

Spatial and temporal distribution of methane in an extensive shallow estuary, south India

A SHALINI¹, R RAMESH^{1,*}, R PURVAJA¹ and J BARNES²

¹*Institute for Ocean Management, Anna University, Chennai 600 025, India.*

²*School of Marine Sciences and Technology, University of Newcastle-upon-Tyne, Newcastle-upon-Tyne, NE1 7RU United Kingdom.*

**e-mail: rramesh_au@yahoo.com*

Sedimentary methane (CH₄) fluxes and oxidation rates were determined over the wet and dry seasons (four measurement campaigns) in Pulicat lake, an extensive shallow estuary in south India. Dissolved CH₄ concentrations were measured at 52 locations in December 2000. The annual mean net CH₄ flux from Pulicat lake sediments was 3.7×10^9 g yr⁻¹ based on static chamber measurements. A further 1.7×10^9 g yr⁻¹ was estimated to be oxidized at the sediment–water interface. The mean dissolved concentration of CH₄ was 242 nmol l⁻¹ (ranging between 94 and 501 nmol l⁻¹) and the spatial distribution could be explained by tidal dynamics and freshwater input. Sea–air exchange estimates using models, account only for ~13% (0.5×10^9 g yr⁻¹) of the total CH₄ produced in sediments, whereas ebullition appeared to be the major route for loss to the atmosphere (~63% of the net sediment flux). We estimated the total atmospheric source of CH₄ from Pulicat lake to be 0.5 to 4.0×10^9 g yr⁻¹.

1. Introduction

Methane (CH₄) is an important atmospheric trace gas with a tropospheric residence time of ~10 years. Mean tropospheric CH₄ increased from about 1650 ppbv in the early 1980s to around 1800 ppbv in 2005; tropospheric growth was variable throughout the 1990s and since then overall, has slowed significantly (<http://cdiac.ornl.gov/trends/otheratg/blake/blake.html>). CH₄ currently accounts for about 20% of anthropogenic greenhouse forcing (IPCC 2001).

CH₄ concentrations in temperate estuaries generally far exceed atmospheric equilibrium values and typically are highest at low salinities, decreasing seaward (de Angelis and Lilley 1987; de Angelis and Scranton 1993; Upstill-Goddard *et al* 2000; Middelburg and Nieuwenhuize 2001); hence estuaries are potentially large sources of atmospheric CH₄.

Estuarine CH₄ is derived both from external sources, i.e., river and open ocean, and via *in situ* production in anoxic micro-environments, such as in organic-rich sediments or O₂ deficient waters (de Angelis and Lee 1994; Chanton *et al* 1989; Iversen *et al* 1987) including turbidity maxima (Upstill-Goddard *et al* 2000). Estuarine CH₄ removal occurs through oxidation in surface sediment (van der Nat *et al* 1997) and in the water column (de Angelis and Scranton 1993), and emission to the atmosphere (Middelburg and Nieuwenhuize 2001).

Coastal environments including estuaries dominate marine CH₄ emissions (Bange *et al* 1994; Upstill-Goddard *et al* 2000) but these sources of atmospheric CH₄ remain poorly quantified (Middelburg *et al* 2002); estimates range from ~1 to 10% of the total global emissions. Although data from tropical/sub-tropical estuaries are scarce, a few that are available suggest that CH₄ emissions

Keywords. Methane emission; spatial and temporal variation; macro-algal zone; estuaries; Pulicat lake.

