# **Roadside Plants as Bio-indicators of Urban Air Pollution**

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**Abstract:** This paper describes air pollution tolerance among roadside plants exposed to varying degrees of vehicular pollutants. Evaluation of air pollution tolerance index (APTI) of 10 selected wild plant species was carried out to assess their response to ambient levels of air pollutants along the busy roadways of Bangalore. Four parameters namely total chlorophyll, ascorbic acid, pH of leaf extract and relative water content were determined and computed together to signify air pollution tolerance index (APTI) of plants. The observed significant reduction in total chlorophyll, ascorbic acid and relative water content showed inverse relationship with traffic density. Similarly, pH of leaf extract followed an exponential decrease with increase in traffic density and drifted towards acidic range. Comparison of APTI values from control to polluted sites revealed maximum reduction in Bougainvillea spectabilis while least change was noted in Peltophorumpterocarpum. Among the plants studied maximum net per cent reduction of APTI over control was seen in Bougainvillea spectabilis and Ageratum conyzoides and are considered to be sensitive species. While Peltophorumpterocarpum and Portulacaoleraceaeare tolerant species since they have shown least per cent reduction in APTI.

Keywords: APTI, chlorophyll, ascorbic acid, relative water content, Bougainvillea spectabilis

# I. Introduction

The atmosphere is a complex and dynamic gaseous system that is essential to support life on planet Earth. Rapid industrialization and urbanization coupled with increase in vehicular traffic in the urban areas has become a great threat to air quality, threatening the very existence of the living beings. Among the various categories, air pollution by automobiles is the most insidious one, which exerts highly detrimental effects on living organisms. Ambient air pollution in several large cities of India is amongst some of the highest in the world [1]. Bangalore too with its rapid growth and commercial activities harbours approximately 41.71 lakh vehicles releasing large quantities of pollutants such as oxides of nitrogen, sulphur, carbon, heavy metals, dust and particulate matter.

Among various strategies of controlling atmospheric pollution, absorption of gaseous pollutants by plants provide one of the natural ways of cleansing the atmosphere [2] and they act as effective indicators of air pollution [3]. Recent studies have explored the possibility to find out the ability of plants to remove and also act as sinks for air pollutants [4, 5].Biomonitoring is a low cost and valuable method to evaluate the effect of different air and environment pollutants [6]. There is hardly any data available on plants which act as sensitive, tolerant and resistant species to urban pollution, particularly vehicular pollution.

The air pollution tolerance index (APTI) based on four parameters, namely total chlorophyll, ascorbic acid, pH and relative water content have been used for identifying tolerance levels of these species [7, 8].APTI has been extensively used to rank plant species in their order of tolerance to air pollution [7, 9]. In a similar attempt the present study was carried out to assess APTI of 10 angiospermic plants covering the busy roadways of Bangalore city.

# 2.1 Study site

# II. Materials and methods

With a population of 8.5 million, Bangalore is the fifth largest city in India, positioned at  $12^{\circ}58^{\circ}N$ ,  $77^{\circ}34^{\circ}E$ ,  $12.97^{\circ}N$ ,  $77.56^{\circ}E$  and covers an area of  $741 \text{ km}^2$ . Situated at 1000 meters above sea level, the city experiences salubrious savanna climate with distinct rainy, winter and summer seasons. Bangalore harbours 41.71 lakh vehicles, of which 28.81 lakh are two wheelers, 7.92 lakh cars, 1.62 lakh most polluting autos and 3.36 lakh other vehicles. The sites selected for the present study includes a control site (Kommasandra - S I), a moderately polluted residential site (Ramamurtynagar - S II) and a highly polluted site (Bangalore city railway station - S III) having an approximate traffic density of <300, 43,500 and 73,800 vehicles respectively.

# 2.2 Plants

The plant species selected for the present study are

Species Family Ageratum conyzoidesL.Asteraceae BambusabambosL. VossPoaceae Bougainvillea spectabilisComm. Ex. Juss.Nyctaginaceae CynodondactylonL. Pers.Poaceae FicusreligiosaL.Moraceae MangiferaindicaL.Anacardiaceae PeltophorumpterocarpumDC. K. HeyneFabaceae PortulacaoleraceaeL.Portulacaeae RicinuscommunisL.Euphorbiaceae TerminaliacatappaL.Combretaceae

Fresh leaf samples of these plants were collected and analysed for the following parameters.

