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## Greenhouse Gas Diffusive Flux Assessment

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#### Abstract

There is a growing interest and concern regarding Green House Gas (GHG) emissions as these is the major contributors of global warming. Carbon dioxide (CO<sub>2</sub>) and Methane (CH<sub>4</sub>) are two main GHGs which get emitted from both natural aquatic and terrestrial ecosystems as well as from anthropogenic activities. In natural aquatic system water storage is an important aspect for meeting the requirements of drinking water, food, and energy. However, development of such water bodies will impact the environment. Recent studies have shown that water bodies play a significant role as the sources of GHG emission, particularly in tropical climatic zones. One possible reason for this is the annual water temperature is much higher in tropical climates. This means that the rate of decomposition is faster leading to higher CO<sub>2</sub> and CH<sub>4</sub> flux in the water. Indian reservoirs indicate the complete spectrum of different types of reservoir found in the world. Their performance in terms of emission of GHGs is more difficult to trace out. In this paper pathways of GHG emission from a reservoir have been discussed and a tool as suggested by UNESCO/IHA has been used to assess the GHG emission from four existing reservoirs in India. These reservoirs are of different age and are located in different parts and climatic zones of India. Predicted diffusive fluxes in CO<sub>2</sub>eq have been estimated for the year 2013 as well as over the 100 years of their existence in terms of Tonnes CO<sub>2</sub> eq.

**Keywords:** Greenhouse gas; Global warming; Natural aquatic; Terrestrial ecosystems; Hydroelectricity

#### Introduction

The increasing anthropogenic activities have nowadays resulted in increasing concentration of natural gases  $\mathrm{CO}_2$  and  $\mathrm{CH}_4$  resulting in GHG effect [1]. According to the European Environment Agency (EEA),  $\mathrm{CO}_2$  emissions account for the largest share of GHGs equivalent of 80–85% of the emissions. Fossil fuel combustion for transportation and electricity generation are the main sources of  $\mathrm{CO}_2$  contributing to more than 50% of the emissions [2]. In India generation of electricity with coal based thermal power plant contributing to more than 55% [3], Hydroelectricity and natural gases represent respectively more than 15% and 5% of electric generation capacity. So far hydro power has been consider as the clean source of energy. Nevertheless, for the last few years GHG emission from freshwater reservoirs and their contribution has been a big issue regarding generation of electricity [4].

Recent studies showed that the carbon which is transferred to water body will undergo decomposition under oxic and anoxic conditions and produces CO<sub>2</sub> and CH<sub>4</sub> [5]. Once CO<sub>2</sub> and CH<sub>4</sub> are produced, they are not immediately released into the atmosphere, this gases are soluble in the water until a chemical event occurs that causes the gases to be released [6]. In this paper it briefly discusses exactly how reservoirs become a greenhouse gas and the mechanism behind the emission are been pointed out clearly and the predicted emissions of CO<sub>2</sub> and CH<sub>4</sub> in the form of diffusive flux from Indian reservoirs located in different climatic zones are been assessed using UNESCO/IHA GHG Risk Assessment Tool.

### GHGS emission by creating reservoir

While considering without a reservoir creation over a flowing water bodies only natural emission like conduction, deposition and emission will take place. On creation of a reservoir, emission from different parts of the reservoir will takes place Figure 1 shows detail sources of GHGs emission from the reservoir. The OM (Organic Matter) which present in the soil and plants is imported from the catchment in addition to that OM which preexisting in the reservoir together will decomposes aerobically and anaerobically and emits CO<sub>2</sub> and CH<sub>4</sub> gases to the atmosphere with the help of some parameters (primary and secondary) [2]. Macrophytes which are present on the surface of water are also responsible for some amount of CH<sub>4</sub> emission to the atmosphere.

## Reactions involved in emissions

The OM which is present in the water bodies and which has been inputted by surface and subsurface runoff decomposes under oxic condition and produce  $\mathrm{CO}_2$  (equation 1). And at the bottom the OM which is stored in the sediments decompose under anoxic conditions and produces  $\mathrm{CO}_2$  and  $\mathrm{CH}_4$  (equation 2).

