

**FORMULATION AND EVALUATION OF MICRONUTRIENT MIXTURE
FOR FOLIAR APPLICATION IN TC BANANA (*Musa* sp.)
var. NENDRAN**

by

PREMALATHA. A

(2014 - 11 - 211)

THESIS

**Submitted in partial fulfilment of the
requirements for the degree of**

MASTER OF SCIENCE IN AGRICULTURE

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**DEPARTMENT OF SOIL SCIENCE AND AGRICULTURAL CHEMISTRY
COLLEGE OF AGRICULTURE
PADANNAKKAD, KASARGOD – 671 314
KERALA, INDIA**

2016

DECLARATION

I, hereby declare that this thesis entitled “**FORMULATION AND EVALUATION OF MICRONUTRIENT MIXTURE FOR FOLIAR APPLICATION IN TC BANANA (*Musa* sp.) var. NENDRAN**” is a bonafide record of research work done by me during the course of research and the thesis has not previously formed the basis for the award to me of any degree, diploma, associateship, fellowship or other similar title, of any other University or Society

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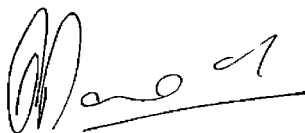
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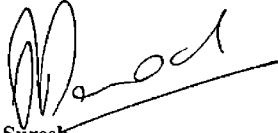
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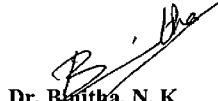
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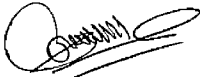
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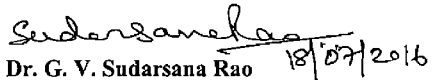
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LIST OF ABBREVIATIONS

%	-	Per cent
@	-	at the rate of
°brix	-	Degree brix
B	-	Boron
BCR	-	Benefit / cost ratio
Ca	-	Calcium
CD	-	Critical difference
CEC	-	Cation exchange capacity
Cl	-	Chlorine
cm	-	Centimeter
Cu	-	Copper
dSm ¹	-	deci Siemens per meter
EC	-	Electrical conductivity
<i>et al</i>	-	And others
Fe	-	Iron
Fig	-	Figure
g	-	Gram
ha ¹	-	Per hectare
K	-	Potassium
KAU	-	Kerala Agricultural University
kg	-	Kilogram
kg ha ⁻¹	-	Kilogram per hectare
l	-	Litre
m	-	Meter
MAP	-	Months after planting
Mg	-	Magnesium

mg g ¹	-	Milli gram per gram
mg kg ¹	-	milligram per kilogram
ml	-	Milli litre
mm	-	Milli meter
mmol g ⁻¹	-	Milli moles per gram
Mn	-	Manganese
Mo	-	Molybdenum
MT/ha	-	Metric tonnes per hectare
N	-	Nitrogen
NS	-	Not significant
pH	-	Soil reaction
ppm	-	parts per million
S	-	Sulphur
SE	-	Standard error
Rs ha ¹	-	Rupees per hectare
TC	-	Tissue culture
t ha ¹	-	Tonnes per hectare
TSS	-	Total soluble solids
viz	-	namely
Zn	-	Zinc

Introduction

1. INTRODUCTION

Micronutrients are essential for crop growth and are equally important as primary and secondary nutrients. Though their requirement is low, they often make a huge variation in yield and difference in quality of crop produce if there is a deficiency. Micronutrients have an important role in balanced plant nutrition for the stabilization of crop yield and quality of produce. These nutrients play a vital role in regulating the metabolic activities of plants and animals including human beings. Micronutrient deficiencies in soils and crops have become prevalent in the recent years due to several factors like intensive cropping, loss of top soil by erosion, leaching, decreased use of farmyard manure, increased use of high analysis fertilizers and lack of proper liming of acid soils (Singh, 2003, Rattan and Sharma, 2004, De and Rai, 2005). The deficiency of micronutrients has become major constraint to productivity, stability and sustainability of soils (Bell and Dell, 2008).

Systematic soil studies and analysis of more than 2.50 lakh soil samples in 20 states of India by All India Coordinated Research Project indicated the extent of deficiency in zinc (49%), B (33%), Fe (13%), Mo (7%), Mn (4%) and Cu (3%) (Singh, 2009). Another study by Shukla *et al.*, 2014 indicated that micronutrient deficiencies are rampant in India and on an average 43.0, 12.1, 5.4, 5.6 and 18.3 per cent soils are deficient in Zn, Fe, Cu, Mn and B, respectively. Presently, the trend of micronutrients deficiency is changing from single nutrient deficiency to multi-micro nutrients deficiencies due to depletion in soil fertility and this becomes an emerging issue in agriculture, relating to crop health and human health.