# 2.3. Total chlorophyll

The total chlorophyll was estimated principally by the method of Arnon [10]. One gram fresh leaf was macerated with 80% (v/v) chilled acetone and a pinch of magnesium carbonate in a prechilled pestle and mortar. The extract was centrifuged at 2500 rpm for 10 minutes. The process was repeated till the extract becomes colourless and the extracts were pooled and the volume was made up to 15mL. All operations were carried in the ice bath under dark condition. The absorbance was measured at 645, 663 and 750nm using UV-visible spectrophotometer.

TCh =  $20.2(A_{645})+8.02(A_{663})xV/(1000x W)$ Where, TCh= Total chlorophyll in mgg<sup>-1</sup>  $A_{645}$  = Absorbance at 645nm minus the absorbance at 750nm  $A_{663}$ =Absorbance at 663nm minus the absorbance at 750nm V = Total volume of the extract in mL W = Weight of the sample in g

# 2.4. Ascorbic acid

One gram fresh leaf was homogenized in 4mL of freshly prepared oxalic acid (0.4% w/v), filtered and centrifuged at 1000 rpm for 20 minutes. Final volume was made up to 10mL using oxalic acid. About 5mL of the extract was titrated against standardized 2,6-dichlorophenol-indophenol [11]. AA = IxSxD/Ax1/W

Where.

AA = Ascorbic acid in  $mgg^{-1}$ I = mL of indophenol used for titration S = mg of ascorbic acid reacting with 1mL indophenol D = total volume of the extract in mL A = Aliquot titrated in mL

W = Weight of the sample in g

# 2.5. Relative Water Content

The fresh weight of the leaves was determined and immersed in water over night. Turgid weights of the leaves were measured after blotting the leaves. The leaves were dried overnight in an oven at 80°C and the dry weight was determined [12]. RWC = (F-D/T-D)x100

Where,

RWC = Relative water content in %

F = Fresh weight in g

T = Turgid weight in g

D = Dry weight in g

## 2.6. pH of the leaf samples

Five grams of fresh leaves were homogenized in 10mL deionised water. This was filtered and the pH of the leaf extract was determined after calibrating pH meter using buffer solution [12].

# 2.7. APTI

The air pollution tolerance indices were determined following the method of Singh and Rao [7]. APTI = AA(TCh+pH)+RWC/10Where,

APTI = Air pollution tolerance index AA = ascorbic acid content in mgg<sup>-1</sup>

 $TCh = total chlorophyll in mgg^{-1}$ 

pH = pH of leaf extract

RWC = relative water content in %

# III. Results and Discussion

The results of analysis of total chlorophyll and ascorbic acid are presented in table 1 and pH of leaf extract and relative water content table 2.

### 3.1. Total chlorophyll

The concentration of total chlorophyll decreased in all the species growing at polluted sites when compared to the controls, with maximum reduction at S III which harbours highest vehicular density. Maximum total chlorophyll reduction was seen in *Bougainvillea spectabilis* and *Portulacaoleraceae*, moderate reduction in *Peltophorumpterocarpum, Ricinuscommunis* and *Ficusreligiosa*, while minimum reduction was observed in *Mangiferaindica, Bambusabambos* and *Cynodondactylon*. Since chlorophylls are the chief photosynthetic pigments, their content signifies growth and development of biomass and overall health status of plants. Decrease in chlorophyll content has been suggested as an indicator of SO<sub>2</sub> pollution. High amount of gaseous SO<sub>2</sub> causes destruction of chlorophyll and that might be due to the replacement of Mg<sup>++</sup> by two hydrogen atoms and degradation of chlorophyll molecules to phaeophytin [13]. A considerable loss of total chlorophyll in the plants such as SPM, SO<sub>2</sub> and NO<sub>x</sub> [5]. Pollutants such as SO<sub>2</sub>, NO<sub>2</sub> and O<sub>3</sub> cause damage to membranes and associated molecules including chlorophyll pigments [14].

