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# EFFECT OF SULPHUR DIOXIDE ON PLANT CHLOROPHYLL ON THE FAMILY OF BRASSICACEAE Manoj Kumar<sup>1</sup>, Anjali<sup>2</sup>, Narayan<sup>3</sup>, Sheetal Chaudhary<sup>4</sup> and Krishan Pal<sup>\*5</sup>



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# Abstract

The effect of SO<sub>2</sub> on chlorophyll in (*Brassica juncea* [*L.*] *Czern.and Coss.cv.Pusa Bold; Raphanus sativus I Mino Early Long White*).In exposed seedlings SO<sub>2</sub> concentration ranging (653, 1306, 2612&3918  $\mu$ g m<sup>-3</sup> S resulted in a sharp decrease in total chlorophyll content. Both chlorophyll a and chlorophyll b got reduced declined following the exposure of SO<sub>2</sub>. This decrease could be due to disturbance in chlorophyll b got reduced because of conversion of chlorophyll into phaeophytin upon exposure to SO<sub>2</sub>. After spraying of calcium hydro the yield in C+ SO<sub>2</sub> exposed set was higher than the set of plant exposed to SO<sub>2</sub> alone. Maximum reductions v noticed at 3918  $\mu$ g m<sup>-3</sup> of SO<sub>2</sub>. The breakdown of chlorophyll molecules by SO<sub>2</sub> as measured by loss of Mg<sup>++</sup> total chlorophyll.

Keywords: - Sulphur Dioxide, Chlorophyll, Brassicaceae

# Introduction

Sulphur dioxide is one of the major air pollutants in industrialized areas that can damage vegetation. The process of photosynthesis appears to be mainly affected. A current view is that plant must exhibit visible symptoms for injury to exist. Many studies have tried to establish certain relationship between visible symptoms caused by exposure to  $SO_2$  and injury. However several glasshouse exposure studies (Bogorad L 1966; Reinert RA and JS Sanders, 1982) have shown a net reduction of growth and yield without the development of visual symptoms. The present study was initiated to determine if low concentration of SO<sub>2</sub> have any effect on the process photosynthesis without producing visual of symptoms. The effect of SO<sub>2</sub> was studied on the following aspects of pigment metabolism : (a) Effect on chlorophyll conversion into phaeophytin,(b)Effect on the activity of chlorophyllase and (c) Effect of low cytoplasmic pH caused by  $SO_2$  on (a) and (b).

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# Growth Conditions

Plants from the family Brassicaceae (Brassica juncea (L.) Czern.and Coss.cv.Pusa Bold; Raphanus sativus L.cv.Mino Early Long White) seeds were washed with sterile distilled water and then treated with 0.1% mercuric chloride for 5 minutes and finally washed with sterile distilled water for 15 minutes. Surface sterilized seeds were allowed to imbibe water for 6 hrs.and thereafter sown on petriplates lined with cotton over which Whatman no.40 filter paper was placed. Seeds were placed on filter paper. For each variety, five sets each having 150 seeds were maintained. Then seeds were sown in polythene bags/earthen ware containing garden soil. The soil was sandy loam and homogenously mixed with farmyard manure. All experiment will carried out in a fumigation chamber (1m\*1m\*1m) made of iron rods. Each chamber was portable and covered with transparent polythene sheets. Then Plants were exposed to  $SO_2$  from 5<sup>th</sup> day onwards. Plants of each species were divided into five sets. Out of five four sets were exposed to four different concentration of Sulphur dioxide (653, 1306, 2612, 3918  $\mu$ g m<sup>-3</sup>) while the fifth one was used as control.

# Sulphur dioxide Treatment

Sulphur dioxide was prepared by allowing a reaction of dilute sulphuric acid (10%  $H_2SO_4$ ) and sodium

sulphite  $(NA_2SO_3)$  under controlled condition of temperature and humidity.Complete reaction of 1M  $NA_2SO_3$  with 10%  $H_2SO_4$  produces 1M  $SO_2$  or 126 mg of  $NA_2SO_3$  yields 64 mg  $SO_2$ .