According to KSPB (2013), after a detailed study in all 14 districts reported that Kerala soils are deficient in Zn (12%), Cu (15%) and B (59%). The acid leaching environment of Kerala soils is not conducive for retention of boron arrested by highly porous nature dominated by low activity clays, which resulted in

widespread deficiency of B Iron and manganese deficiency is noticed some zones in like Onattukara and coastal sandy tracts Keeping this in view now Government of Kerala had notified the use of zinc, copper and boron for foliar as well as soil application

Banana (*Musa* sp) is the second most important fruit crop of India after mango It is one of the oldest fruit crop known to mankind and it is a rich source of carbohydrates, vitamins and minerals which are essential components of human diet Because of their commercial status, banana and plantains are referred usually as “Poor man’s apple”

At present, banana is grown in around 150 countries across the world on an area of 4.84 million ha producing 95.6 million tonnes of fruits (FAO Stat, 2011) Among horticultural crops, contribution of banana to Agricultural Gross Domestic Product (AGDP) is the highest (Smgh, H P, 2007) India is world’s leading producer of banana with an annual production of 297.24 lakh tonnes which accounts for 33.4 per cent of total fruit production with a productivity of 37.0 MT/ha In Kerala banana is grown in 34.5 lakh ha with a production of 52.8 million tones and productivity of 15.3 t/ha (Saxena, 2015) and the productivity is low

In vitro multiplication is an advanced technology over traditional method of banana propagation with respect to uniformity, vigor, disease-free planting material and true to type plants Tissue culture plants produces 20 per cent higher yield than conventional method due to good quality planting material Nendran is a popular variety of banana extensively cultivated in Kerala which is relished as fruit and vegetable as well as used for processing

Banana requires more nutrients than any other commercially cultivated crop and the various nutritional deficiencies and disorders affect its growth and yield Micronutrients such as Zn, B, Mo and Cu had been reported to be essential for the growth and development of banana plants (Srivastava, 1964a & b) Deficiencies of

Zn, Cu, Fe and Mo affect the growth and production of banana (Charpentier and Martin, 1965) Application of micronutrients in banana increases the growth, yield and quantity of banana (Pathak *et al* , 2011)

Application of essential nutrients in an appropriate balance is fundamental for various physiological processes in plants Primary nutrients play a vital role in promoting the plant vigour and productivity, whereas micronutrients like zinc, boron, copper and molybdenum perform a specific role in the growth and development of plant, quality of produce and uptake of major nutrients

The fertilizers applied through soil are needed in higher quantities because some portion leaches down and some does not become available to the plants due to complex chemical reactions happening in soil or adverse soil conditions hindering uptake The foliar application, therefore, offer a viable alternative way of applying nutrients to fruit plants in such conditions Micronutrients availability can be enhanced by foliar application of the appropriate mineral forms (Alloway, 1986, House and Welch, 1989) Micronutrient content and uptake by plants is better enhanced with foliar application

The multi-micronutrients mixture facilitates the application of wide range of plant nutrients in the correct proportion and to suit the specific requirements of a crop in different stages of growth and are more relevant under site specific nutrient management (Hegde *et al* , 2007)

Multi-nutrient foliar feeding products are the most effective way to correct nutrient deficiencies and disorders and thus resulting in increased crop growth (Mona *et al* , 2012)

Keeping all these points in view, the present study was undertaken with the following objectives

- ❖ To Prepare a micronutrient mixture formulation containing zinc, iron, boron, copper, manganese and molybdenum for foliar spray
- ❖ To evaluate stability of formulation and its keeping quality
- ❖ To investigate the effect of this formulation on growth, yield and quality of fruits in banana (*Musa* sp) var Nendran

Review of Literature

2. REVIEW OF LITERATURE

Micronutrients exist in very small amounts in both soils and plants, but their role is frequently as important as the primary and secondary nutrients. Essential micronutrients include six elements viz zinc, boron, manganese, iron, copper, and molybdenum (Stevenson, 1986). Micronutrients have assumed increasing importance in crop production under present day exploitative agriculture. Intensive cultivation of high yielding varieties and use of high analysis fertilizers and limited use of manures along with restricted recycling of plant residues are some important factors which have led to accelerated exhaustion of soil micronutrients and this in turn limits the crop production. The availability of the essential micronutrients to plants is often poorly related to their total quantity in the soil.