Table 1:Mean con	ncentration of tota	al chlorophyll and	d ascorbic acidof selected	plant species at different sites

	Total	Total chlorophyll (mgg <sup>-1</sup> )			Ascorbic acid (mgg <sup>-1</sup> )		
Plants	S I	S II	S III	S I	S II	S III	
A.conyzoides	2.443±0.023	1.823±0.260	1.643±0.296	0.263±0.291	0.216±0.092	0.142±0.208	
B.bambos	1.991±0.156	1.610±0.301	1.382±0.316	0.111±0.256	0.098±0.276	0.093±0.104	
B. spectabilis	2.901±0.347	1.899±0.098	1.544±0.089	0.433±0.333	0.311±0.198	0.081±0.273	
C.dactylon	2.050±0.256	1.523±0.125	1.401±0.059	0.158±0.089	0.121±0.387	0.101±0.124	
F.religiosa	2.512±0.176	1.645±0.287	1.584±0.138	0.150±0.302	0.102±0.207	0.083±0.242	
M.indica	2.822±0.067	2.237±0.311	2.086±0.207	0.210±0.099	0.189±0.088	0.159±0.255	
P.pterocarpum	2.527±0.219	1.664±0.200	1.512±0.273	0.222±0.103	0.198±0.276	0.142±0.087	
P.oleraceae	2.926±0.311	2.086±0.291	1.635±0.345	0.197±0.168	0.148±0.157	0.132±0.182	
R.communis	2.323±0.249	2.088±0.089	1.403±0.301	0.261±0.298	0.255±0.298	0.232±0.076	
T.catappa	2.401±0.110	1.895±0.196	1.601±0.204	0.116±0.345	0.102±0.289	0.093±0.156	

n=15

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Table 2:Mean concentration of pH of leaf extracts and relative water content of selected plant species at different sites

	рН		Relative water content (%)			
Plants	S I	S II	S III	S I	S II	S III
A. conyzoides	6.71±0.302	$6.14 \pm 0.142$	$5.43 \pm 0.086$	85.95±0.124	71.72±0.134	68.81±0.148
B. bambos	5.68±0.178	5.24±0.257	5.31±0.244	45.37±0.323	36.36±0.366	34.21±0.198
B. spectabilis	5.76±0.079	$5.00 \pm 0.278$	4.74±0.194	67.41±0.258	53.10±0.234	49.20±0.234
C. dactylon	6.71±0.355	$6.45 \pm 0.340$	6.26±0.350	25.12±0.209	21.40±0.098	20.36±0.246
F. religiosa	7.59±0.198	7.01±0.089	6.43±0.254	75.97±0.136	61.10±0.124	55.98±0.378
M. indica	6.93±0.154	6.59±0.129	6.02±0.187	73.97±0.378	61.71±0.186	52.02±0.093
P. pterocarpum	7.01±0.257	$6.68 \pm 0.108$	6.01±0.108	69.23±0.214	62.14±0.099	50.54±0.355
P. oleraceae	6.80±0.224	$6.62 \pm 0.386$	$6.40 \pm 0.088$	76.23±0.125	70.14±0.100	64.26±0.236
R. communis	5.76±0.197	5.70±0.134	5.32±0.148	88.4±0.263	78.76±0.189	70.16±0.248
T. catappa	5.50±0.173	$5.48 \pm 0.284$	5.39±0.247	68.71±0.298	58.21±0.294	52.22±0.248

n=15

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# 3.2. Ascorbic acid

Traffic density dependent decrease in ascorbic acid content of all the species was noted. Maximum reduction in ascorbic acid content was recorded in *Bougainvillea spectabilis, Ageratum conyzoides* and *Ficusreligiosa*, moderate reductionwas noted in*Cynodondactylon, Peltophorumpterocarpum* and *Portulacaoleraceae*. While *Ricinuscommunis, Bambusabambos, Terminaliacatappas* howed least reduction. Reduction in ascorbic acid is attributed to increased rate of production of reactive oxygen species (ROS) during photo-oxidation of SO<sub>2</sub> to SO<sub>3</sub>[15]. Ascorbic acid is an antisorbic vitamin, strong reducing agent reported to play an important role in SO<sub>2</sub> reduction and it activates many physiological and defence mechanism, also maintains the stability of plant cell membranes during pollution stress. Its reducing power is directly proportional to its concentration [9].

## 3.4. pH

All the plant samples collected from polluted site exhibited a shift in pH towards acidic range. High acidic pH was found in *Bougainvillea spectabilis*(4.74),*Bambusabambos*(5.31) *and Ricinuscommunis*(5.32). The presence of SO<sub>2</sub> and NOx in the ambient air causes a change in pH of the leaf sap towards acidic range [17]. Upon diffusion of SO<sub>2</sub> through stomata, gaseous SO<sub>2</sub> dissolves in water to form sulphites, bisulphate and their ionic species with the generation of protons influencing the cellular pH [18]. It is therefore opined that the pH change towards acidic range observed in most species is due to entry of SO<sub>2</sub> into leaf mesophyll tissue.