Decomposition under oxic conditions:

$$C_6H_{12}O_6 + 6O_2 \rightarrow 6CO_2 + 6H_2O$$
 (1)

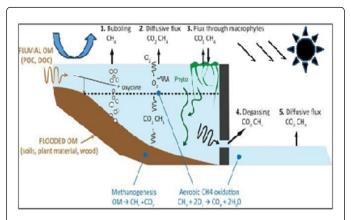
Decomposition in anoxic conditions (Methanogenesis):

$$C_6H_{12}O_6 \rightarrow 3 CO_2 + 3 CH_4$$
 (2)

CO<sub>2</sub> and CH<sub>4</sub> emissions to the atmosphere from reservoirs include:

- Bubble fluxes (ebullition) from the shallow part of water bodies
- Diffusive fluxes which are emitted from water surface of the reservoir

- Diffusion through macrophytes
- Degassing at downstream of reservoir outlet (s)
- Increased diffusive fluxes along the downstream part of the reservoir



**Figure 1:** Pathways of GHG emissions from a reservoir.

#### Main parameters/factors influencing GHG emissions

Parameters that effect in the production of  ${\rm CO_2}$  and  ${\rm CH_4}$  are divided into two types (Table 1)

- 1. Primary Parameters
- 2. Secondary Parameters

Primary parameters	Secondary parameters
Biomass of plants, algae, bacteria and animals in the water bodies	Wind speed and direction
Sediment load, Stratification of the water body OM storage, concentrations and C/N, C/P and N/P ratios in water and in sediments	Reductions in hydrostatic pressure as water are released through low level outlets
Nutrients supply; Temperature of water	Water current speeds
Light (absence of turbidity)	Rainfall
Dissolved oxygen concentrations	Water body depth and changes in water body depth

**Table 1:** Types of parameters that effect in the production of  $CO_2$  and  $CH_4$ .

## Calculation of Diffusive Flux from Aquatic Ecosystem

At Air water interface this both  $\rm CO_2$  and  $\rm CH_4$  will be transferred by diffusion from the aquatic ecosystems. This pathway happens at reservoir upstream and downstream and it is based on the Henry's law difference of partial pressure of a gas between the air (Pa) and the water (Pw). If Pw is higher than Pa the gas diffuses from the water to the atmosphere because a chemical compound always diffuses from the most concentrated layer to the less concentrated [5]. Several parameters control the intensity of the diffusive fluxes and the level of diffusive flux emissions can be estimated using the UNESCO/IHA Risk Assessment Tool with a confidence interval of 67% from the reservoir by giving the required inputs into the model.

# UNESCO/IHA GHG Risk Assessment Tool Model Formulas

Several alternative formulations were attempted by the UNESCO/IHA GHG emissions from freshwater reservoirs research project the following general expression has been given as the best fitting expressions (equation 3), (equation 4), and (equation 5) which consider the parameters which are responsible for the emission of  $\rm CO_2$  and  $\rm CH_4$  ( $\rm CCO_2$ ,  $\rm CCH_4$  in  $\rm mg/m2^*d1$ ) from the reservoir by considering the age of reservoir.

Flux 
$$C - CO_2 = 186.0 + 0.148 \times R + (944.485 + 1.91 \times T + 0.09727 \times T^2)$$
  
  $\times e^{-0.044} \times [52.339 - 0.7033 \times T - 0.0358 \times T^2]$  (3)

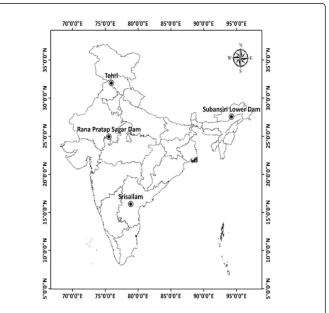
Formula for reservoir aged  $\leq 32$  years

$$C - CH_4 = 10 \left( 1.46 + 0.056 \times T - 0.00053 \times P - 0.0186 \times Age + 0.000288 \times Age^2 \right)$$

Formula for reservoir aged >32 years up to 100 years

$$C - CH_{A} = 10(1.16 + 0.056 \times T - 0.00053 \times P)$$
 (5)

R-Runoff (mm/year), Age-Age of the reservoir, T-Mean annual Temperature (°C), P-Mean annual Precipitation (mm/year).



