{-1} in August 2010 to 87 μmol L^{-1} in June 2012.

The diffusive fluxes ranged between 0.03 and 605.38 mmol m^{-2} d^{-1} (Figure 6b and Table 2). On average, the CH_4 diffusive fluxes at RES9 were two to forty times higher than at the other stations in the CD, WD and WW season. Diffusive fluxes at this station are usually higher than 10 mmol m^{-2} d^{-1} from April to July that corresponds to the WD season and the very beginning of the WW season. In 2010, diffusive fluxes were on average 241 ± 219 and 239 ± 228 mmol m^{-2} d^{-1} respectively for the WD and WW seasons. In 2011 and 2012, the fluxes dropped down by a factor of two in the WD season (112 ± 110 mmol m^{-2} d^{-1}) and almost by a factor of forty in the WW season (6.8 ± 14.4 mmol m^{-2} d^{-1}). Overall, emissions at RES9 decreased by a factor of two between 2010 and 2012.

At the water intake, CH_4 diffusive fluxes during the transition between the WD and WW seasons (up to 600 mmol m^{-2} d^{-1}) are the highest reported at the surface of an aquatic ecosystem (Abril et al., 2005;Guerin et al., 2006;Bastviken et al., 2011;Barros et al., 2011;Deshmukh et al., 2015).

4. Discussion

4.1. CH_4 dynamic in the reservoir water column

The gradual decrease of the CH_4 concentration from the anoxic bottom water column to the metalimnion and the sharp decrease around the oxicline in the metalimnion (Figure 2) is typical in reservoirs and lakes where CH_4 is produced in anoxic sediments and flooded soils (Guerin et al., 2008;Sobek et al., 2012;Maeck et al., 2013), and where most of it is oxidized at the oxic-anoxic interface (Bedard and Knowles, 1997;Bastviken et al., 2002;Guerin and Abril, 2007;Deshmukh et al., 2015).

CH_4 concentrations and storage increase concomitantly with the surface water temperature and the establishment of the thermal stratification during the WD season and peak at the end of the WD season-beginning of the WW season (Figure 2 and 3). During the WW season, CH_4 concentrations and storage decrease slowly (Figure 3) while aerobic methane oxidation reaches its maximum (Figure 4). When the reservoir overturns at the beginning of the CD season, the CH_4 hypolimnic concentrations and storage reach their minima (Figure 3). The overturn favours the penetration of oxygen down to the bottom (Figure 2 and 3b). The sharp decrease of the CH_4 concentrations and CH_4 storage during this period is expected to result from sudden outgassing (Section 4.2) together with an enhancement of the aerobic CH_4 oxidation as already observed in lakes that overturn (Utsumi et al., 1998b;Utsumi et al., 1998a;Kankaala et al., 2007;López Bellido et al., 2009;Schubert et al., 2010;Schubert et al., 2012;Fernández et al., 2014). A large increase of the aerobic methane oxidation was only observed in the CD season in the dry year 2012 (Figure 4) because the amount of hypolimnic CH_4 to be oxidized at the beginning of the CD season was still high in the water column (Figure 3c,d).

As the reservoir overturns during the period over which the water residence time is the longest in the reservoir, the temporal evolution of the concentrations is anti-correlated with the residence time (Figure 3c,d). The seasonal dynamics of the CH_4 in the monomictic NT2R differs from permanently stratified reservoirs like Petit Saut Reservoir where CH_4 concentration increased with retention time (Abril et al., 2005). However, at the annual scale the water residence time has a strong influence on CH_4 concentration and storage in the reservoir. Before the reservoir was commissioned (April 2010), the water residence time was up to 4 years and the CH_4 storage was up to four times higher than in 2011 and 2012 (Figure 3d). Although a decrease of concentration and storage with the age of the reservoir was expected (Abril et al., 2005), the storage in the dry year 2012 was twice higher than in the wet year 2011 due to an increase of the water residence time by 25% between 2011 and 2012. In wet years like 2011, the thermal stratification is weaker than in dry years since the warming of surface water is less efficient and the high water inputs alters the stability of the reservoir thermal stratification as shown by the sharper decrease and the larger range of ΔT in 2011 than in 2012 (Figure 3a). As a consequence, the oxygen diffusion to the hypolimnion was higher in 2011 than in 2012 (Figure 3b) and it enhanced aerobic methane oxidation by 20% in the water column in the WW season in 2011 as compared to 2012 (Figure 4). It therefore suggests that the hydrology affects both the thermal stratification and the hypolimnic storage

of CH_4 in reservoirs, indirectly controls aerobic methane oxidation, and ultimately influences emissions.

4.2. Hot moments of emissions during sporadic destratification and reservoir overturn

The figure 7 illustrates the evolution of the diffusive fluxes, the stratification index (ΔT), the CH_4 storage and the aerobic CH_4 oxidation at the stations RES1, RES3, RES7 and RES8. These four stations were selected for their contrasting skewness (Figure S3) which gives an indication on the occurrence of extreme events and the facts that they are representative for all station characteristics (Table 1). It shows that the large bursts of CH_4 (from 5 up to 200 mmol m^{-2} d^{-1}) always occurred when ΔT decreased sharply (>4°C, Figure 7a,d,g,j) and are usually followed by a sharp decrease of the CH_4 storage in the water column (Figure 7b,e,h,k). These hot moments of emissions occurred mostly in the CD at the stations RES1 and RES3 whereas it was in the WW season at the stations RES7 and RES8 (Figure 7). In the WD season, diffusive fluxes gradually increased together with the CH_4 storage in the water column (Figure 7a,d,g,j) and they remained always lower than 20 mmol m^{-2} d^{-1}. These sporadic high fluxes occurred in the WD season at RES3, RES7 and RES8 (Figure 7d,g,j). They are usually associated with ΔT variations lower than 2°C and the CH_4 storage decrease that is associated with these fluxes is not as sharp as the one observed in the CD and WW season (Figure 7e,h,k).

