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## Optimization of elevated CO<sub>2</sub> levels and nutrient management for lowland rice ecosystem

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### Abstract

Experiments were conducted to assess the optimum CO<sub>2</sub> levels and nutrient management strategy for enhancing the rice yield and soil fertility status with rice (var.CO. 50) as test crop in open top chambers (OTC). The treatments comprised of three levels of CO<sub>2</sub> enrichment viz., Atmospheric CO<sub>2</sub> (C<sub>0</sub>): 370 ppm (Control); OTC I (C<sub>1</sub>): 550 ppm and OTC II (C<sub>2</sub>): 750 ppm. Both the OTC chambers were connected to a control monitoring system and the each chamber has a regulator to adjust CO<sub>2</sub> flow. Rice crop was fertilized with four varying nutrient management techniques viz., inorganic (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup>), organic (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis), IPNS (Integrated Plant Nutrient System) (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>) and untreated control. Maximum grain and straw yields were recorded due to the application of Integrated Plant Nutrient System (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>) in the CO<sub>2</sub> concentration of 750 ppm. Highest acid phosphatase activity was registered at the CO<sub>2</sub> concentration of 750 ppm followed by 550 ppm and 370 ppm. Acid phosphatase activity was the highest due to the application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis) followed by the application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>. With increase in CO<sub>2</sub> concentration an increase in CO<sub>2</sub> emission and microbial biomass were observed.

**Keywords:** Elevated CO<sub>2</sub> level, nutrient management, soil enzyme, microbial biomass, CO<sub>2</sub> emission

### Introduction

Rice (*Oryza sativa* L.) is one of the most important crops in the world and the primary staple food in Asia. Cultivation practices such as crop rotations, soil tillage, fallow periods, and water management have been known to affect crop yields and soil carbon sequestration. Among the GHGs CO<sub>2</sub> is the main one, the effect of which should be related to the crop production. Carbon serves as the primary substrate for photosynthesis and is the one to contribute to the yield formation in plants. Rice responses to climate change vary with region and rice cultivars. There is a need for more experiments on various cultivars of rice under global warming situation in many areas. Hence this study was conducted with the objectives to determine the effect of elevated CO<sub>2</sub> levels on growth, yield and soil fertility status under lowland rice ecosystem. Depending on population growth and energy use scenarios, atmospheric CO<sub>2</sub> concentration is expected to rise from about 370 μmol<sup>-1</sup> currently to about 485 to 1000 μmol<sup>-1</sup> by 2100 (Prentice *et al.*, 2001)<sup>[9]</sup>. Elevated CO<sub>2</sub> typically increases plant photosynthesis and biomass production, whereas increasing temperatures might either decrease or increase photosynthesis and production (Baker *et al.*, 1993)<sup>[2]</sup>.

### Materials and Methods

Pot experiments were conducted with rice (var.CO. 50) as test crop in open top chambers (OTC). The treatments comprised of three levels of CO<sub>2</sub> enrichment viz., Atmospheric CO<sub>2</sub> (C<sub>0</sub>): 370 ppm (Control); OTC I (C<sub>1</sub>): 550 ppm and OTC II (C<sub>2</sub>): 750 ppm. Both the OTC chambers were connected to a control monitoring system and the each chamber has a regulator to adjust CO<sub>2</sub> flow. Rice crop was fertilized with four varying nutrient management techniques viz., inorganic (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup>), organic (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis), IPNS (Integrated Plant Nutrient System) (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>) and untreated control.

Mean monthly climatic variables at different stages of crop growth viz., active tillering, panicle initiation, flowering and harvest stages were measured. Grain and straw yields were recorded. Field moist soil samples were analyzed for acid and alkaline phosphomonoesterase activity with a buffered disodium p-nitrophenyl phosphate tetrahydrate solution (Tabatabai, 1982) [12]. Soil dehydrogenase and urease activities were measured (Von Mersi and Schinner, 1991 [14]; Tabatabai and Bremner, 1972) [13]. Soil CO<sub>2</sub> emissions (Carter and Gregorich, 2008) [5] by alkali trap method, soil organic carbon (Walkely and Black, 1934) [15], microbial biomass C in soil by chloroform fumigation method (Beck *et al.* 1997) [3] and water soluble carbon were analysed.

## Results and Discussion

The mean monthly climatic variables recorded at different stages of crop growth indicated that air temperature was the highest at harvest stage and humidity per cent was the highest at active tillering stage (Table 1).

Maximum grain and straw yields were recorded due to the application of Integrated Plant Nutrient System (150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>) in the CO<sub>2</sub> concentration of 750 ppm. Application of inorganics alone recorded higher yield than the application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis). The untreated control recorded the lowest yield of 33.6 g pot<sup>-1</sup> (Table 2). Baker *et al.* (1992) [1] showed that shoot and root biomass increased under increasing CO<sub>2</sub> concentration. According to Yang *et al.* (2006) [16], the final total biomass of rice was increased by 16% under elevated CO<sub>2</sub>.

