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# EFFECTS OF GA3 AND THIOUREA ON PROTEIN, PEROXIDASE ACTIVITY AND CHLOROPHYLL STATUS IN SEEDLINGS OF ANOLA

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#### ABSTRACT

The experiment was conducted in the laboratories of the Department of Horticulture. G.A.U., Anand during 1993. Three concentration each of GA<sub>3</sub> and thiourea, distilled water and control (total 8 treatments) were given to the seeds of Anola with 36 h. of subsequent soaking and drying thrice. Observations were recorded 6, 8 and 10 week intervals on protein peroxidase specific activity and chlorophyll content. GA<sub>3</sub> 500 ppm was effective in increasing the protein content of leaves and stem whereas no treatment was effective in enhancing the peroxidase specific activity in leaves, stems and roots. GA<sub>3</sub> 250 ppm was again helpful in increasing the total chlorophyll content of the leaves at all the three intervals.

KEY WORDS: Anola, seedling, protein, peroxidase activity, chlorophyll

Anola (Phyllanthus emblica Linn.) is one of most important fruit crops of the tropics and ubtropics and broadly cultivated for its fruits rich n vitamin c and commercially exploited for hair lyes, medicine, ink etc. (Ram, 1983). Till recently. he main emphasis has been given on the yield and ruit preservation and very little on improvement in eed germination in anola raised for rootstocks. The henomenon of seed germination is associated with hanges in several protein constituents. Proteins are during germination down oncomitant rise in amino acids and amides ollowed by de novo synthesis of protein and nzymes in the germinating parts of embryo. eroxidases widely distributed in plants and eroxidase activity is related to cell growth and ifferentiation. The aim of the present investigation as to find out the possible role of protein and croxidase enzyme for altering seed germination.

# MATERIALS AND METHODS

Present investigations were carried out at the Department of Horticulture, Gujarat Agricultural University, Anand Campus during 1993 in randomised design with four completely replications and eight treatment viz., concentrations each of GA3 and Thiourea (250, 500 and 750 ppm), distilled water and control (untreated). Pre-sowing treatments of seed soaking and subsequent drying processes were carried out for 36 hr followed by a surface sterilisation with 0.2 percent mercuric chloride. Treated seeds were allowed to germinate in pots and for chemical analysis samples were taken at 6 weeks (Ist stage), 8 weeks (IInd stage) and 10 weeks (IIIrd stage) interval. Protein percentage peroxidase specific activity (Change in O.D/hr./mg. protein) in leaves. stem, leaves and roots and chlorophyll content (mg/g fresh weight) were determined by employing normal methods.

| Table 1.  | Effect of GA  | and thioures on  | protein content | (%) at differe     | nt stages in anola seedling |          |
|-----------|---------------|------------------|-----------------|--------------------|-----------------------------|----------|
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| 1 37 50                |       | Leaves |       |       | Stem    |       |       | Roots  |        |
|------------------------|-------|--------|-------|-------|---------|-------|-------|--------|--------|
| Treatments             |       | Stages |       |       | Stages  |       |       | Stages |        |
| 7. <del></del>         | 1st.  | IInd   | Hird  | . Ist | Hnd     | IIIrd | lst   | IInd   | filird |
| Control                | 15.43 | 12.15  | 10.17 | 7.00  | 9.22    | 8.70  | 7,74  | 8.50   | 11.46  |
| Distilled water        | 17.15 | 15.26  | 12.26 | 8.75  | 10.17   | 8.90  | 8.74  | 9.88   | 10.40  |
| GA <sub>3</sub> 250ppm | 25.20 | 24.37  | 22.36 | 17.50 | 21.30   | 19.70 | 18.65 | 20,10  | 21.94  |
| GA <sub>3</sub> 500ppm | 26.60 | 25.40  | 24.50 | 16.24 | 20.15   | 18.55 | 17.54 | 21.20  | 22.08  |
| GA <sub>3</sub> 750ppm | 21.70 | 19.80  | 13.00 | 16.00 | 19.70   | 15.67 | 14.26 | 17.00  | 19.60  |
| Thiourea 250ppm        | 24.52 | 22.93  | 18.82 | 14.35 | . 15.26 | 12.88 | 15.00 | 17.80  | 18.60  |
| Thiourea 500ppm        | 25.20 | 24.60  | 22.64 | 15.00 | 16.58   | 15.50 | 16.67 | 20.18  | 21.80  |
| Thiourea 750ppm        | 23.00 | 21.92  | 15.05 | 14.00 | 15.40   | 10.08 | 16.00 | 19.20  | 20.00  |

### RESULTS AND DISCUSSION

#### Protein content

Leaf protein content (Table 1) decreased with advancement in period of germination. Maximum protein content was recorded with GA<sub>3</sub> 500 ppm at all three stages while it was minimum in control. The influence of GA<sub>3</sub> on enzymes of carbohydrate, protein and nucleic acid metabolism has been explained by Key (1969). The protein in stem was found highest at second stage under all treatments. The level of protein in roots increased as germination period advanced. It appears that increase in enzyme activity which is associated with protein synthesis resulted in protein level changes in the plants. The decrease in leaf protein at third stage may be due to utilisation of protein in growth and development processes of plant.

### Peroxidase specific activity

It is evident (Table 2) that control recorded the highest peroxidase activity in leaves at all three stages. Under thiourea treatments 750 ppm gave highest activity and increase in GA3 level upto 750 ppm increases the activity at all three stages.