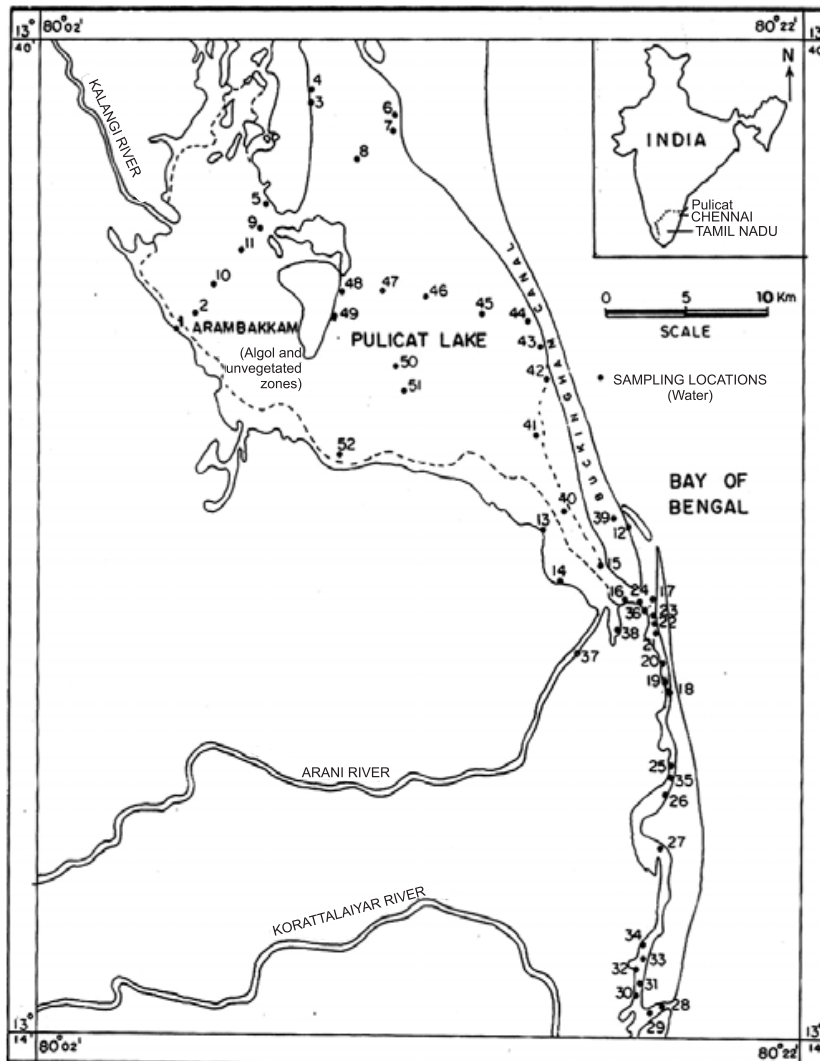


Figure 1. Study area: Arambakkam, Pulicat lake and surface water sampling locations (December 2000).

from polluted tropical estuaries are up to 20 times greater than from unpolluted ones and exceed those from temperate estuaries (Friborg *et al* 1997; Granberg *et al* 1997; Ramesh *et al* 1997; Purvaja and Ramesh 2000). A study of CH_4 emissions from coastal wet lands of south India showed that CH_4 emission rates from a polluted ecosystem (Adyar Estuary) were twice as high as in an unpolluted mangrove ecosystem (Purvaja and Ramesh 2000).

In this paper, we discuss seasonal (wet and dry season) CH_4 emission fluxes from Pulicat lake, an extensive shallow estuary in southeast India; estimated from static flux chamber data and through applying a gas exchange–wind speed relation to surface water dissolved CH_4 concentrations, and rates of potential sediment CH_4 oxidation determined from core incubations under aerobic and anaerobic incubations. Direct ebullition was estimated on a single occasion. Using these estimates we derive a mean annual CH_4 flux for the whole estuary. A generic CH_4 cycling budget for Pulicat

lake was constructed based on these data, which combine spatial and seasonal CH_4 distribution with assessments of fluxes at the sediment–water and air–sea interfaces. This provides the opportunity to discuss the relative contributions of different processes, such as the roles of diffusive and bubble-mediated fluxes.

2. Material and methods

2.1 Study area

Pulicat lake (latitude: 13°24'N; longitude: 80°02'E) is a shallow (~1–2.5 m) estuarine ecosystem that provides important support to the local economy via the exploitation of its fish stocks (figure 1). The mixing regime and biogeochemistry of Pulicat lake are highly seasonal. Freshwater input occurs from three small rivers, the Arani,

Kalangi and Swarnamukhi. Of these, the Kalangi is quantitatively the most important mainly due to its high discharge $\sim 700 \text{ m}^3 \text{ s}^{-1}$ during the monsoon season. Pulicat lake behaves as a classical estuary with open connection to the sea and salinity increasing seaward. Conversely, due to low freshwater inputs and high evaporation during the dry season (summer) Pulicat lake shrinks from its maximum surface area of $\sim 460 \text{ km}^2$ during the wet season to $\sim 250 \text{ km}^2$ during summer, with a drop in water level by ~ 1.5 metres. At this time the system behaves as a negative estuary; salinities within Pulicat lake exceeding those in surrounding coastal waters (Barnes 2003 unpublished). Occasionally (every two years or so) Pulicat lake is completely isolated from the Bay of Bengal during summer, effectively becoming a hyper-saline lagoon.