We therefore confirm the occurrence of hot moments of emissions during the reservoir overturn in the CD season as already observed in lakes that overturn in temperate regions (Kankaala et al., 2007;López Bellido et al., 2009;Schubert et al., 2010;Schubert et al., 2012;Fernández et al., 2014). The highest emissions determined at NT2R are one order of magnitude higher than previously reported outgassing during overturn and they occur mostly in the section of the reservoir that has the longest water residence time (RES1-3, Table 1) and the largest CH_4 storage (Figure 7b,e,h,k). This suggests that the impact of reservoir overturn can be very critical for the whole-reservoir CH_4 budget in tropical hydroelectric reservoirs and especially in young ones where hypolimnic concentration could reach up to 1000 μmol L^{-1}. Hot moments of emissions also occur during sporadic destratifications in the WW season in the inflow region (RES4 and RES6-8) where the inflow of cool water from the watershed might disrupt the thermal stratification in reservoirs. This is contrasting with the observations in older reservoir than NT2R where high emissions from the inflow region were recently attributed to an enhancement of CH_4 production fuelled by the sedimentation of organic

459 matter from the watershed (Musenze et al., 2014). The high emissions in the WD seasons
460 were associated with early rains and associated high winds that occur sometimes in the last
461 fifteen days of May. This shows that a moderate erosion of the stratification when hypolimnic
462 CH_4 concentrations are high could enhance vertical transport of CH_4 toward the surface and
463 emissions to the atmosphere. Basically, this intense monitoring shows that spatial and
464 temporal variations of CH_4 emissions are largely controlled by the hydrodynamics of the
465 reservoir with extreme emissions occurring mostly in the inflow region during the wet season
466 and mostly in area remotely located from the inflow zone and the riverbed during reservoir
467 overturns in the CD season. Even if less frequent, moderate erosion of the stable and steep
468 thermal stratification during warm seasons, could also lead to high emissions.

469 The evolution of depth-integrated aerobic CH_4 oxidation is not clearly related with the
470 reservoir overturns and the CH_4 burst (Figure 7). Significant increases in the aerobic CH_4
471 oxidation occurred mostly during the first half of the WD season when the stratification was
472 unstable and at the very beginning of the destratification in the WW, when ΔT started to
473 decrease. The oxidation could reach high values (up to 380 mmol m^{-2} d^{-1}) during these two
474 periods since the yield of CH_4 in the water column to sustain the activity of methanotrophs is
475 higher than in the CD season when the reservoir overturns. It shows that in reservoirs or lakes
476 like NT2R that destratify progressively before the overturn, there is no substantial increase of
477 the CH_4 oxidation when the water body overturns as it could be observed in lakes that
478 overturn within a few days (Kankaala et al., 2007). In addition, the contribution of CH_4
479 oxidation to the total loss of CH_4 (sum of diffusion and oxidation) in the WD and WW
480 seasons was 90-95% during the entire monitoring whereas it was 85% in the CD season.
481 During overturns, a significant amount of CH_4 is oxidized (Utsumi et al., 1998a;Utsumi et al.,
482 1998b;Kankaala et al., 2007;Schubert et al., 2012) but it also indicates that the removal of
483 CH_4 during overturn is not as efficient as during seasons with a well established thermal
484 stratification.

485 During the periods with major loss in the CH_4 storage with concomitant CH_4 burst, we
486 compared the change in the yield of CH_4 with the sum of emissions and oxidation. Most of
487 the time, the emissions alone and/or the sum of emissions and oxidation were significantly
488 higher than the amount of CH_4 that was lost from the water column. At the Pääjäri Lake in
489 Finland (López Bellido et al., 2009), the fact that measured or calculated emissions exceed the
490 loss of CH_4 in the water column was attributed to a probable underestimation of the CH_4

storage in the lake by under-sampling the shallow area of the lake. In this study, emissions, storage and oxidation were estimated at the same stations, avoiding such sampling artefacts. Therefore, it suggests that CH_4 is provided by lateral transport or by production in the flooded soil and biomass (Guerin et al., 2008) at a higher rate than the total loss of CH_4 from the water column by emissions and oxidation. This hypothesis could only be verified by a full CH_4 mass balance including production and total emissions from the reservoir, which is beyond the scope of this article.

4.3. Hot spot of emissions at the water intake (RES9)

After the commissioning of the reservoir, the temperature and the oxygen and CH_4 concentrations were constant from the surface to the bottom of the reservoir at the vicinity of the water intake. On the basis of physical modelling and measurements of water current velocities (Chanudet et al., 2012), the vertical mixing at this station was attributed to the water withdrawal at the intake generating turbulence and water currents over a surface area of 3 km^2. At this station, CH_4-rich water from the reservoir hypolimnion reached the surface and led to diffusive fluxes up to 600 mmol m^{-2} d^{-1} in the WD-WW seasons (Figure 6b) whereas fluxes are 3 orders of magnitude lower in the CD season. To the best of our knowledge, this is the first time that a hotspot of emissions is reported upstream of a dam or an intake bringing water to the turbines. At NT2, the intake is located at the bottom of a narrow and shallow channel (depth =9-20 m) on the side of the reservoir. This design enhances horizontal water current velocities, the vertical mixing and therefore the emissions. The existence of such a hotspot at other reservoirs might be highly dependant on the design of the water intake (depth among other parameters) and its effect on the hydrodynamics of the reservoir water column.

4.4. Estimation of total diffusive fluxes from the reservoir

Yearly emissions by diffusive fluxes peaked at more than 9 Gg(CH_4) in 2010 when the reservoir was commissioned and they decreased down to ≈ 5 Gg(CH_4) in 2011 and 2012 (Figure 8a and Table 3). Yearly integrated at the whole reservoir surface, these emissions correspond to diffusive fluxes of 1.5 to 4 mmol m^{-2} d^{-1}. These emissions are significantly lower than diffusive fluxes measured at the Petit Saut Reservoir during the first two years after flooding but similar to those determined in the following years (Abril et al., 2005) and values reported for diffusive fluxes from tropical reservoirs in Barros et al. (2011). In absence of the extreme emissions (both hotspots and hot moments), diffusive emissions from NT2R

522 would have been one order of magnitude lower than emissions from tropical reservoirs as
523 expected from the lower flooded biomass compare to Amazonian reservoirs (Descloux et al.,
524 2011). Due to the specific dynamic of diffusive fluxes at NT2R, diffusion at the reservoir
525 surface contribute 18 to 27% of total emissions (Table 3) that is significantly higher than at
526 other reservoirs tropical reservoirs where it was measured (See Deshmukh et al., 2015 for a
527 detailed discussion).

528 Most of the increase of CH_4 emissions by diffusive fluxes from 4 to 9 Gg(CH_4) between 2009
529 and 2010 is due to very significant emissions of 2-3 Gg(CH_4) at the water intake (Figure 8a).
530 This outgassing of CH_4 was triggered by the vertical mixing generated by the withdrawal of
531 water from the reservoir to the turbines. Although the area under the influence of the water
532 intake is less than 1% of the total area of the reservoir, emissions at the water intake
533 contributed between 13 and 25% of total diffusive emissions and 4 to 10 % if considering
534 both ebullition and diffusion (Table 3). It is worth to note that emissions at this site are only
535 significant within 3-5 month per year at the end of the WD season-beginning of the WW
536 season when the storage of CH_4 reach its maximum in the reservoir (Figure 8b). This new
537 hotspot equals 20 to 40% of downstream emissions and contributes between 4 and 7% of total
538 emissions from the NT2 reservoir surface when including ebullition and downstream
539 emissions (Table 3 and Deshmukh et al. (2015)). Very localized perturbation of the
540 hydrodynamics, especially in lakes or reservoirs with CH_4-rich hypolimnion, can generate
541 hotspots of emissions contributing significantly to the total emissions from a given ecosystem.
542 These hotspots could be found upstream of dams and water intake in reservoirs but also
543 around aeration stations based on air injection or artificial mixing that could be used for
544 improving water quality in water bodies (Wüest et al., 1992).