The micronutrients are essential for the proper biochemical transformations within the plant body, so as to yield the desired end products. Zn is essential for protein and auxin production, Cu is a constituent of cytochrome oxidase, Fe helps in photosynthesis while Mn is essential for photosynthesis, carbon assimilation and nitrogen metabolism (Mandavgade *et al*, 2015). Widespread micronutrient deficiencies have been found in many countries all over the world and their deficiency scenario has changed very much in Indian soils and crops during the last 4 decades from 1968 to 2008 (Singh, 2009).

2.1 BANANA

Banana is an important commercial fruit crop in tropical and sub-tropical regions of the world. In India, it is grown in different states under different climatic conditions (Butani *et al*, 2012). The edible banana is believed to have originated in the hot tropical regions of South-East Asia (Spiden, 1926, Suar, 1952). It comes under the family *Musaceae*. It is generally harvested when green between 70 to 100 per cent maturity and ripened before consumption (Thomas *et al*, 1968a).

Banana is globally ranked fourth, next to rice, wheat and maize in terms of gross value of production. Presently, it has emerged as the major cash-subsistence crop across all parts of the world (Robinson, 1996) as it is a complete fruit-food with delicious taste, necessary energy and health giving nutrients (Anonymous, 1969) along with pleasant flavor. On account of these properties, it is a staple food for millions of people all around the world. It is in great demand in fresh as well as processed form all over the world and gained commercial popularity in the international fruit trade (Thomas *et al* , 1968b). Bananas provide a good source of nutrients for both human and animal consumption.

Among the major producers in the world, India alone accounts for 27.43 per cent of fruits (26.2 million tonnes) followed by Philippines, producing 9.01 million tonnes and China, Brazil and Ecuador, altogether with production ranging from 7.19 to 8.21 million tonnes (FAO Stat, 2011).

Generally, Banana is a heavy consumer of nutrients and requires large quantities of nutrients for its growth, development and yield (Hazarika and Ansari, 2010) and so it is considered as an exhaustive or nutrient mining crop.

2.1.1. Tissue culture banana

Traditionally, banana is grown as a perennial crop where the plant is allowed to produce continuous shoots from an underground stem. But, the yield falls after three to five years and declines rapidly after ten to fifteen years. So the need of shifting to cyclic replacement with a new plantation comprising cycles of one crop and one ratoon has been realized only recently in most Asian countries.

Through conventional propagation techniques, there is a difficulty with obtaining large number of uniform disease free plants with high yielding potential which in turn limits the productivity of banana. Another important problem faced by the growers is the staggered flowering (variability in time of flowering). Tissue

culture technology enabling the rapid production of a large quantity of uniform disease free plants from a single plant showing good genetic potential is gaining importance in recent days (Sheela and Nair, 2001)

With the increasing demand and vast export potential coupled with the farmer's desire to grow *m-vitro* propagated banana on a large area, it is becoming increasingly important for rapid multiplication of quality planting material (Ray *et al* , 2006)

In micropropagation technique, it is desirable to produce plantlets that can grow better after transplanting into the soil. So, acclimatization is the most crucial process during banana micropropagation as the *m vitro* raised plantlets are not readily adapted for *m vivo* conditions (Vasane and Kothari, 2006)

Geetha *et al* (2004) reported that the average bunch weight (12.43 kg) and number of hands per bunch of TC-derived Nendran was 35 per cent higher than that of the conventional suckers

Robinson *et al* (1993) claimed that there was an increase in height and girth of tissue culture plants over the suckers. An increase in yield of 25.63 per cent was obtained in tissue culture plants over the plants from suckers (Sheela and Nair, 2001)

2.1.2. Nendran banana

Among the banana varieties grown in India, the French Plantain cultivar 'Nendran' belonging to the 'Plantain' group (*Musa* AAB) is the most popular variety among growers and consumers, particularly in Tamil Nadu and Kerala for domestic and export markets (Mulagund *et al* , 2015). The cultivators of Agasthiyamalai ranges call this variety as "King of Banana". The shelf life of Nendran banana is high as compared to other cultivars. So, the Nendran fruits are exported to European and Arabian countries (Das, 2010)

Nendran fruit is used for culinary purposes as well as for processing. Bunch has average of 5-6 hands weighing about 12-15 kg in the popular types.

Venugopal (2008) reported that Nendran banana is mostly cultivated in homesteads and in well-drained rice fields by small and marginal farmers of Kerala. It is the most popular commercial cultivar which is loved much by cultivators and fruits have excellent fruit quality, multiple uses and sustained income.