Pulicat lake sediments are all generally rich in organic carbon; mean value is $4.5 \pm 0.4\%$ (Shalini 2002). We estimated seasonal CH_4 fluxes at two contrasting sediment sites located about 300 m apart (Arambakkam; figure 1), selected to span the range of benthic environments found at Pulicat. One site was in an area of exposed silty sand with high populations of burrowing macro infauna including mole crabs, amphipods and polychaetes; we subsequently refer to this as the bioturbated zone. The other site was in an area of silty clay dominated by seagrasses (*Halophila ovalis*) and green algae (*Enteromorpha*, *Chaetomorpha*), and macrofauna; we subsequently refer to this as the macro-algal zone.

2.2 CH_4 analysis

Surface water samples (0.5 m) for dissolved CH_4 analyses were collected in 100 ml serum bottles at 52 locations during December 2000 (figure 1). Samples were immediately inoculated with 0.1 ml of saturated HgCl_2 to arrest microbial activity, sealed and stored in the dark in an incubated icebox prior to analysis. In the laboratory, 50 ml sub-samples were transferred to 100 ml sealed vials pre-flushed with ultra pure N_2 . The vials were then shaken vigorously for 10 minutes to equilibrate CH_4 between the two phases (Upstill-Goddard *et al* 1996), and analyzed by flame ionization gas chromatography (HP 6890); separation was done on 80–100 mesh Porapak Q, a 6 ft-long column. Column, injector and detector temperatures were 60, 100 and 250°C respectively, and N_2 (flow rate 30 ml min^{-1}) was used as a carrier gas (Ramesh *et al* 1997). Calibration standards were 1.77 ppmv CH_4 in N_2 (Matheson Inc., USA), and 116 ppmv CH_4 in N_2 (National Physical Laboratory, New Delhi, India). Details of calculations are described elsewhere (Upstill-Goddard *et al* 1996).

2.3 CH_4 flux measurements (Chamber method)

Sediment CH_4 effluxes from the macroalgal and bioturbated sediment sites were estimated with a static perspex chamber ($53 \times 37 \times 71 \text{ cm}$, $l \times w \times h$) in four seasons: pre-monsoon (September 1999), winter monsoon (December 1999), post-monsoon (March 2000) and inter-monsoon (June 2000). The chamber was secured to an aluminum base collar inserted to a sediment depth of $\sim 10 \text{ cm}$, three hours prior to sampling. A battery-operated air circulation pump with a flow rate of 1.5 l min^{-1} (M/S Aerovironment Inc., Monrovia CA, USA) was used to mix the chamber headspace and to transfer headspace samples to glass sampling bottles at 15-minute intervals over a 1-hour period. The sampling bottles were fitted with two stopcocks and were initially filled with N_2 -purged distilled water. Sample collection involved connecting the pump outlet to one stopcock and displacing the water with the sample air. CH_4 emission fluxes were determined from the increase in the headspace CH_4 mixing ratio with time, and changes in the chamber headspace volume, as determined by continuously monitoring changes in chamber water levels due to tidal height (Ramesh *et al* 1997; Purvaja and Ramesh 2000; Purvaja *et al* 2004). Surface water and sediment (10 cm depth) temperatures were also measured *in situ*.

2.4 Core fluxes

Perspex[®] cores ($15 \times 3 \text{ cm}$, length \times diameter) were used to collect sediments from the two different zones. Four replicate cores from each zone were collected to a depth of 10 cm leaving a 5 cm headspace to enable flushing with air and nitrogen.

The cores were transferred to the laboratory at *in situ* temperature in an insulated icebox containing ambient water collected on site. In the laboratory, replicate cores were incubated with either aerobic or anaerobic headspaces to determine potential oxidation rates of CH_4 . For aerobic incubations, the headspace was flushed with ambient air for 5 minutes and sealed. For anaerobic incubations the procedure was identical except that flushing was with N_2 for ~ 20 minutes. For each, gas samples (1 ml) were collected at 15-minute intervals for an hour and analysed for CH_4 as described above. Fluxes were estimated from temporal increases in headspace CH_4 mixing ratios, as for the incubation chamber. Since CH_4 oxidation is generally possible only under aerobic conditions, the anaerobic incubations represent gross CH_4 production in each core whereas the aerobic incubations estimate the fraction of CH_4 produced that escaped oxidation and subsequently released into the overlying water (Conrad and Rothfuss 1991).

The difference between the anaerobic and aerobic CH_4 flux thus represents the potential rate of CH_4 oxidation at the sediment–water interface.