Highest acid phosphatase activity was registered at the CO<sub>2</sub> concentration of 750 ppm followed by 550 ppm and 370 ppm. Acid phosphatase activity was the highest due to the application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis) followed by the application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>. Application of inorganics alone recorded higher acid phosphatase activity than the untreated control. A similar pattern of results was obtained with alkaline phosphatase, dehydrogenase and urease activity also (Table 3). Larson *et al.* (2002) [7] and Lipson *et al.* (2005) [8] found a significant stimulation of soil extracellular enzyme activities under elevated CO<sub>2</sub>.

With increase in CO<sub>2</sub> concentration an increase in CO<sub>2</sub> emission was observed. At flowering stage the emission was more followed by tillering and harvest stages. Application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis) recorded higher CO<sub>2</sub> emission followed by the application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>. The application of organic substances or manure at high rates in the fields caused the deposition of soil and organic substances. The organic substances reacted with soil particles to form complex compounds that were hardly decomposed into carbon dioxide (Suwannarit, 2008) [11]. The optimum application rates of manure are between 3.13-6.25 t ha<sup>-1</sup> because those application rates have the least effect on quantities of greenhouse gas emissions (Sampanpanish, 2012) [10]. Application of inorganics alone and untreated control recorded lowest CO<sub>2</sub> emission (Table 4).

With increase in CO<sub>2</sub> concentration, an increase in organic C content was observed. The labile soil C pools, namely microbial biomass C, readily mineralizable C, and potassium permanganate oxidizable C were increased by 27, 38 and 37% respectively under elevated CO<sub>2</sub> concentration of 550 μmol mol<sup>-1</sup> than ambient CO<sub>2</sub>, 394 μmol mol<sup>-1</sup>. Application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis) recorded higher organic carbon content followed by the application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>. Application of inorganics alone and untreated control recorded lowest organic carbon content. Water soluble carbon content was not remarkably changed due to the treatment effects (Table 5).

With increase in CO<sub>2</sub> concentration, an increase in microbial biomass C was observed. Soil microbial biomass C under elevated CO<sub>2</sub> was significantly larger (by 18.9–25.2%) than that under ambient CO<sub>2</sub> (Inubushi *et al.* 2010) [6]. Application of organics (1/3<sup>rd</sup> FYM+ 1/3<sup>rd</sup> Vermicompost + 1/3<sup>rd</sup> Neem cake on N equivalent basis) recorded higher microbial biomass C content followed by the application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup>. Application of inorganics alone and untreated control recorded lowest microbial biomass C content (Table 5).

With CO<sub>2</sub> enrichment of 750 ppm and application of 150:50:50 kg N, P, K ha<sup>-1</sup> +25 kg ZnSO<sub>4</sub> ha<sup>-1</sup> +FYM @ 12.5 t ha<sup>-1</sup> + Azospirillum @ 2 kg ha<sup>-1</sup> enhanced the growth, yield and soil fertility status under lowland rice ecosystem.

**Table 1:** Mean monthly climatic variable during crop growing stages

Variables	Stage of the crop			
	Active tillering	Panicle Initiation	Flowering	Harvest
Air temperature ° C	35.1	33.0	35.5	40.2
Humidity %	49.9	48.6	35.3	34.2
Soil temperature ° C	36.8	33.8	28.8	35.7
Canopy temperature ° C	32.2	29.6	28.5	32.2

**Table 2:** Effect of elevated CO<sub>2</sub> and nutrient management on grain and straw yield of rice (g pot<sup>-1</sup>)

Treatments	Grain			Straw		
C <sub>0</sub> : 370 ppm S <sub>1</sub>	33.6			48.6		
S <sub>2</sub>	54.3			68.6		
S <sub>3</sub>	42.0			63.6		
S <sub>4</sub>	81.0			98.6		
Mean	52.7			69.9		
C <sub>1</sub> : 550 ppm S <sub>1</sub>	46.6			65.3		
S <sub>2</sub>	73.3			110.6		
S <sub>3</sub>	67.0			71.6		
S <sub>4</sub>	96.6			128.0		
Mean	70.9			93.9		
C <sub>2</sub> : 750 ppm S <sub>1</sub>	54.0			79.3		
S <sub>2</sub>	98.3			144.3		
S <sub>3</sub>	69.0			98.6		
S <sub>4</sub>	142.6			167.6		
Mean	91.0			122.5		
CD(P:0.05)	C	N	CXN	C	N	CXN
	23.4**	27.0**	NS	17.5**	20.2**	NS