545 The contribution of extreme diffusive fluxes (> 5 up to 200 mmol m^{-2} d^{-1}) to total emission by
546 diffusion range from 30 to 50% on a yearly basis (Figure 8a) and from 40 up to 70% on a
547 monthly basis (Figure 8b) although these hot moments represent less than 10% of the
548 observations during the monitoring. In the literature, the statistical distribution of CH_4
549 emissions dataset always follows heavy-tailed and right skewed distribution like the log-
550 normal, the Generalized Pareto Distribution (Windsor et al., 1992;Czepiel et al., 1993;Ramos
551 et al., 2006;DelSontro et al., 2011) or loglogistic (this study) which indicates that CH_4
552 emissions are always characterized by high episodic fluxes. The quantification of emissions
553 thus requires the highest spatial and temporal resolutions in order to capture as many hot
554 moments as possible. At a single station, extreme emission events never lasted more than 2

555 months (3 consecutive sampling dates) and probably lasted less than 15 days most of the time
556 (Figure 7). The auto-correlation function of the concentration time series indicate that a
557 minimum sampling frequency of 1 month is required in this monomictic reservoirs for an
558 accurate description of the change in the surface concentrations and estimation of the
559 emissions (Figure S1). A lower temporal resolution can significantly affect (positively or
560 negatively) the emissions factors of non-permanently stratified freshwater reservoirs. This is
561 particularly critical in the inflow regions when water inputs from the watershed increase in the
562 rainy season in all reservoirs and at the beginning of the overturn in regions of the world
563 where reservoirs are not permanently stratified like in Asia (Chanudet et al., 2011) which
564 concentrate 60% of the worldwide hydroelectric reservoirs (Kumar et al., 2011).

565 ## *5. Conclusion*

566 The fortnightly monitoring of CH_4 diffusive emissions at nine stations revealed complex
567 temporal and spatial variations that could hardly been characterized by seasonal sampling.
568 The highest emissions occur sporadically during hot moments in the rainy season and when
569 the reservoir overturns. In the rainy season, they mostly occur in the inflow region because the
570 increase of the discharge of cool water from the reservoir tributaries contributes to sporadic
571 thermal destratification. During the reservoir overturn, extreme emissions occur mostly in
572 area remotely located from the inflows and outflows that are supposed to have the highest
573 water residence time. It shows that diffusive emissions can be sporadically as high as
574 ebullition and that these hot moments could contribute very significantly to the total emissions
575 from natural aquatic ecosystems and reservoirs. Our results showing that a monthly frequency
576 monitoring is the minimum required to capture all emissions is probably not applicable to
577 every aquatic ecosystem. However, it suggests that quantification of emissions based on 2-4
578 campaigns in a year might significantly affect emissions factors and carbon budgets of
579 ecosystems under study.

580 We also identified a new hotspot of emissions upstream of the water intake resulting from the
581 artificial destratification of the water column due to horizontal and vertical mixing generated
582 by the water withdrawal. In the case of the NT2R, emissions from this site contribute up to
583 25% of total diffusive emissions over less than 1% of the total reservoir area. We highly
584 recommend measurements of diffusive fluxes around water intakes (immediately upstream of
585 dams, typically) in order to evaluate if such results can be generalized.

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757
758

759
760 Table 1: Characteristics of the nine monitoring stations in the Nam Theun 2 Reservoir
761

Station	Flooded ecosystem[1]	Hydrology	Water residence time[2]
RES1	Dense forest	100 m upstream of the Nakai Dam	**
RES2	Dense forest	Thalweg of the Nam Theun River	**
RES3	Dense forest	Embayment	***
RES4	Degraded forest	Confluence Nam Theun-Nam Xot Rivers	**
RES5	Degraded forest	Aside from the main stream	**
RES6	Degraded forest	Thalweg of the Nam Theun River	*
RES7	Swamp	Between inflows and water intake	*
RES8	Agricultural soils	Between inflows and water intake	*
RES9	Civil construction	Water intake	*

762 [1]Descloux et al. (2011)

763 [2]Water residence time in arbitrary units, (***) stands for long residence time, (**) for
764 intermediate residence times and (*) for short residence times
765

766
767 Table 2 : Median, average, ranges and proportion of diffusive fluxes (F_{CH4}) < 1 and > 5 mmol
768 m^{-1} d^{-1} for three seasons
769

Station		Warm Dry (WD)	Warm Wet (WW)	Cool Dry (CD)
RES1-RES8	n	212	252	217
	range	0.01-102.59	0.01-201.86	0.01-94.64
	median	1.08	0.64	0.20
	Average ± SD	2.23±7.37	3.12±14.58	3.04±12.89
	% F_{CH4} < 1	48%	63%	86%
	% F_{CH4} > 5	6.6%	7.5%	7.4%
	Mediane F > 5	10.67	13.80	23.75
	Average F > 5	16.69±25.04	30.23±45.99	36.45±33.19
RES9	n	39	45	36
	range	0.24-342.00	0.03-605.38	0.07-17.62
	median	40.81	1.23	0.48
	average ± SD	83.33±15.57	78.58±24.73	2.21±0.69

770
771

772 Table 3: Methane emissions from the Nam Theun 2 Reservoir between 2009 and 2012.

$Gg(CH_4)$ $year^{-1}$	2009	2010	2011	2012
Emission from reservoir				
Diffusion at RES9 only	0.02±0.01	2.33±0.21	0.86±0.12	0.66±0.11
Total diffusion	4.45±1.01	9.34±2.32	3.71±0.81	4.95±1.09
Contribution of RES9 to diffusion (%)	0.4	24.9	23.2	13.3
Ebullition[1]	11.21±0.16	14.39±0.11	14.68±0.10	12.29±0.09
Total emissions from reservoir	15.66±1.02	23.73±2.32	18.39±0.82	17.25±1.09
Contribution of RES9 (%)	0.1	9.8	4.7	3.8
Total downstream emissions[2]	7.79±0.90	10.73±0.83	2.29±0.41	2.00±0.32
Total emissions (reservoir + downstream)	23.45±1.36	34.46±2.46	20.67±0.92	19.24±1.14
Contribution of diffusion to total emission	19%	27%	18%	26%
Contribution of RES9 to total (%)	<0.1	6.8	4.2	3.4

773 [1]Deshmukh et al. (2014)
774 [2]Deshmukh et al. (2015)
775
776

Figure captions

Figure 1: Map of the sampling stations and civil structures at the Nam Theun 2 Reservoir (Lao PDR).