2.5 CH_4 emission estimates (air–sea gas exchange models)

CH_4 emissions from the surface of Pulicat lake were also estimated from:

$$F = kL\Delta C$$

where F is the interfacial gas flux ($\text{mol unit area}^{-1} \text{unit time}^{-1}$), k is the gas transfer velocity (cm hr^{-1}), L is the gas solubility, and ΔC is the CH_4 partial pressure difference between the air ($1.7 \mu\text{atm}$) and water phases. Various empirical relationships have been derived for estimating k . The two most widely used are those of Liss and Merlivat (1986) and Wanninkhof (1992), which are often assumed to define the upper and lower limits for k . Clark *et al* (1995) derived a relationship for estuaries (using data from the Hudson River estuary), which takes into account turbulence due to interaction with the seabed without wind. For this reason, all three methods are included here to help a comparison with similar studies, although the model developed by Clark *et al* (1995) for estuaries is probably the most suitable to Pulicat lake. A mean wind speed (4 m s^{-1}) obtained from the Chennai harbour buoy data for December 2000 was used in the present calculations.

2.6 Collection of gas bubbles

Gas bubbles trapped in the sediments were collected during December 2000 using an inverted glass funnel (20 cm diameter) fitted with a silicone rubber septum in its stem between 10 and 20 m from the location where the static chamber was placed. The sediment was stirred up to collect the gas bubbles into the funnel. The gas was withdrawn using a disposable syringe and transferred into 10-ml Venoject[®] (Terumo, Belgium) tubes, which were pre-filled with a saturated NaCl solution. Excess salt solution was expelled during injection through an additional needle (Krüger *et al* 2001).

3. Results and discussion

3.1 Spatial variation of dissolved CH_4

Pulicat lake was a source of CH_4 to air on all sampling occasions. The mean dissolved CH_4 concentration for all samples were 242 nmol l^{-1} ($n = 52$) with distinct spatial variation, which ranged from

Table 1. Concentrations of dissolved CH_4 and associated environmental parameters in the surface water of Pulicat lake in December 2000.

Parameters	Min	Max	Mean
Dissolved CH_4 (nmol l^{-1})	94.14	500.62	241.91
SO_4 (mM)	2.45	16.42	9.30
Dissolved oxygen (mg l^{-1})	1.92	8.57	5.31
Standard deviation	45.94	265.83	117.02

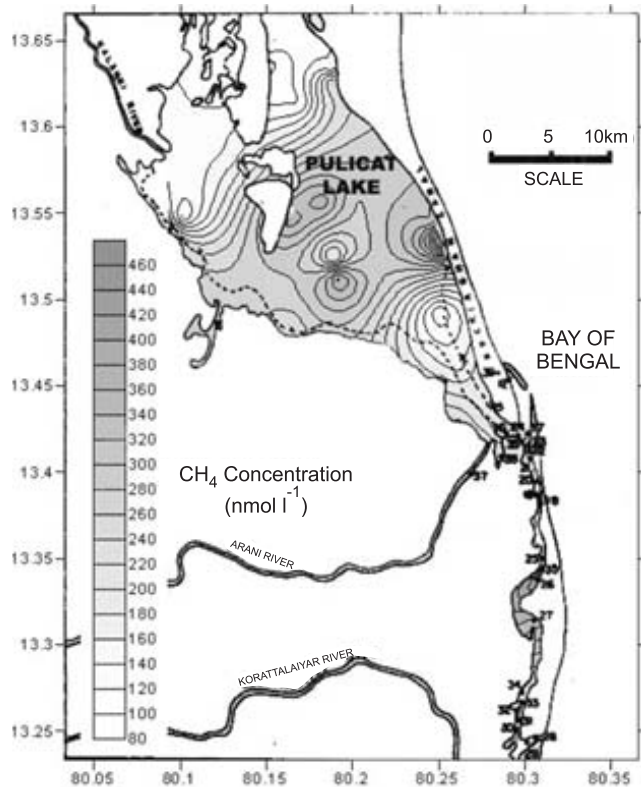


Figure 2. Spatial variation of dissolved CH_4 concentrations in the surface water (December 2000).

94 nmol l^{-1} to 500 nmol l^{-1} (table 1). Dissolved CH_4 concentration maxima therefore appear to be a feature of the centre of the estuary where mixing processes from seawater and freshwater inputs have least impact. The concentration variation was attributed to a variety of influences notably inputs from the Kalangi river which appeared to dilute dissolved estuarine CH_4 concentrations. Furthermore, tidal dilution occurs more readily at the Bay of Bengal area of the estuary resulting in lower observed dissolved CH_4 concentrations (figure 2).