### 3.3. Relative water content

Plants growing in polluted site showed reduced relative water content, with maximum reduction in *Mangiferaindica, Bougainvillea spectabilis,* moderate reduction was observed in *Peltophorumpterocarpum, Ficusreligiosa, Bambusabambos* and minimum reduction in *Portulacaoleraceae* and *Cynodondactylon.* Water content is an important factor which determines the physiological status of the plant. The relative water content is associated with protoplasmic permeability in cells.Loss of water and dissolved nutrients results in early senescence of leaves. The relative water content in a plant body helps in maintaining its physiological balance under stress conditions including air pollution stress [16]. The reduced relative water content indicates disturbed physiological status in the plants due to pollution [14].

#### **3.5.** Air Pollution Tolerance Index

The calculated APTI decreased progressively from control to highly polluted sites, table 3.

Plants	S I	S II	S III	
	APTI	APTI	APTI	% ROC
Ageratum conyzoides	9.15	6.48	5.12	44.06±0.251
Bambusabambos	4.62	3.70	3.48	24.64±0.109
Bougainvillea spectabilis	9.21	5.52	4.57	50.41±0.307
Cynodondactylon	2.65	2.33	2.27	14.22±0.298
Ficusreligiosa	7.74	6.19	5.66	26.87±0.273
Mangiferaindica	8.60	6.33	5.03	41.51±0.271
Peltophorumpterocarpum	7.13	6.67	6.36	10.84±0.346
Portulacaoleraceae	7.81	7.16	6.93	11.28±0.271
Ricinuscommunis	8.90	8.07	8.07	19.43±0.192
Terminaliacatappa	6.96	5.94	5.28	24.13±0.278

Table 2. ADTI and their	non contradivation area	acentral (0/ DOC)
Table 3: APTI and their	per cent reduction over	COILLOI (% KOC)

n=15

The APTI of *Bougainvillea spectabilis* showed maximum decrease from S I (9.21) to S III (4.57), *Cynodondactylon*showed least difference from control to polluted site with 2.65 to 2.27 respectively. The mean per cent reduction of APTI over control showed highest reduction in *Bougainvillea spectabilis* (50.41), *Ageratum conyzoides*(44.06)and *Mangiferaindica*(41.51), while least reduction was seen in *Cynodondactylon* (14.22), *Portulacaoleraceae*(11.28) and *Peltophorumpterocarpum*(10.84).

The results obtained for *Cynodondactylon* and *Bougainvillea spectabilis* are comparable to the results reported by Ramakrishnaiah and Somashekar [14]. The observed APTI values for *Ficusreligiosa* and *Mangiferaindica* closely agrees with Avnish [19]. The APTI results obtained for *Bambusabambos* (3.48 - 4.62) and *Terminaliacatappa* (5.28 - 6.96) in the present study were lower when compared to that reported by Agbaire [20].

Many reports have indicated that the species with low index values are sensitive to air pollution and *vice versa*[21, 22, 23]. The level of APTI exclusively depends on the intrinsic nature of each species since the level of total chlorophylls, ascorbic acid, pH and relative water contents varies greatly from species to species

and they are not directly comparable. It is important to draw conclusions based on the differences in the amount of changes (%ROC) observed within the species.

Due to their higher reduction in air pollution tolerance index over their control counterparts *Bougainvillea spectabilis* and *Ageratum conyzoides* are considered as more sensitive species, whereas *Ficusreligiosa, Bambusabambos* and *Terminaliacatappa* with moderate changes are regarded as tolerant species. While *Peltophorumpterocarpum Portulacaoleraceae* with least APTI reduction over control counterparts are regarded as relatively resistant species. The plants of the former category can be effectively used as bio-indicators of automobile exhaust pollution, whilst the resistant plants can be employed as sinks for vehicular pollutants. It is evident that the plants growing alongside busy roadways have exhibited significant reduction in total chlorophyll, ascorbic acid, relative water content and pH of leaf extracts and has resulted in the substantial reduction of air pollution tolerance in plants. This is because of the pollutants released by automobiles.

#### IV. Conclusion

Air pollution has been considered as a potential selection force for plants, therefore the species growing in such adverse roadside environment present the best material to ascertain the levels of susceptibility rather than drawing interpretations from measurements obtained from lab conditions. Furthermore, APTI determination provides a reliable method for screening large number of plants with respect to their response to air pollutants.

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