Figure 2: Vertical profiles of temperature (°C), oxygen (μmol L^{-1}) and methane (μmol L^{-1}) at the stations RES1, RES3, RES7, RES8 and RES9 in the Nam Theun 2 Reservoir. Representative profile of the years 2010 (circle), 2011 (square) and 2012 (triangle) are given for each seasons: cool dry in blue, warm dry in red, and warm wet in grey.

Figure 3: (a) Stratification index (ΔT, see text), (b) O_2 concentration in the hypolimnion (μmol L^{-1}), (c) CH_4 concentration in the hypolimnion (μmol L^{-1}) and (d) CH_4 storage in the water column (Gg(CH_4) $month^{-1}$, bars) and water residence time (days, black line with circles) in the Nam Theun 2 Reservoir (Lao PDR) between 2009 and 2012. The red, grey and blue colours indicate the warm dry (WD), warm wet (WW) and cool dry (CD) seasons, respectively. For the panels (a), (b) and (c), the boxes show the median and the interquartile range, the whiskers denote the full range of values and the plus sign (+) denotes the mean.

Figure 4: Seasonal variations between 2010 and 2012 of the depth-integrated aerobic CH_4 oxidation (mmol m^{-2} d^{-1}) at the stations RES1-RES8 calculated from the aerobic oxidation rates determined by Deshmukh et al. (2015). WD stands for warm dry (in red), WW for warm wet (in grey) and CD for cool dry (in blue). The boxes show the median and the interquartile range, the whiskers denote the full range of values and the plus sign (+) denotes the mean.

Figure 5: Frequency distribution of the log of CH_4 concentrations (μmol L^{-1}) at the nine monitoring stations of the Nam Theun 2 Reservoir. The red, grey and blue colours indicate the warm dry (WD), warm wet (WW) and cool dry (CD) seasons, respectively.

Figure 6: (a) Surface concentrations and (b) diffusive fluxes between June 2009 and December 2012 at the station RES9 located at the water intake. Julian day 0 is 1st of January, 2009. The red, grey and blue colours indicate the warm dry (WD), warm wet (WW) and cool dry (CD) seasons, respectively.

Figure 7: (a, d, g, j) stratification index (ΔT, red line, see text) and diffusive fluxes, (b,e,h,k) CH_4 storage and (c,f,i,l) depth-integrated aerobic CH_4 oxidation (mmol m^{-2} d^{-1}, black line) calculated from the aerobic oxidation rates determined by Deshmukh et al. (2015) and ΔT (red line) between June 2009 and December 2012 at the stations RES1, RES3, RES7 and RES8 at the Nam Theun 2 Reservoir. Julian day 0 is 1st of January, 2009. The red, grey and blue colour dots indicate the warm dry (WD), warm wet (WW) and cold dry (CD) seasons, respectively.

Figure 8: (a) Total emissions by diffusive fluxes in 2009, 2010, 2011 and 2012, and (b) monthly emissions by diffusive fluxes between May 2009 and December 2012. Emissions from RES9 (water intake) are shown in black, emissions resulting from diffusive fluxes lower than 5 mmol m^{-2} d^{-1} from the stations RES1 to RES8 are shown in white and emissions resulting from diffusive fluxes higher than 5 mmol m^{-2} d^{-1} from the stations RES1-RES8 are shown in grey.