The chemical reaction for  $SO_2$  preparation is as follows-

Hence, 1.968 of NA<sub>2</sub>SO<sub>3</sub> is required to produce 1 mg (1000  $\mu$ g m<sup>-3</sup>) of sulphur dioxide. Therefore, on the basis of this equation 1.285, 2.571, 5.142, 7.713 mg of NA<sub>2</sub>SO<sub>3</sub> were used to obtain 653, 1306, 2612, 3918  $\mu$ g m<sup>-3</sup> of sulphur dioxide, respectively, inside the exposure system. The plants were given the treatment of SO<sub>2</sub> on alternate day for 2 hrs.

#### Calcium hydroxide treatment

Calcium hydroxide (0.5% aqueous solution) was used for amelioration studies. Results of germination studies showed that lower concentration of sulphur dioxide (653 and 1306  $\mu$ g m<sup>-3</sup>) did not cause any appreciable reduction. On the contrary, higher concentration (2612 and 3918  $\mu$ g m<sup>-3</sup>) of sulphur dioxide proved to be highly toxic. Plants treated with 2612 and 3918  $\mu$ g m<sup>-3</sup> of SO<sub>2</sub> were selected for amelioration studies and calcium hydroxide was used as an ameliorating agent.

For ameliorating study set treated with 0.5%  $Ca(OH)_2$ , 0.5%  $Ca(OH)_2+2612 \ \mu g \ m^{-3} \ SO_2$ , 2612  $\ \mu g \ m^{-3} \ SO_2$ , 0.5%  $Ca(OH)_2+3918 \ \mu g \ m^{-3} \ SO_2$  and 3918  $\ \mu g \ m^{-3} \ SO_2$  were designated as  $C,C+T_1$ ,  $T_1,C+T_2$  and  $T_2$  sets, respectively. The calcium hydroxide solution was sprayed on sulphur dioxide (2612  $\ \mu g \ m^{-3} \ and 3918 \ \mu g \ m^{-3} \ SO_2$ ) treated plants once in a week with the help of a sprayer.

The study on both plants was carried out till maturity (90 d in *Brassica juncea* and 120 d in *Raphanus sativus*).the observations for various attributes were recorded in 15 d, 30 d, 45 d, 60 d, 75 d, 90 d old plants of *Brassica juncea* and *Raphanus sativus* respectively.

Chlorophyll content was estimated according to Arnon's (1949) method. Fresh leaves (100 mg) were homogenized with acetone (80%) and a pinch of sodium bicarbonate was added. The homogenate was centrifuged for 5 minutes and final volume of supernatant was made to 10 ml. by adding 80% acetone. The optical density (OD) of the extract was nm and recorded at 645 663 nm on а spectrophotometer against a blank (80 % acetone).

The amount of chlorophyll a, chlorophyll b and total chlorophyll was calculated using the following formulae-

Chlorophyll a=  $(12.7 A_{663}-2.69 A_{645}) * V/W*1000(mg g^{-1} fresh weight)$ 

Chlorophyll b=  $(22.9 A_{645}-4.68 A_{663}) * V/W*1000(mg g^{-1} fresh weight)$ 

Total chlorophyll =  $(8.02 \ A_{663}-20.2 \ A_{645}) * V/W*1000(mg g^{-1} fresh weight)$ 