**Figure 2:** location of reservoirs studied.

Reason for consideration of these parameters is:

- Max CO<sub>2</sub> emission occurs after flooding so positive factor of temperature.
- The new long term equilibrium emission (after the initial pulse) is a positive factor of runoff.
- Higher the runoff higher the CO<sub>2</sub> emission from the reservoir.
- The steepness of the initial decline (the exponential term) is a negative function of temperature.

 For older reservoirs (>32 years), diffusive CH<sub>4</sub> emissions are constant in time at a level which is determined by temperature and precipitation only.

## Range of variability of the estimates

The predicted values "lower limit" and the "upper limit" can be estimated as a function of the predicted values of gross GHG fluxes (of  $CH_4$  and  $CO_2$ ) and the mean square errors. Table 2 expresses how to estimate the values of the limits of the 67% confidence interval, for the models adopted in GHG Risk Assessment Tool.

Predicted Value	Lower limit	Upper limits			
Gross CCO <sub>2</sub> Flux	1/2.3* "Predicted Gross C-CO <sub>2</sub>	2.3* "Predicted Gross CCO <sub>2</sub> Flux"			
Gross CCH <sub>4</sub> Flux	1/3.55* "Predicted Gross CCH <sub>4</sub> Flux"	3.55* "Predicted Gross CCH <sub>4</sub> Flux"			

**Table 2:** Limits of predicted values of the 67% confidence interval.

#### Prediction of Diffusive Flux from Indian Reservoirs

Indian reservoirs indicate the complete spectrum of different types of reservoir found in the world. Some are located in a tropic climate which can release a significant amount of GHG and some in arid environments, where sequestration probably dominates over release of carbon [7]. Between these extremes are reservoirs located in wet, humid or dry tropical environments. Their performance in terms of emission of GHGs is more difficult to trace out. The data of the four Indian reservoirs which are located in different regions shown in the Figure 2 have been collected according to the latitude and longitude basics, the mean annual daily air temperature and Mean annual precipitation from 2 meters above the located surface has been analyzed by collecting the data from 19972013 from NASA Prediction of Worldwide Energy Resource (POWER). Runoff data are obtained from UNH/GRDC composite runoff fields V 1.0. and the analyzed values are shown in the Table 3 [8]. The predicted values of that particular year as well as the expected lower and upper range of CO2 and CH<sub>4</sub> with a confidence interval of 67 percent are listed in the Table 4 [9-11]. And mean emissions over reservoir life time (100 years) is shown in the Table 3.

S. No	Stations	Age	DMAP (Mm/Yr) (1997-2013)	R (mm/yr)	DMAT (0C) (1997-1912)	Lat.	Long.	
1	Srisailam	31	919	200	25	16005'13"N	78053'50"E	
2	Tehri	7	980	405	14.57	30022'40"N	78028'50"E	
3	RanaPratap Sagar	43	852	315	26	24055'04"N	75034'53"E	
4	Subansiri Lower	1	1766.5	500	9	27033'13"N	94015'31"E	
DMAP and DMAT Daily Mean Annual Precipitation and Temperature R – Runoff, Lat. Latitude, Long. – Longitude								

Table 3: Details of parameters which are required for estimating diffusive fluxes by using GHG risk assessment tool.