In the present study, the correlation between salinity and dissolved CH_4 ($r^2 = -0.63$) was found to be weak and negative. Similar linear correlation of estuarine CH_4 concentrations with increasing salinity were observed by Purvaja and Ramesh (2000) for the Pichavaram mangrove and estuary in south India and in several temperate estuaries

Table 2. Seasonal and annual flux of CH_4 from the Pulicat Lake sediments calculated from temporal increase in headspace mixing ratios using a static flux chamber.

Seasons	Algal zone		Bioturbated zone	
	$\text{mg m}^{-2} \text{h}^{-1}$	$\mu\text{mol m}^{-2} \text{h}^{-1}$	$\text{mg m}^{-2} \text{h}^{-1}$	$\mu\text{mol m}^{-2} \text{h}^{-1}$
Wet (September 1999)	1.15	71.88	1.00	62.50
Wet (December 1999)	0.52	32.50	0.76	47.50
Dry (March 2000)	0.56	35.00	NA	NA
Dry (July 2000)	0.85	53.13	1.41	88.13
Mean	0.77	48.13	1.06	66.04
Annual flux from both the zones				
Min	2.10		3.06	
Max	4.63		5.68	
Mean	3.36		4.37	
Overall mean CH_4 flux from macro-algal and bioturbated zones	0.91	57.08		
Annual flux from Pulicat lake (10^9 g yr^{-1})	3.68			

(Upstill-Goddard *et al* 2000; Middelburg *et al* 2002).

Dissolved CH_4 showed a linear correlation with dissolved sulphate (-0.70) in the estuary. Sulphate inhibition of methanogenesis is widely known as a mechanism for reduced CH_4 production by marine sediments. Conversely dissolved O_2 showed a strong positive correlation with salinity (0.98), which appeared to be a direct function of tidal flushing. The Bay of Bengal exhibited dissolved O_2 (5.9 mg l^{-1}) concentrations higher than in the inner zone of the estuary ($\sim 2.7 \text{ mg l}^{-1}$). There was a negative correlation (-0.64) between dissolved O_2 and CH_4 (figure 5) similar to that observed between CH_4 and salinity. This may be a direct influence of tidal flushing, which dilutes the estuarine CH_4 and increases dissolved oxygen concentrations in Pulicat lake. An additional mechanism could be due to a reduction in CH_4 oxidation in the sediments, which could occur due to lower dissolved O_2 levels. Table 1 shows the concentrations of dissolved sulfate and oxygen that have a possible influence on surface dissolved CH_4 concentrations in Pulicat lake. Tidal flushing is a consistent phenomenon in the Pulicat lake, which plays a major role in the spatial variation of the dissolved CH_4 concentration. Although the dissolved CH_4 concentrations measured in the Bay of Bengal (50 nmol l^{-1}) during this study were lower than those measured in the estuary proper, they are considerably higher than data published for surrounding coastal waters/oceans (Upstill-Goddard *et al* 2000) suggesting that the system is also a source of CH_4 to the surrounding coastal waters.

3.2 Temporal variation of CH_4 emission from sediments (flux chamber)

Mean sediment–water–air CH_4 fluxes determined with the static chamber are shown in table 2.

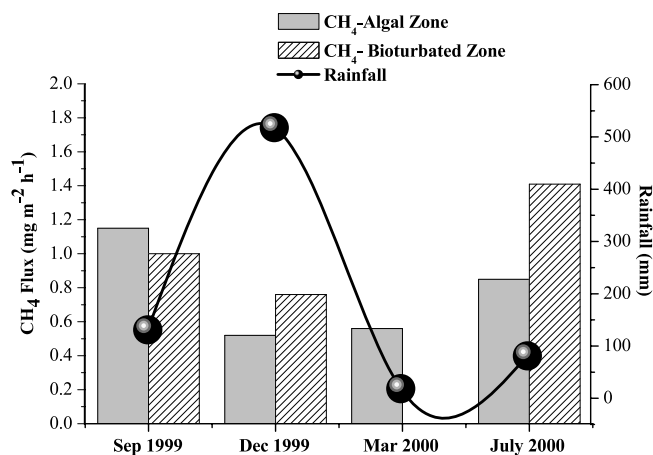


Figure 3. Seasonal variations in CH_4 emission and rainfall in the algal and unvegetated zones of Pulicat lake.