#### **RESULT & DISSCUSION**

Reduction in chlorophyll a and chlorophyll b content of treated seedlings in comparison to control ones was observed at higher SO<sub>2</sub> level.SO<sub>2</sub> also reduced the stability index of chlorophyll. Chlorophyll content was estimated to determine the effect of SO2 on photosynthetic machinery of plants. Data analysis reveals that so2 exposures cause significant loss of chlorophyll in both experimental crops. Amount of chlorophyll a, chlorophyll b &total chlorophyll increased rapidly first 30 days and then remained almost changed but thereafter a sharp decline was observed in all sets. Max reduction in chlorophyll a content was recorded at 3918 µg m<sup>-3</sup> of SO<sub>2</sub> in 60 d old leaves. Maximum % decrease in chlorophyll a content was 33.67 in Brassica juncea and 58.33 in Raphanus sativus. On other hand maximum decline in chlorophyll b content was 28.24 and 39.73 % in Brassica juncea and Raphanus sativus respectively at 3918  $\mu$ g m<sup>-3</sup> of sulphur dioxide. Influence of SO<sub>2</sub> was also observed on stability index of chlorophyll. Stability index of chlorophyll in treated plant was lower in comparison to control one. Maximum reduction noticed at 3918  $\mu$ g m<sup>-3</sup> of SO<sub>2</sub>

#### Chlorophyll Extraction & Determination

1 0			2.
	Table 3a:	Estimation of some biochemical components in leaves of Brassica junc	ea fumigated with
		different concentration of sulphur dioxide at 45 d and 60 d plant	age.
Plant age(d)	)		

			45 d	l		60 d											
Attribute	concen	tration of ş	ulphur diox	ide(µg m⁻³)	CD	concent	tration of	sulphur, d	ioxide(µg n	n-*)	CD						
		0	653	1306	2612	3918	596	1%	0	653	1306	2612	3918	596	1%		
Chlorophyll a (mg.g <sup>-1</sup> f.wt.)		4.861	4.973	3.230**	3.440**	2.697**	1.193	1.275	4.086	4.166	2.877**	2.876**	2.669**	0.890	0.952		
		±0.358	±0.281	±0.272	±0.541	±0.1754			±0.283	±0.283	±0.302	±0.102	±0.046				
Chlorophyll b		1.406	1.403	1.321	1.123	1.204	0.824	0.882	1.813	1.904	1.373**	1.323**	1.301++	0.382	0.407		
(mg.g <sup>-1</sup> f.wt.)		±0.056	±0.231	±0.125	±.025	±0.102			±0.099	±0.023	±0.178	±0.178	±0.043				
Stability index chlorophyll	of	100.00	101.739	72.618	72.809	62.246	-	-	100.00	102.895	72.029	78.669	73.495	-	-		

#### Volume-3, Issue-2, April-2012

Table 3b:	Estimation of some biochemical components in leaves of Raphanus sativus
fumigated	with different concentration of sulphur dioxide at 45 d and 60 d plant age.

Plant sge(d)																		
			45 d				60 d											
Attribute	concen	tration of ş	ulphur diox	ide(µg m³)	CD	concen	stration of <u>sulphur</u> dioxide(ug m <sup>-1</sup> ) CD											
			650	1000		2010												
		U	053	1300	2012	3918	390	190	U	053	1300	2012	3918	390	190			
Chloraphyll a																		
(ma al furt)		1.121	0.983**	0.780**	0.681**	0.587**	0.000	0.120	0.948	0.864**	0.735**	0.526**	0.395**	0.028	0.083			
(mg.g., t'mt)							0.098											
Chlorophyll b		±0.022	±0.008	±0.037	±0.041	±0.013			±0.035	±0.032	±0.011	±0.038	±0.020					
c altri payn b		0.354	0.353	0.390	0.238**	0.263**	0.090	0.108	0.307	0.286*	0.255**	0.186**	0.186**	0.010	0.052			
(mg.g <sup>-1</sup> f.yd.)		±0.005	±0.002	±0.029	±0.051	±0.005			±0.021	±0.014	±0.014	±0.011	±0.009					
Stability index chlorophyll	x of	100.00	89.843	82.882	62.058	57.653	-	-	100.00	91.669	79.384	57.044	46.582	-				
				1			10	11	-1 × 1			NA	/					