827
828 Figure 1
829

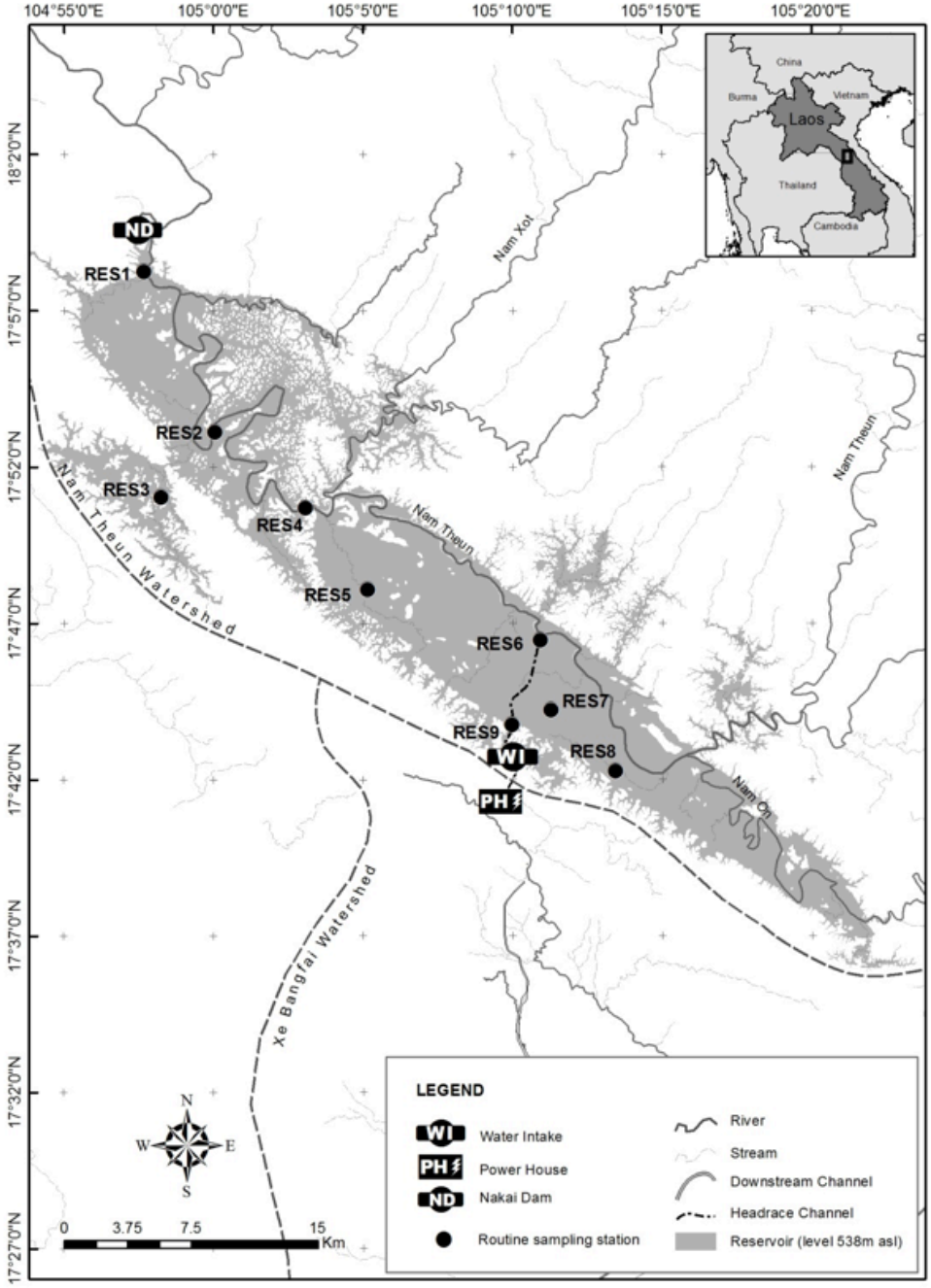

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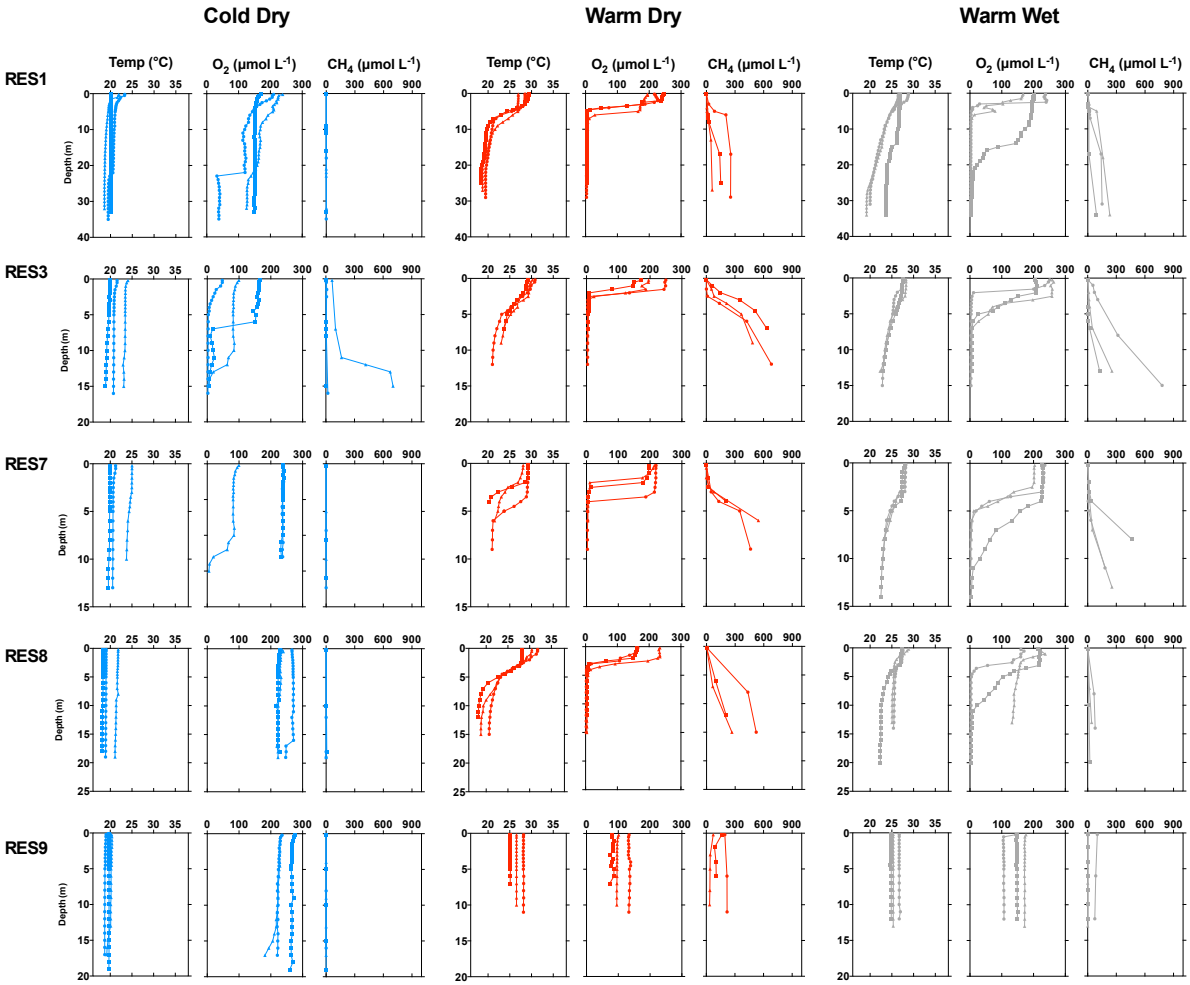
Figure 2


Cold Dry
Warm Dry
Warm Wet
RES1
RES3
RES7
RES8
RES9
Temp (°C)
O2 (µmol L-1)
CH4 (µmol L-1)
Depth (m)