S. No Stations	Stations	Predicted gross* annual CO <sub>2</sub> diffusive flux (mg CCO <sup>2</sup> m <sup>2</sup> d1) 67% CI			(mg C	oss* annual ( sive flux CH <sub>4</sub> m <sup>2</sup> d1) 57% Cl	Remarks		
	Predicted value	Lower limit	Upper limit	Predicted value	Lower limit	Upper limit	CO <sub>2</sub> emission	CH <sub>4</sub> emission	
1	Srisailam	410	178	943	118	33	420	М	Н
2	Tehri	812	353	1868	114	32	404	Н	Н
3	Rana Pratap Sagar	397	173	913	146	41	518	М	Н
4	Subansari	1223	532	2814	55	15	194	Н	Н
H = High,	M = Medium, CI = Cor	nfidence Interval	1	1	1		1	1	1

Table 4: Average Diffusive Flux over 100 years with 67% confidence interval.

## Conclusions

In this case study, four reservoirs from different regions of India have been selected and the emission through diffusive flux has been

estimated. According to the study,  $CH_4$  emissions are high for all the reservoirs and  $CO_2$  emissions are high for Tehri, Subansari and moderate for Srisailam and Rana Prathap Sagar when compared with threshold limits of the model. While considering throughout the life

time assessment of the reservoir (100 years), the emission of  $CH_4$  is high for all reservoirs except Subansari and  $CO_2$  emissions are in a limit and medium except Srisailam (Table 5). Even though these are predicted values, the  $CH_4$  emission is high for all the reservoirs and hence mitigation measures must be taken to reduce the emission since GWP of  $CH_4$  is 25 times higher than the  $CO_2$ . Water bodies have the potential to emit large amounts of  $CO_2$  and  $CH_4$  and contribute to global warming [12]. The decomposition of organic matter is the main reason for the production of these GHGs so we have to control the

entrance of OM into water bodies, maybe up to some extent. Another possibility is logging trees before starting the flooding process so that less organic matter is available for decomposition. Due to the fact that the oxidation of CH<sub>4</sub> through Methanotrophic bacteria seems to be a key factor to decrease the amount of CH<sub>4</sub> released into the atmosphere, this mechanism should be supported somehow to minimize the emissions from water bodies. There is still a need for lot of research to understand all the important processes.

State	Dam	Area (Km <sup>2</sup> )	IC (MW)	Predicted value (mg C m² d¹)		at of CCO <sub>2</sub> /yr btC	btCO <sub>2</sub> eq/yr ct of CCH <sub>4</sub> /yr	dtCH <sub>4</sub> /yr	et of CO <sub>2</sub> eq/yr	Total	
				CO <sub>2</sub>	CH <sub>4</sub>						
AP	Srisailam	800	1670	413	130	120596	442185	37960	50613	1E+06	2E+06
Uttarakhand	Tehri	52	100	372	82	7060.6	25889	1556.4	2075.1	51879	77767
Rajasthan	Rana P Sagar	198.2	172	479	160	34652	127058	11575	15433	385829	512888
Assam	Subansiri	33.5	2000	400	31	2556.5	9373.7	198.1	264.2	6604.2	15978

alncludes conversion of predicted value of CO2 into Tons over the complete surface area per year (Surface area x predicted value of CO2 x 0.001 x 365)

Table 5: Emissions calculated from total surface area and converting into T of CO<sub>2</sub> eq averaged over 100 years.

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<sup>&</sup>lt;sup>b</sup>Converting CCO<sub>2</sub> into CO<sub>2</sub>eq by multiplying with GWP of CO<sub>2</sub>(t CCO<sub>2</sub> x 3.6 x 1)

cludes conversion of predicted value of CH4 into Tons over the complete surface area per year (Area x predicted value OF CH4 x0.001 x 365)

dIncludes conversion of CCH<sub>4</sub> into CH<sub>4</sub> (t CCH4 x 1.3)

<sup>&</sup>lt;sup>e</sup>Converting CCO<sub>2</sub> into CO<sub>2</sub> eq by multiplying with GWP of CO<sub>2</sub> (tCH<sub>4</sub> x 25)