#### Table 4a: Estimation of some biochemical components in leaves of Brassica juncea treated with SO2 and calcium hydroxide at 15 d and 30 d plant age

Plant aga(d)

Tiant age(u)																
	30 d															
Attribute	Treatment				CD			Ti	reatment		CD					
		-								-			_	-		
	Contro 1	c	C+T <sub>1</sub>	C+T2	T <sub>1</sub>	T2	596	196	Contr ol	c	C+T1	C+T2	T <sub>1</sub>	T <sub>2</sub>	5%	1%
Chlorophyll a	1.336	1.453	1.460	1.266	1.054	1.073			3.055	2.562	2.755	2.349	2.468	1.738		
(mg.g <sup>-1</sup> f.ut.)							0.595	0.837							1.333	1.882
	±0.294	±0.236	±0.225	±0.108	±0.124	±0.036	•	0.007	±0.315	±0.430	±0.101	±0.484	<b>±0.46</b> 7	±0.274		
Chlorophyll b	0.766	0.202	0.766	0.572	0.650	0.415			1.005	1 2 2 2	1.100	1266	1.050	1.069	0.724	1.026
(mg.g <sup>-1</sup> f.nt.)	0.700	0.785	0.700	0.575	0.050	0.415	0.564	0.796	1.225	1.282	1.159	1.500	1.059	1.008	0.754	1.050
	±0.252	±0.222	±0.252	±0.093	±0.096	±0.087			±0.237	±0.318 9	±0.087	±0.225	±0.077	±0.205		
Stability index of chlorophyll	100.00	100.17	99.726	82.388	81.060	70.789		-	100.00	89.813	92.140	86.790	82.406	65.560		
1																

# Table 4b: Estimation of some biochemical components in leaves of Raphanus sativus treated with SO<sub>2</sub> and calcium hydroxide at 15 d and 30 d plant age

Plant age(d)																
	:	15 d				30 d										
Attribute	Treatment				CD			Ti	reatment			CD				
			_					_	_					_	_	
	Contro 1	с	C+T1	C+T2	T <sub>1</sub>	T2	5%	1%	Contr ol	с	C+T1	C+T2	T <sub>1</sub>	T2	5%	1%
Chlorophyll a	0.567	0.557	0.499	0.499	0.436*	0.404*			0.814	0.793	0.693**	0.642**	0.641*	0.562*	0.082	0.116
(445-2, 1700-)			0.092	0.129					-	-	0.082	0.110				
	±0.005	±0.024	±0.049	±0.048	±0.007	±0.011			±0.003	±0.015	±0.035	±0.056	±0.008	±0.015		
Chlorophyll b	0.134	0.125	0.105*	0.107*	0.108*	0.085*			0.201	0.233	0.221	0.189	0.157	0.139	0.094	0.133
(mg.g <sup>-1</sup> f.ut.)						٠	0.026	0.036								
	±0.001	±0.014	±0.004	±0.061	±0.002	±0.005			±0.003	±0.032	±0.012	±0.045	±0.017	±0.013		
Stability index of	100.00	02.612	06 200	06 200	26.052	21.001			100.00	00.222		01.044	20.602	60.060		
cnioropnyii	100.00	97.517	80.729	80.329	70.953	71.291	-	-	100.00	99.737	88.910	81.844	78.095	09.000	-	-

Chlorophyll Tables show that content of C+T1&C+T2 sets are higher than the T1&T2 sets respectively. On 60<sup>th</sup> day % increase in chlorophyll a content of C+T1 set was 36.50 in Brassica juncea and 32.09 in Raphanus sativus. Over their T1sets.the corresponding value for C+T2 sets were 27.32 &22.58 in Brassica juncea and Raphanus sativus respectively. Level of chlorophyll b also increased in  $C+SO_2$  set than set treated with  $SO_2$  alone. The total chlorophyll contents of C+ SO<sub>2</sub> set were higher than those of SO<sub>2</sub> treated plants. The application of calcium hydroxide reduced the loss in chlorophyll stability.