839
840 Figure 3
841

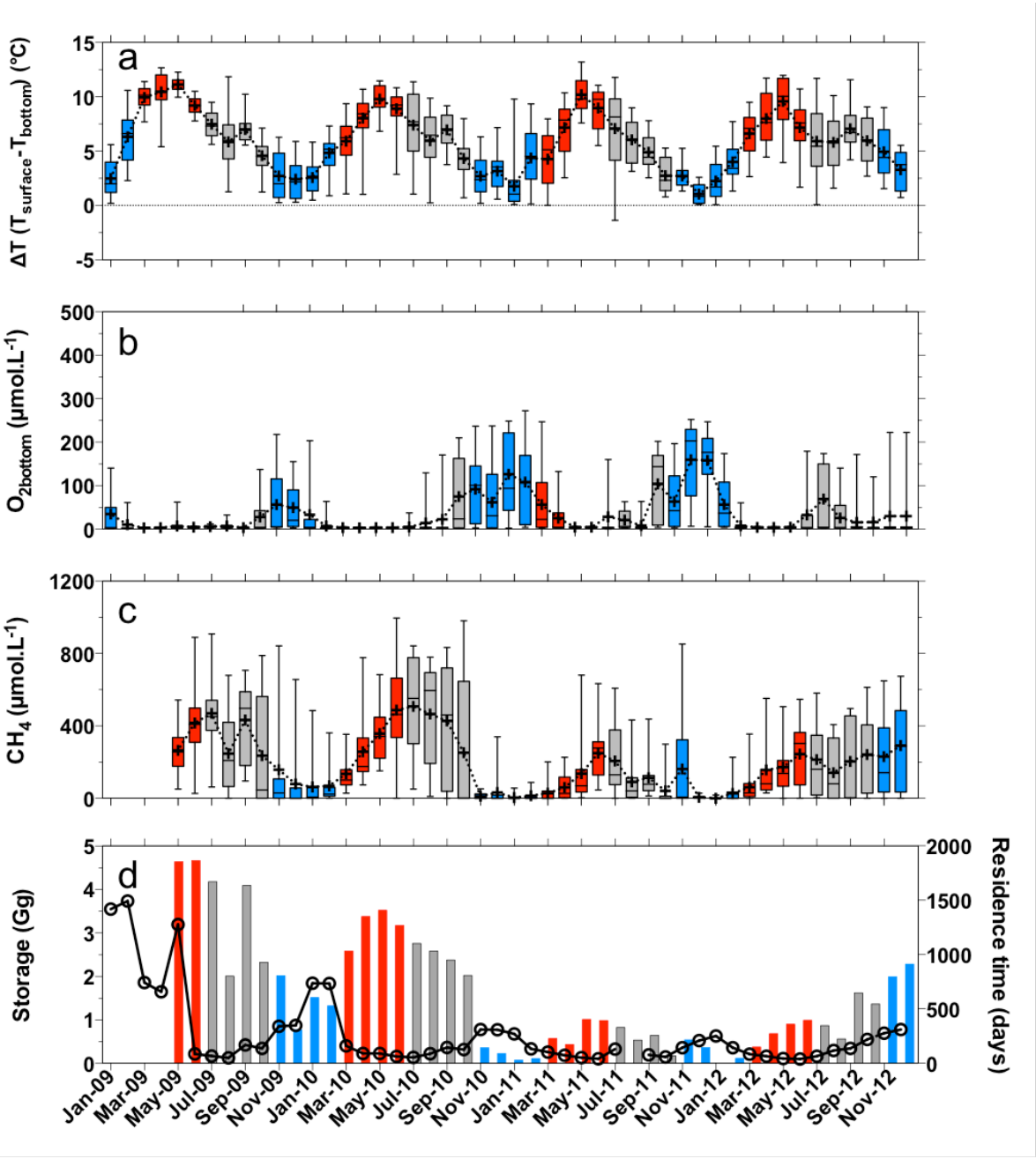


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Figure 4

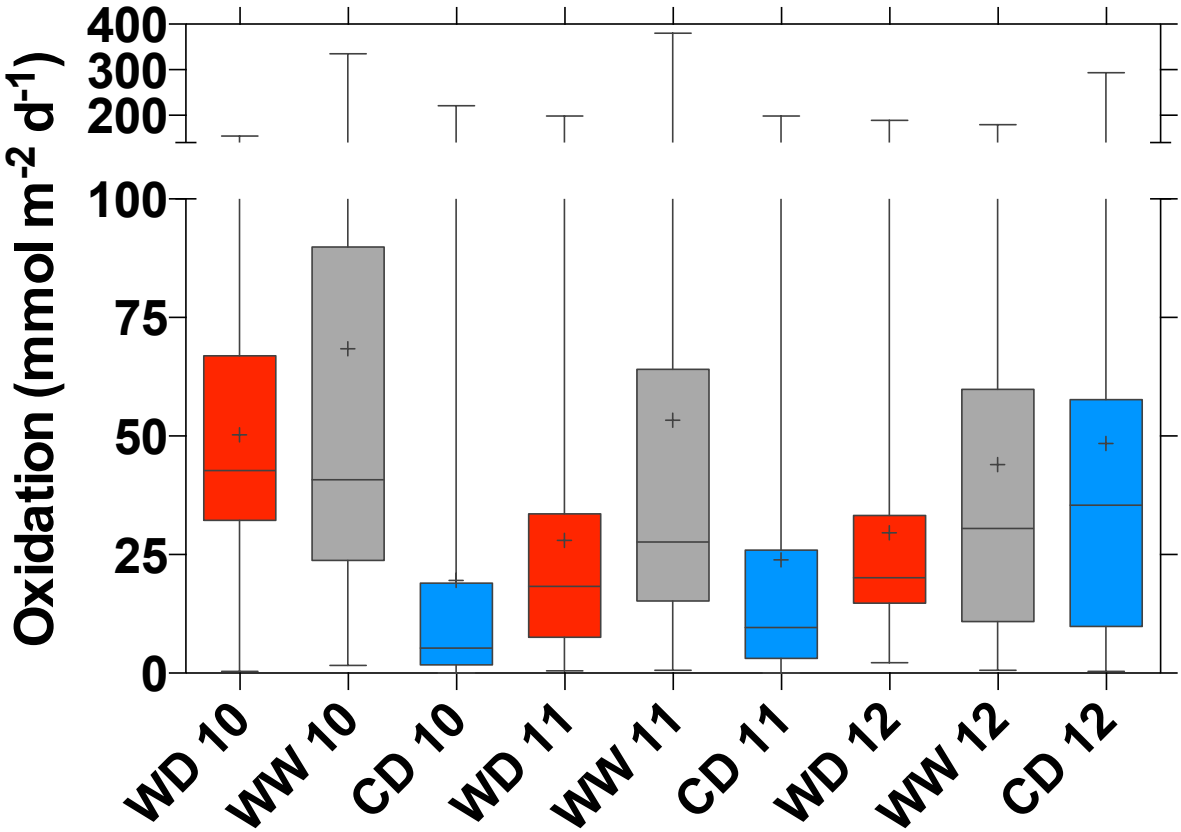
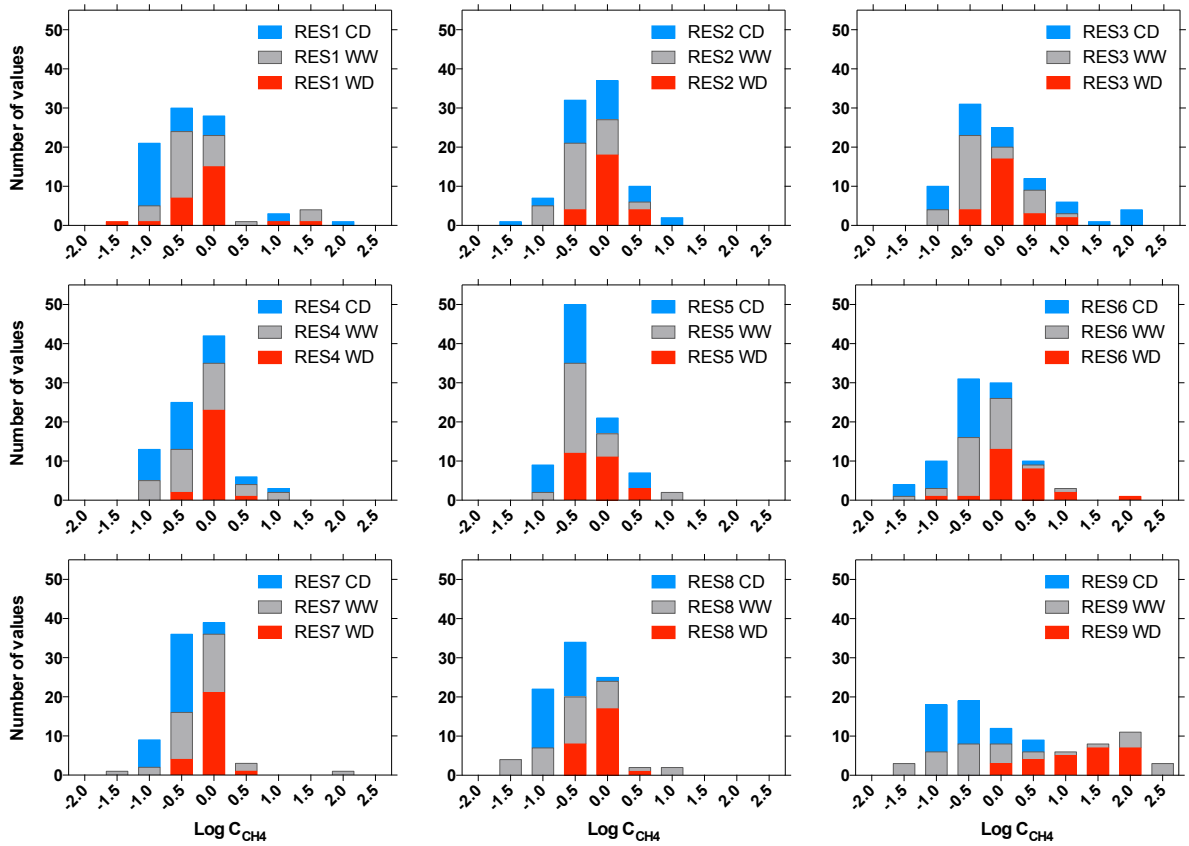