At the macro-algal site CH_4 emissions varied from $35 \mu\text{mol m}^{-2} \text{h}^{-1}$ during the dry season (March 2000) to a maximum of $72 \mu\text{mol m}^{-2} \text{h}^{-1}$ in the wet season (September 1999), whereas at bioturbated site CH_4 fluxes ranged from $48 \mu\text{mol m}^{-2} \text{h}^{-1}$ in the wet season (December 1999) to $88 \mu\text{mol m}^{-2} \text{h}^{-1}$ in the dry season (June 2000) (figure 3). This observed seasonal variation in CH_4 fluxes could be attributable to a variety of environmental influences notably an increase in ambient temperatures and decreased flushing of the estuary with freshwater. When all seasonal and spatial CH_4 flux data are pooled, the mean observed rate is $57 \mu\text{mol m}^{-2} \text{h}^{-1}$ ($0.91 \text{ mg m}^{-2} \text{h}^{-1}$). Comparison of the present mean CH_4 flux ($8.00 \text{ g m}^{-2} \text{yr}^{-1}$) with other aquatic ecosystems revealed that the flux rates were comparable to the rates observed in pichavaram mangroves ($10.00 \text{ g m}^{-2} \text{yr}^{-1}$); higher in highly polluted Adyar estuary and the Sunderban mangroves and lower in all the other estuarine areas. The gross sediment CH_4 production was

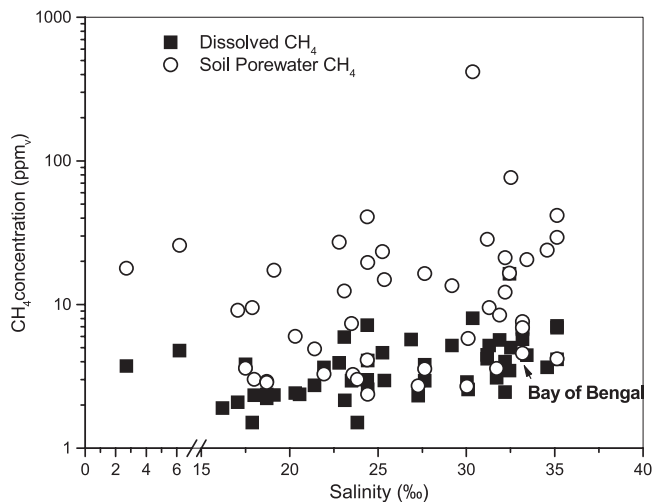


Figure 4. Salinity stratification of dissolved and soil pore water CH_4 in Pulicat lake.

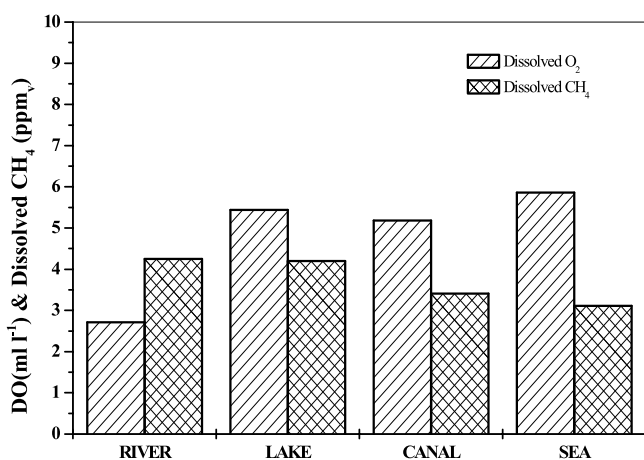


Figure 5. Spatial variation of dissolved oxygen with CH_4 in Pulicat lake.

calculated as $3.68 \times 10^9 \text{ g yr}^{-1}$ for the entire area of Pulicat lake (460 km^2).

3.3 CH_4 flux from the sediment cores (aerobic-anaerobic fluxes)

Aerobic CH_4 fluxes in cores from the macroalgal site were $0.26\text{--}21.1 \mu\text{mol m}^{-2} \text{ h}^{-1}$; these were generally lower than the anaerobically measured fluxes of $0.32\text{--}37.35 \mu\text{mol m}^{-2} \text{ h}^{-1}$; hence 19–67% (mean: 42%) CH_4 oxidation is indicated by these data. At the bioturbated site, potential CH_4 oxidation rates were similar to those at the macro-algal site with aerobic flux rates ($0.12\text{--}6.05 \mu\text{mol m}^{-2} \text{ h}^{-1}$) being lower than the anaerobic flux rates ($0.15\text{--}22.9 \mu\text{mol m}^{-2} \text{ h}^{-1}$). Thus potential oxidation rates ranged from 17 to 73% (mean: 45%). These data indicate that 41–45% of the potential diffusive CH_4 flux through the

Table 3. CH_4 fluxes calculated using air-sea gas exchange models based on mean surface water dissolved CH_4 concentration in Pulicat lake.