Das (2010) reported that fruit pulp of Nendran contains water soluble vitamins like thiamin (Vitamin B₁), Riboflavin (Vitamin B₂), Niacin (Vitamin B₃), Ascorbic acid (Vitamin C), various amino acids, proteins and nutrients in substantial amount which support the daily diet needed for human beings.

2.2 ESSENTIALITY OF MICRONUTRIENTS

Micronutrients are required in minute quantities as compared to those of macro nutrients but these nutrients play a vital role in plant metabolism (Benepal, 1967, Katyal, 2004). Also scarcity of these trace elements in soils resulted in poor crop yields (Udode- Haes *et al* , 2012). Micronutrient plays many complex roles in plant nutrition and plant production, as most of micronutrients are specific to functioning of various enzyme systems (Kazi *et al* , 2012). Micronutrients play a catalytic role in nutrient absorption and balancing of other nutrients (Singh and Kalloo, 2000).

The essential micronutrients are metals (except B and Cl) and its uptake is affected by soil (Lindsay, 1991, Lake *et al* , 1984), plant (Barber, 1995, Marschner, 1995) microbial and environmental factors (Romheld and Marschner, 1986, Clark and Zeto, 2000).

2.2.1. Essentiality of Zinc

Zinc (Zn) was discovered as an essential plant nutrient by Sommer and Lipman in 1926. Zinc is essential for proper growth and development of plant. Zinc is now being regarded as the third most important limiting nutrient element in crop production after N and P. Synthesis of plant hormones and balancing intake of P and K inside the plant cells are depending upon Zn nutrition.

Zinc is important as a component of enzymes for protein synthesis and energy production and maintains the structural integrity of biomembranes. Zn is required for the synthesis of carbohydrate metabolism, protein synthesis, internode elongation for stem growth and in pollen formation (Shukla *et al*, 2009). It is also vital for the oxidation processes in plant cells and helps in the transformation of carbohydrates and regulates sugar in plants.

Under Zn deficient conditions, flowering and fruit development are reduced and maturity is delayed which resulted in lower yield, poor quality and sub-optimal nutrient use efficiency (Gupta, 1995). Zn^{2+} ions at low concentration (0.01 ppm) slightly enhance the activity of tryptophan synthesis leading to biosynthesis of auxin (Horak *et al*, 1976).

Zinc deficiency in plants resulted in stunted growth, little leaf and fruit sizes which are attributed with IAA metabolism (Marschner, 1995). Application of zinc was found to increase the green pigments of necrotic leaf of plants (Srivastava and Singh, 2003).

2.2.2. Essentiality of Boron

Boron has been known to be a constituent of plants since 1857 (Tandon, 1989). Warrington (1923) was the first to really prove the essentiality of boron. According to Truog (1940), the importance of boron is spelt as follows, ‘ plants

will not make growth without boron any more than without phosphorus or potassium which they require in considerable amounts'

Boron is essential for plant growth, new cell division in meristematic tissue, translocation of sugar, starch, nitrogen, phosphorus, certain hormones, synthesis of amino acids and protein, regulations of carbohydrate metabolism, development of phloem. In the absence of adequate supply, middle lamella of new cell develops poorly and phloem tubes break down (Edmond *et al* , 1997). The primary role of B in plants is to improve Ca metabolism and improved solubility and mobility of Ca and helps the absorption of nitrogen.

It also involves in lignification, growth regulatory metabolism, phenol metabolism and integrity of membranes, root elongation, DNA synthesis, pollen formation and pollination (Shukla *et al* , 2009). Increased B application enhances root elongation in acidic and high-aluminium soils (Blevins and Lukaszewski, 1998).

2.2.3. Essentiality of Iron

The essentiality of iron (Fe) for plant growth was established by Gris as early as 1843. Iron acts as a constituent of various enzymes such as cytochrome oxidase, catalase and nitrogenase. It helps in the synthesis of chlorophyll. 60-80 per cent of the iron content of leaves is found in the chloroplast. Fe is a key element in various redox reactions of respiration, photosynthesis and reduction of nitrates and sulphates (Wallihan *et al* , 1958, Reddy and Reddi, 2002).

It is a component of flavoprotein like FMN (Flavin Mono Nucleotides) and FAD (Flavin Adenosine Dinucleotide). The leghaemoglobin present in the root nodules of leguminous crops contains iron as an essential constituent (Gupta and Gupta, 2005). Iron is a constituent of a large number of metabolically active compounds like cytochromes, haeme and non-haeme enzymes and other functional metalloproteins such as ferredoxin and haemoglobin.