Oxidation (mmol m-2 d-1)
400
300
200
100
75
50
25
0
WD 10
WW 10
CD 10
WD 11
WW 11
CD 11
WD 12
WW 12
CD 12

Figure 5


RES1 CD
RES1 WW
RES1 WD
RES2 CD
RES2 WW
RES2 WD
RES3 CD
RES3 WW
RES3 WD
RES4 CD
RES4 WW
RES4 WD
RES5 CD
RES5 WW
RES5 WD
RES6 CD
RES6 WW
RES6 WD
RES7 CD
RES7 WW
RES7 WD
RES8 CD
RES8 WW
RES8 WD
RES9 CD
RES9 WW
RES9 WD
Number of values
Log C CH4

857
858 Figure 6
859

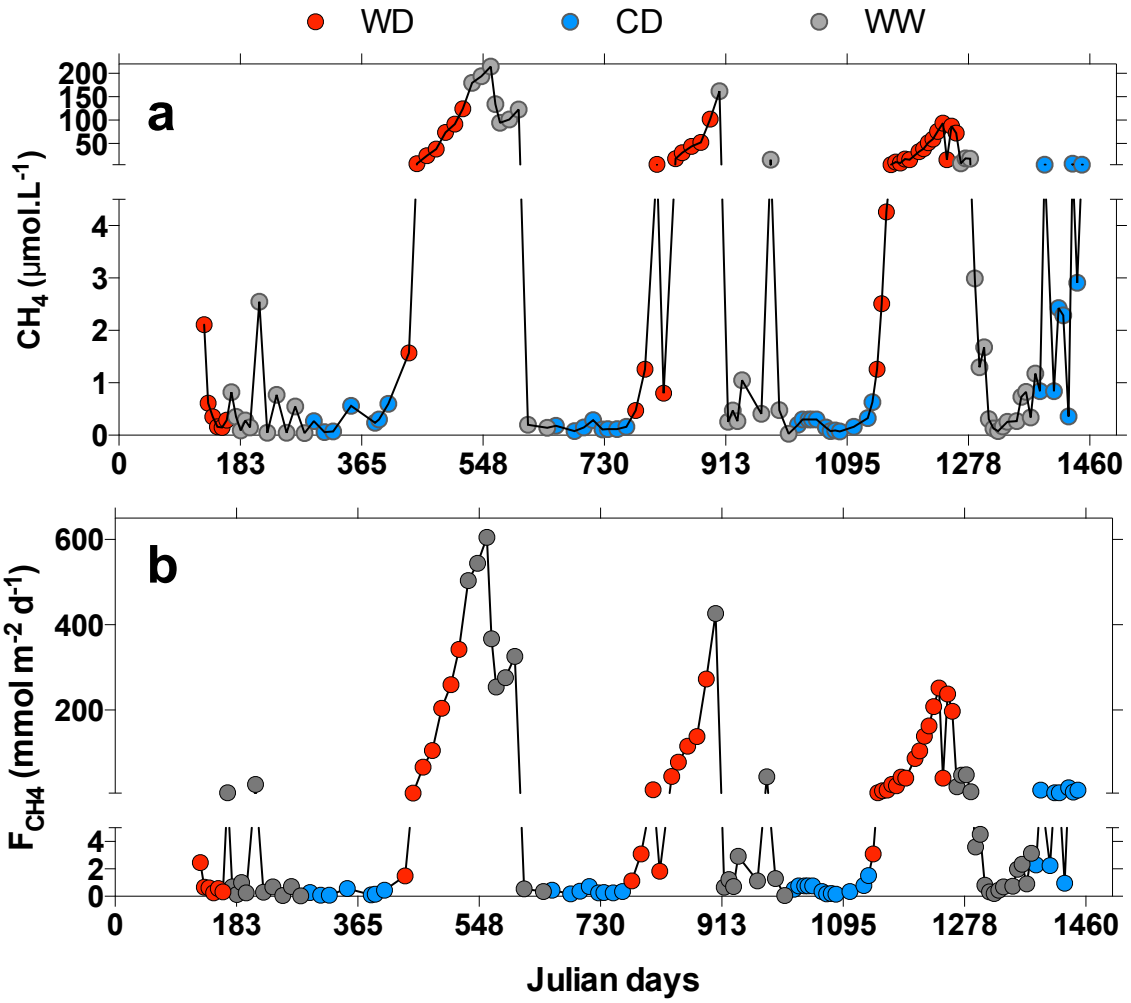