**Table 3:** Effect of elevated CO<sub>2</sub> and nutrient management on Soil enzyme activity: Acid and alkaline phosphatase activity ( $\mu\text{g PNP g}^{-1}$ ); dehydrogenase activity (DHA) ( $\mu\text{g TPF g}^{-1} \text{ day}^{-1}$ ) and urease activity ( $\mu\text{g NH}_3 \text{ g}^{-1}$ ) (Harvest stage)

Treatments	Acid phosphatase			Alkaline phosphatase			DHA			Urease		
C <sub>0</sub> : 370 ppm S <sub>1</sub>	60.75			132.45			27.20			10.25		
S <sub>2</sub>	60.66			142.13			30.75			12.12		
S <sub>3</sub>	74.50			154.50			36.83			13.85		
S <sub>4</sub>	63.17			162.29			37.51			14.03		
Mean	64.62			147.84			33.07			12.56		
C <sub>1</sub> : 550 ppm S <sub>1</sub>	67.16			151.27			25.09			11.57		
S <sub>2</sub>	75.32			158.61			28.66			12.78		
S <sub>3</sub>	86.62			161.29			28.81			17.03		
S <sub>4</sub>	76.13			162.76			30.11			21.85		
Mean	76.31			158.48			28.00			15.81		
C <sub>2</sub> : 750 ppm S <sub>1</sub>	72.15			157.97			28.76			12.59		
S <sub>2</sub>	77.80			162.10			35.53			14.29		
S <sub>3</sub>	91.66			166.25			36.92			22.54		
S <sub>4</sub>	80.20			175.00			39.13			27.10		
Mean	80.45			165.33			35.09			19.13		
CD(P:0.05)	C	N	CXN	C	N	CXN	C	N	CXN	C	N	CXN
	4.0	4.7	NS	8.1	9.3	NS	2.9	3.5	NS	3.0	3.5	6.0

**Table 4:** Effect of elevated CO<sub>2</sub> and nutrient management on CO<sub>2</sub> emission ( $\text{mg CO}_2 \text{ m}^{-2} \text{ day}^{-1}$ )

Treatments	Tillering			Flowering			Harvest		
C <sub>0</sub> : 370 ppm S <sub>1</sub>	16.0			39.0			11.0		
S <sub>2</sub>	14.8			33.2			8.7		
S <sub>3</sub>	17.6			41.5			15.6		
S <sub>4</sub>	14.8			35.0			10.2		
Mean	15.8			37.2			11.4		
C <sub>1</sub> : 550 ppm S <sub>1</sub>	31.2			55.4			26.5		
S <sub>2</sub>	19.2			36.2			12.1		
S <sub>3</sub>	53.2			76.1			44.3		
S <sub>4</sub>	26.4			48.5			20.2		
Mean	32.5			54.0			25.8		
C <sub>2</sub> : 750 ppm S <sub>1</sub>	58.2			78.7			50.7		
S <sub>2</sub>	29.2			49.2			23.2		
S <sub>3</sub>	87.0			98.0			66.7		
S <sub>4</sub>	36.8			53.4			27.6		
Mean	52.8			69.8			42.0		
CD(P:0.05)	C	N	CXN	C	N	CXN	C	N	CXN
	1.5**	1.7**	3.0	2.1**	2.4**	4.2	1.4**	1.7**	2.9

**Table 5:** Effect of elevated CO<sub>2</sub> and nutrient management on organic C, water soluble C ( $\text{g kg}^{-1}$ ) and microbial biomass C ( $\mu\text{g kg}^{-1}$ )

Treatments	Organic C			Microbial biomass C			Water soluble C		
C <sub>0</sub> : 370 ppm S <sub>1</sub>	5.1			275			12.3		
S <sub>2</sub>	5.5			437			12.4		
S <sub>3</sub>	6.5			520			12.6		
S <sub>4</sub>	6.2			464			12.5		
Mean	5.8			424			12.4		
C <sub>1</sub> : 550 ppm S <sub>1</sub>	6.2			390			12.3		
S <sub>2</sub>	6.6			545			12.4		
S <sub>3</sub>	7.3			688			12.5		
S <sub>4</sub>	6.8			626			12.5		
Mean	6.7			562			12.4		
C <sub>2</sub> : 750 ppm S <sub>1</sub>	6.6			437			12.3		
S <sub>2</sub>	7.2			680			12.3		
S <sub>3</sub>	7.5			931			12.5		
S <sub>4</sub>	7.4			807			12.4		
Mean	7.2			714			12.4		
CD (P:0.05)	C	N	CXN	C	N	CXN	C	N	CXN
	0.16**	0.18**	0.31	11.61**	13.40**	23.22	NS	NS	NS

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