	Clark	Liss and Merlivat	Wanninkhoff
Flux ($\text{mol m}^{-2} \text{ d}^{-1}$)	0.00018	0.000054	0.00028
Flux for the Pulicat lake (10^9 g yr^{-1})	0.48	0.14	0.75

sediment-water interface gets oxidized in the oxic surface layers of the Pulicat lake.

3.4 Bubble ebullition

One of the main pathways of exchange of CH_4 between air and water is ebullition of CH_4 from the sediment through the water column (Sebacher *et al* 1985). The total concentration of CH_4 in bubbles was found to be 18% in the macroalgal ($546\text{--}14210 \mu\text{mol}$) and 41% in the bioturbated zones ($602\text{--}33390 \mu\text{mol}$). Accurate quantification of ebullition rates was only achieved successfully in December 2000, where 0.8 ml was collected in 3 hours from an area of 1 m^{-2} .

3.5 Calculation of emissions using air-sea gas exchange models

Table 3 shows CH_4 fluxes calculated using gas exchange models. A mean wind speed for December 2000 was used (4 m s^{-1} ; wind speed average obtained from the Chennai harbour buoy data) along with the mean observed dissolved CH_4 concentrations (242 nmol l^{-1} ; $n = 52$) and an area of 460 km^2 to estimate whole estuary fluxes.

The calculated sea-air fluxes of CH_4 varied between 0.1 and $0.8 \times 10^9 \text{ g yr}^{-1}$. The number calculated using the method of Clarke lies in the middle of these values ($0.48 \times 10^9 \text{ g yr}^{-1}$) is preferred for this study. This agrees within a range of contemporary data published by Purvaja and Ramesh (2000) for similar environments.

Diffusive CH_4 flux data available for the surrounding coastal Bay of Bengal are included here for comparison purposes. CH_4 concentration in the surface water is close to atmospheric equilibrium in equatorial areas but increases towards the Ganges/Bramaputra mouth where CH_4 flux to the atmosphere reaches up to $30 \times 10^9 \text{ g m}^{-2} \text{ yr}^{-1}$ (Berner *et al* 2003). Adopting the same units for Pulicat lake, based on the model of Clark gives a flux $\sim 1000 \times 10^9 \text{ g m}^{-2} \text{ yr}^{-1}$ demonstrating that although CH_4 atmospheric emission is small in global terms, on a regional basis Pulicat lake is a significant source of CH_4 .

Table 4. Proposed mechanistic CH_4 fluxes in Pulicat lake based on data from this study.

Process no.	Flux pathways	CH_4 flux	Remarks
1	Gross CH_4 production in the sediment ($10^9 \text{ g m}^{-2} \text{ yr}^{-1}$)	5.34	Processes 2+7 (sum of net CH_4 flux from sediment into water column + calculated potential CH_4 oxidation)
2	Net sediment–water–air flux (10^9 g yr^{-1})	3.68	Static chamber measurements
3	CH_4 ebullition as gas bubbles ($10^9 \text{ g m}^{-2} \text{ yr}^{-1}$)	1.83	Rate of gas bubble ebullition was calculated as 0.3 ml h^{-1}
4	Air–sea flux using Clarke’s Model (10^9 g yr^{-1})	0.48	Liss and Merlivat Model: $0.144 (10^9 \text{ g yr}^{-1})$ Wanninkoff Model: $0.755 (10^9 \text{ g yr}^{-1})$
5	Total CH_4 lost via diffusion and bubbles ($10^9 \text{ g m}^{-2} \text{ yr}^{-1}$)	2.31	Processes 3 + 4 (sum of flux from bubble ebullition and air–sea fluxes)
6	Calculated CH_4 oxidation in the water column ($10^9 \text{ g m}^{-2} \text{ yr}^{-1}$)	1.37	Processes 2 – 5 (difference between net CH_4 efflux from sediment into water and total CH_4 lost via diffusion and bubbles)
7	CH_4 oxidation (45%) from core fluxes ($10^9 \text{ g m}^{-2} \text{ yr}^{-1}$)	1.66	45% of Process 2
8	Loss from system to atmosphere (%)	75.90	Processes 9 + 11
9	CH_4 lost as bubbles (%)	62.80	(Process 5/Process 2) · 100
10	Oxidation in the water column (%)	24.10	100 – Process 8
11	Emission to atmosphere from sediment via water (%). Calculated using Clark’s air–sea gas exchange model (diffusive loss)	13.05	(Process 4/Process 2) · 100 (remaining CH_4 in the water column: includes bubbles and diffusion)