Malhotra (1977) & Shimazaki *et.al.* (1980) reported that stress treatment either inhibits or increase its destruction. After treatment with SO<sub>2</sub>, chlorophyll content of plants was positively affected at low conc.; same was found to be decreased at higher conc.

## Chlorophyll Breakdown

In order to understand the mechanism of chlorophyll destruction, the effect of  $SO_2$  on  $Mg^{++}$  loss from chlorophyll (conversion of chlorophyll into pheophytin) was determined. Having entered inside the leaves,  $SO_2$  with the help of water forms sulphurous acid, which then dissociates into H<sup>+</sup> and HSO<sub>3</sub> ions and causes degradation of chlorophyll. chlorophyll a is degraded to phaeophytin through replacement of  $Mg^{+2}$  ions present in chlorophyll molecules by free H<sup>+</sup> ions, while chlorophyll b forms chlorophyllide through the removal of phytol group of the molecule (Rao and LeBlanc, 1966). however the phaeophytinization of chlorophyll caused by SO<sub>2</sub> the pH of outer occurs when solution is lower(Malhotra, 1977). Acids such as hydrofluoric and hydrochloric acids have been shown to convert chlorophyll into pheophytin with the release of Mg<sup>+</sup> (Rao and LeBlanc, 1966). Mg<sup>++</sup> is replaced by two molecules of hydrogen with a resulting change in the light absorption spectral properties of chlorophyll molecule.

## Sulphur dioxide and Chlorophyll destruction

SO<sub>2</sub> exposure resulted in simultaneous loss of both chlorophyll a &chlorophyll b. This implies that the loss of chlorophyll b was resultant upon inhibited biosynthesis of chlorophyll a (Bogorad, 1966). The decrease chlorophyll in content at higher concentration of SO<sub>2</sub> could be due to very low pH which seems to be very fatal as described by Gilbert (1968) and Grunwald (1981). Some other investigator reported that  $SO_2$  is oxidized to  $SO_4^{-2}$  through intermediary sulphite ions. Excess of Sulphite leads to break down of chlorophyll (Ricks&William, acid

1975).  $SO_2$  may react with chlorophyll to form superoxide radicals (Shimazaki et al. 1980; William and Banerjee, 1995). These radicals cause wide spread damage to membrane and associated molecules including chlorophyll pigment (Peiser and Yang, 1978).

#### CONCLUSION

Both crops (Brassica juncea & Raphanus sativus) showed negative response to elevated SO<sub>2</sub> levels, the extend of response was different between both crops. Both plants adopted some protective strategies to fight against SO<sub>2</sub> pollution. Study reports that initial concentration of SO<sub>2</sub> (653  $\mu$ g m<sup>-3</sup>) caused slight stimulation in growth of Brassica juncea thus Brassica juncea can tolerate or grow better up to a certain critical level of  $SO_2$ , the level which is higher than other crops. it was found that photosynthetic pigments were positively affected at low concentration in Brassica juncea, the same found to be decreased at higher concentration reduction in chlorophyll led to impairment of photosynthetic vis-a -vis reduction in growth and yield of plants. Chlorophyll degradation was found more in Raphanus sativus than in Brassica *juncea*. The present investigation led us to believe that calcium hydroxide acts as an antidote against SO<sub>2</sub> stress in two test crops *Brassica juncea* and *Raphanus* sativus. The yield improvement was better in Brassica juncea than that in *Raphanus* sativus, the reason behind this may be that *Raphanus sativus* is very sensitive to SO<sub>2</sub> pollution & is not able to recover from SO<sub>2</sub> stress. On other hand *Brassica juncea* being relatively resistant to SO<sub>2</sub> responded potentially to calcium hydroxide treatment and showed better growth than non sprayed plants.

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Volume-3, Issue-2, April-2012

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