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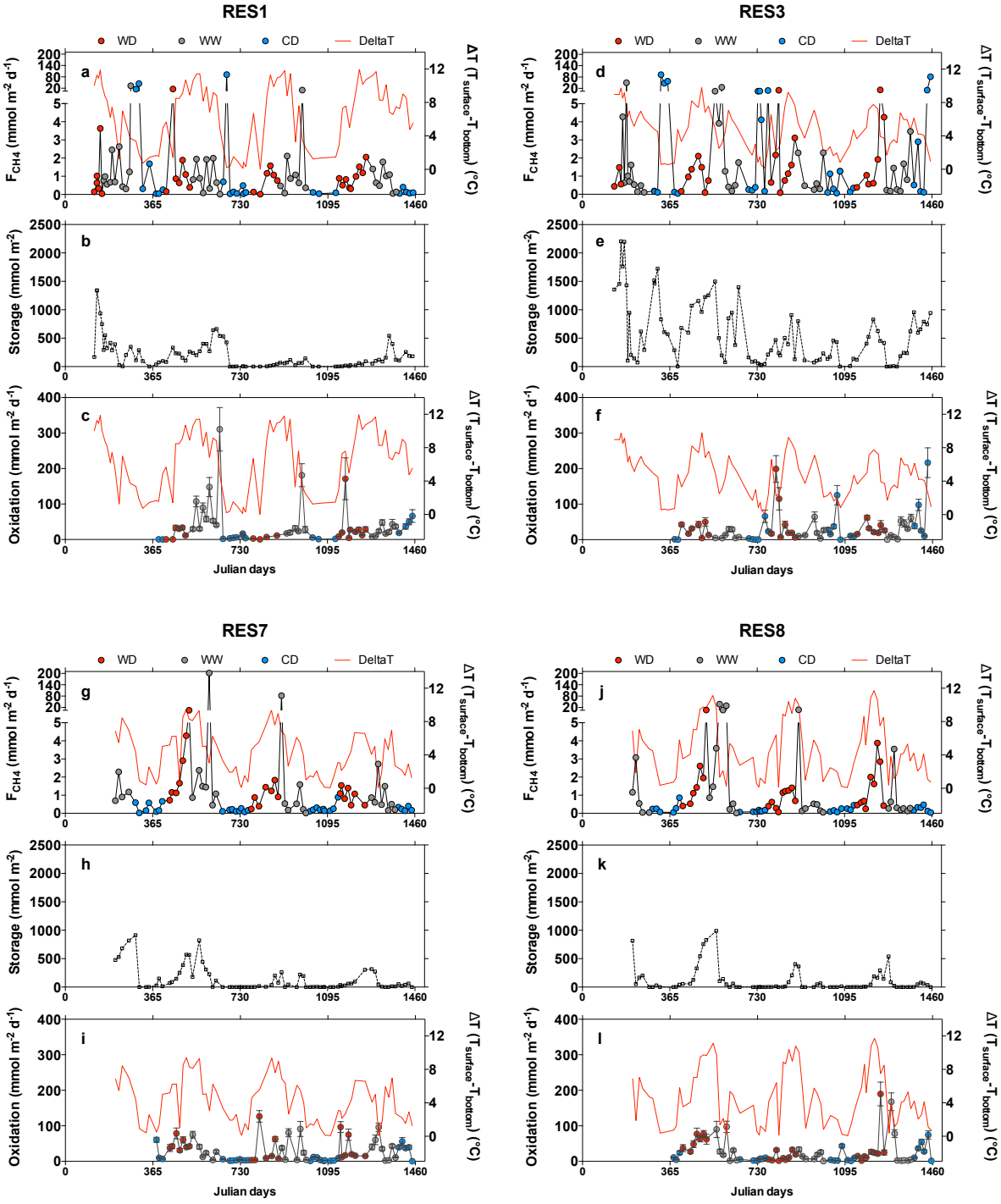
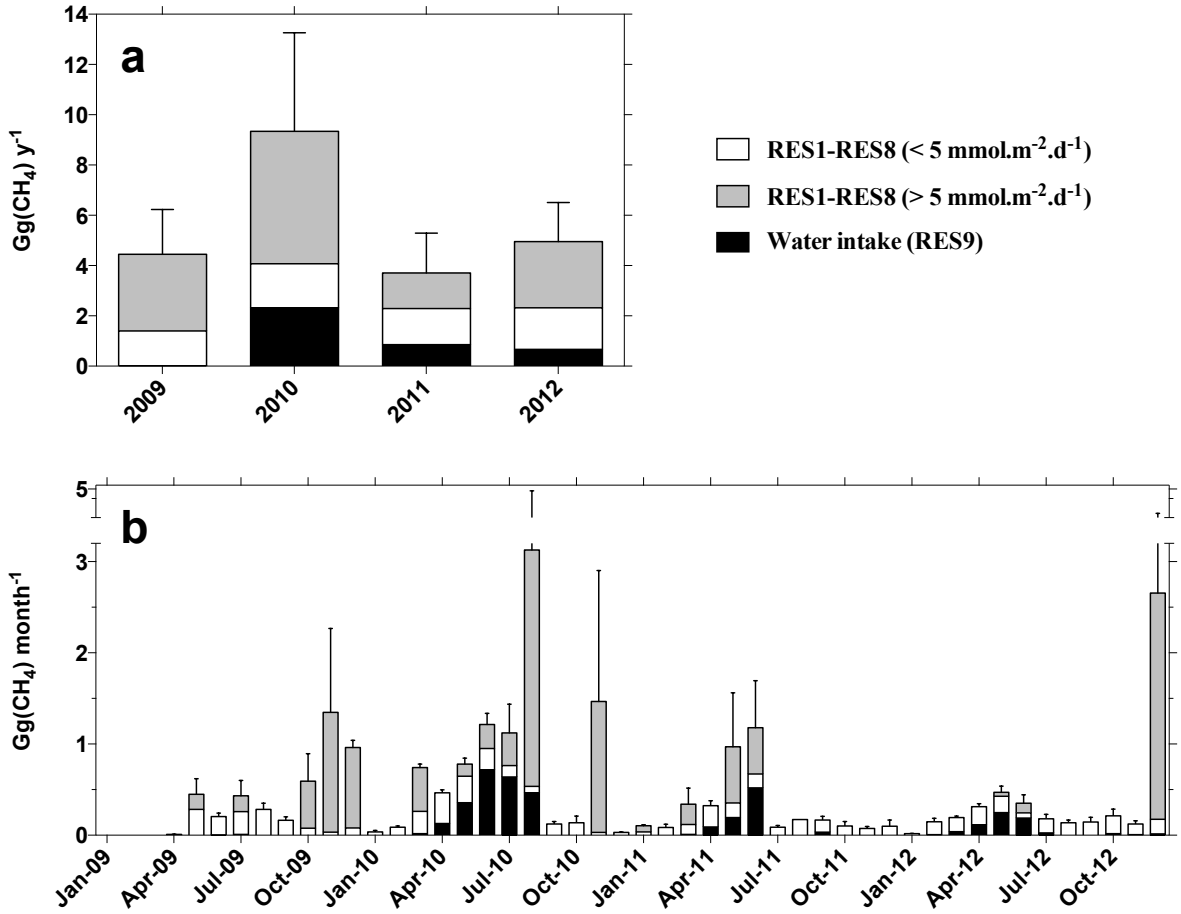
Figure 7


RES1
RES3
RES7
RES8
WD
WW
CD
DeltaT
a
b
c
d
e
f
g
h
i
j
k
l
F_{CH4} (mmol m^{-2} d^{-1})
Storage (mmol m^{-2})
Oxidation (mmol m^{-2} d^{-1})
ΔT ($T_{surface}$-T_{bottom}) (°C)
Julian days

869
870 Figure 8
871



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