4. Conclusions

This study has revealed some interesting insights into mechanisms of CH_4 fluxes in Pulicat lake. Table 4 summarizes these findings, based on the flux calculations shown above. The entire estuarine sediment, regardless of habitat, appears to be a consistent source of methane throughout the year producing $\sim 3.68 \times 10^9 \text{ g yr}^{-1}$, after oxidation of CH_4 at the sediment/water interface which accounts for $\sim 45\%$ of that. However, only 15% of the net flux was calculated to be lost through sea–air gas exchange processes suggesting direct ebullition, water column oxidation and outflows to the Bay of Bengal account for the remainder. Bubble ebullition was observed throughout Pulicat on a seasonal basis and was noted to coincide with tidal minima. Collection of bubbles for concentration analysis was easily achieved (mean = 33% for macro-algal and 41% for bioturbated zone; $n = 10$), however the volume was successfully collected without disturbance only on one occasion ($0.3 \text{ ml m}^{-2} \text{ hr}^{-1}$, December 2000). If this number is tentatively adopted as representative of ebullition rates at Pulicat then, direct loss to the atmosphere through bubble formation would account for $\sim 63\%$ ($2.31 \times 10^9 \text{ g m}^{-2} \text{ yr}^{-1}$) of the total CH_4 produced in the sediments. Even with the inherent shortcomings of this approach it is clear that direct bubble ebullition is a major source of atmospheric CH_4 from Pulicat. Quantitative ebullition data would lead

to a considerable upscaling of the CH_4 estuarine source strength than that based on diffusive fluxes alone. Adopting the calculated flux rate using Clarke’s model ($0.48 \times 10^9 \text{ g yr}^{-1}$) as a minimum and the measured ebullition rate in addition to Clarke’s flux as a maximum we conservatively estimate the total annual atmospheric CH_4 source from Pulicat lake as $0.48\text{--}2.79 \times 10^9 \text{ g yr}^{-1}$.

Table 5 shows the results of this study in the context of other similar contemporary data from tropical coastal areas. Our data are in broad agreement with other static chamber measurements but are generally towards the higher-end of the observed range. The notable exceptions are the Adyar estuary in Chennai, India and the Vembanad lake on the west coast of India. The Adyar estuary is heavily impacted by inputs and flow within the river is mainly due to discharge from sewage treatment plants and discharge of sullage and other wastes from numerous storm water outlets (Ramesh *et al* 1997). Thus, it should be expected that these environments act as a strong source of atmospheric CH_4 . The Vembanad is an extensive shallow estuary geomorphologically similar to Pulicat (Verma *et al* 2001) and reported CH_4 fluxes are higher than those reported for all other estuaries. From the present study it is clear that extensive estuarine sub-tropical and tropical coastal lagoons represent a potentially significant local source of CH_4 to the atmosphere and further work is required to accurately quantify this flux.

Table 5. Methane emissions from tropical coastal wet lands and estuaries (updated from Purvaja et al 2004).

Site and reference	Method	Remarks	Latitude	CH ₄ flux (mg m ⁻² day ⁻¹)		CH ₄ flux (g m ⁻² yr ⁻¹)
				Min	Max	
Estuary: Pulicat lake, south India, this work	Chamber	Sediment-water fluxes	13°N	18.48	25.36	8.00
Estuary: Pulicat lake, south India, this work	Air-sea gas exchange model		13°N			0.75
Adyar Estuary, south India (Ramesh et al 1997)	Chamber	Sediment-water fluxes	13°N	64.08	1103.28	0.48 Clarke L&M Wannin 136.48
Mangrove (<i>Avicennia</i>), south India (Purvaja et al 2004)	Chamber	Flux through pneumatophores	11°N	7.40	63.70	10.00
Mangrove (<i>Bruguiera</i>), Hainan (Lu et al 1999)	Chamber	Sediment-water fluxes	18°N			0.39
Mangrove, Sunderban (Mukhopadhyay et al 2002)	Gradient	Micrometeorological measurement	22°N	-16.00	32.00	43.00
Mangrove (<i>Avicennia</i>), Puerto Rico (Sotomayor et al 1994)	Chamber	Gradient from pristine (low emission to highly polluted area)	18°N	4.00	42.00	n.a.
Ocean waters (Bay of Bengal) (Berner et al 2003)	Vacuum/headspace	Sediment-water fluxes; CH ₄ concentrations in water column; atmospheric exchange	22°N			0.0013 0.145
Ocean waters (Faber et al 1996)	Gradient	CH ₄ concentrations in water column and sediments; isotopic tracing of CH ₄	47°N			0.03

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