Similarly, in stems, control registered the highest at first and third stages whereas, at second stage, distilled water treatment was better over rest of the treatments. The peroxidase activity in roots increased with advances in period of germination. At all three stages control and GA3 supported maximum and minimum peroxidase activity respectively.

In general, different concentrations of GA, and thiourea decreased the enzyme activity in leaves, stem and roots. The presence of peroxidase in seeds resulted into relatively high activity of interaction in plant metabolism which might destroy and cause imbalance in the growth regulator proportions, which results in poor and late germination (Rao, 1975). The findings are in conformity with the result of Chetal and Nainawatee (1976).

## Total chlorophyll

The treatments showed a remarkable difference among the treatments on total chlorophyll content (Table 3) in leaves. The chlorophyll level increased with advancement in the crop growth. The maximum level was recorded

Table 2. Effect of GA3 and thiourea on specific peroxidase activity at different stages of anola seedlings (change in O.D./hr./mg protein)

|                        |        | Leaves |       |        | Stem  |       |        | Roots  |         |
|------------------------|--------|--------|-------|--------|-------|-------|--------|--------|---------|
| Treatments             | Stages |        |       | Stages |       |       | Stages |        |         |
|                        | İst    | Hnd    | Hird  | Ist    | IInd  | Illrd | lst    | lind   | IIIrd   |
| Control                | 0.665  | 2.754  | 2.562 | 4,208  | 2.672 | 3.024 | 0.858  | 11.285 | 15,778  |
| Distilled water        | 0.604  | 2.238  | 1.422 | 3.757  | 3,103 | 2,178 | 0.740  | 9,945  | 13,622  |
| GA3 250ppm .           | D.13X  | 1.470  | 0.210 | 2.576  | 0.262 | 0.582 | 0.184  | 1.412  | 7,986   |
| GA: 500ppm             | 0.258  | 1.925  | 1.326 | 1.418  | 0.354 | 1.464 | 0.480  | 5 257  | 10.848  |
| GA <sub>3</sub> 750ppm | 0.264  | 1.854  | 1.002 | 1.719  | 0.603 | 0.948 | 0.162  | 2.168  | 6.386   |
| Thiourea 250ppm        | 0.397  | 1.134  | 0.342 | 0.984  | 1.762 | 0.714 | 0.288  | 2,727  | 11.252  |
| Thiourea 500ppm        | 0.356  | 1.158  | 0.720 | 1.600  | 1.969 | 1.620 | 0.195  | 3.856  | 15.906  |
| Thiourea 750ppm        | 0.406  | 1.853  | 1.188 | 0.754  | 1,996 | 1.578 | 0.186  | 6.000  | 15 9(y) |

Table 3. Effect of GA3 and thiouren on total chlorophyll content in leaves of anota seedling (mg/g fresh weight)

| Treatments             |       | Stages |         |  |
|------------------------|-------|--------|---------|--|
|                        | İst   | IInd   | Illrd   |  |
| Control                | 0.441 | 1.074  | - 1,436 |  |
| Distilled water        | 0.450 | 1.047  | 1.484   |  |
| GA <sub>3</sub> 250ppm | 0.750 | 1.568  | 1.755   |  |
| GA3 500ppm             | 0.712 | 1.456  | 1.701   |  |
| GA <sub>3</sub> 750ppm | 0.691 | 1.541  | 1.622   |  |
| Thiourea 250ppm        | 0.685 | 1.392  | 1.586   |  |
| Thiomea 500ppm         | 0.664 | 1.371  | 1.532   |  |
| Thiourea 750ppm        | 0.627 | 1.293  | 1.584   |  |

with GA<sub>3</sub> 250 ppm at all three stages and minimum level under distilled water at stage second and control at stage first and third. Similar variation in chlorophyll status was also reported by Shinde et al. (1989).

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# SOIL FERTILITY CAPABILITY CLASSIFICATION OF PROBLEM SOILS OF TIRUNELVELI, TUTICORIN AND KANYAKUMARI DISTRICTS OF TAMIL NADU

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#### ABSTRACT

Surface and sub surface soil samples pertaining to 7, 9 and 4 soil sub groups in Tirunelveli, Tuticorin and Kanyakumari districts respectively were studied during July - September 1994 for the assessment of fertility level and the preparation of the soil fertility capability classification. The main condition modifiers selected were 'e', 'k', 'b', 's', 'h', 'n' and 'g' for Tirunelveli district and 'e', 'k', 'b', 's', 'v', 'h', 'g' and 'n' for Tuticorin district and 'h', 'e', 'g', 'b' and 'k' for Kanyakumari district. All the soils of three districts are low in available nitrogen and hence in addition to the above modifiers, the low available N status was also considered to be the local modifier.

KEY WORDS: Fertility capability classification, problem soils, soil fertility.

The soil fertility capability classification (FCC) system was designed to group soils having similar limitations of fertility management (Buol, 1972). It provides a guide for the extrapolation of the fertilizer response experience. Among the various approaches in providing information on the potential of the soil for crop production, soil fertility capability classification is one which lays emphasis on the components of soil fertility within 50 cm layers from the surface. An attempt has been made to use this concept for the problem soils of

three southern districts of Tamil Nadu viz., Tirunelveli, Tuticorin and Kanyakumari.

# MATERIALS AND METHODS

Seven representative sites of problem soils in Tirunelveli, nine in Tuticorin and four in Kanyakumari districts were selected during July - September, '94 for this purpose. Samples were collected from upper 20cm or plough layer whichever was shallower. Subsoil samples within 50 cm of the surface were collected depending on