As redox-active metal, it is involved in photosynthesis, mitochondrial respiration, nitrogen assimilation, hormone biosynthesis (ethylene, gibberellic acid, and jasmonic acid), production and scavenging of reactive oxygen species, osmoprotection and pathogen defense (Hansch and Mendel, 2009) It may also be associated with organic acid metabolism (citric acid, malic acid and ascorbic acid)

Fe deficiency is common in alkaline soil with typical chlorosis, the young leaves turn yellowish with veins remaining green. Iron application increased the levels of all leaf pigments, but the extent of increase in level depends on the pigment affected (Srivastava and Singh, 2003)

2.2.4. Essentiality of Manganese

The necessity of Mn for the growth of autotrophic and heterotrophic plants was first proposed by Mchargue (1922). Manganese is necessary for chlorophyll formation for photosynthesis, respiration and for the activity of several enzymes like oxidase, peroxidase, dehydrogenase, kinase and decarboxylase. For optimal growth and development plants have to accumulate at least 30 mg Mn kg⁻¹ dry weight in tissues regardless of plant species (Marschner, 1995)

Under deficient condition, the concentration of manganese in leaves is less than 15 ppm and in toxic condition it becomes more than 300 ppm. Mn is involved in the oxygen evolving system in photosynthesis PS II (water oxidizing enzyme complex). It is essential for photolysis of water and also it is involved in nitrogen fixation. It acts as a predominant metal ion in Krebs's cycle (Gupta and Gupta, 2005)

Manganese is only moderately mobile in plant tissues so symptoms appear on younger leaves first, most often in those leaves just reaching their full size. Mn availability is reduced in high pH calcareous soils but is often very high in the acid soil commonly chosen for tropical fruit production. Over liming of the soils as well as well drained, poor, coastal sandy soils can induce Mn deficiency. Mn deficiency

causes a light green mottle between the main veins. A dark green band is left bordering the main veins while the interveinal chlorotic areas become pale green or dull yellowish colour (Gupta and Gupta, 2005)

Soil application of manganese can be ineffective due to immobilization especially in heavier soils or soils which have been over limed. Two to three sprays of 0.1 per cent manganese sulphate can be recommended (Gupta and Gupta, 2005).

2.2.5. Essentiality of Copper

Lipman and MacKinney (1931) demonstrated the necessity of copper for the vegetative growth and reproduction of higher plants. Copper is essential for photosynthesis and mitochondrial respiration, carbon and nitrogen metabolism, oxidative stress protection and cell wall synthesis. It acts as a component of phenolase, lactase, tyrosine and ascorbic acid oxidase. The chloroplast possesses a Cu containing proteins as plastoquinone and plastocyanin, which are essential for electron carrier in photosynthesis (Gupta and Gupta, 2005)

2.2.6. Essentiality of Molybdenum

Arnon and Stout (1939) demonstrated for the first time that molybdenum is essential for higher plants. The essentiality of molybdenum for plant growth can be understood from the expression of Arnon and Stout (1939) as ' a gram of molybdenum may harness more energy, through greater conversion of sunlight into plant materials, than can be obtained from a gram of uranium '. Mo ions are part of enzymes nitrate reductase and nitrogenase. Nitrate reductase is responsible for reduction of nitrate to nitrite during N assimilation in plants. Nitrogenase converts nitrogen gas to ammonia in nitrogen-fixing microorganisms.

Mo plays a role in breakdown of purines such as adenine and guanine because of its essentiality as part of enzyme xanthin dehydrogenase. It is an essential part of oxidase enzyme which converts abscisic acid aldehyde to ABA (Gupta and Gupta,

2005) Mo deficiency is observed in many soils and pasture legumes, vegetables and occasionally in fruits, it is very rare in flower crops

2.3 MICRONUTRIENT DISTRIBUTION IN INDIAN SOILS

Katyal and Sharma (1991) reported that green schist contained the highest concentrations of Zn, Cu, Mn and Fe, while sandstone was the poorest source of these micronutrients. Among the sedimentary rocks, Zn content was highest in limestone. Shale contained considerably higher content of Fe, Mn and Cu as compared to limestone.

Soils developed on flood plain alluvium (Fluvents) and derived largely from silicious sandstone exhibited strikingly lower concentrations of Zn, Cu, Mn and Fe than Chromusterts developed on basaltic alluvium. In case of B, sedimentary rocks contain more B than igneous rocks (Gupta and Gupta, 1985).

Molybdenum (Mo) is the least abundant among micronutrients in the lithosphere (Mortvedt, 2000) and its deficiency largely occurs in acid soils and also in the soils formed from parent materials low in Mo, such as sedimentary, basalt and granite. Peaty, alkaline and poorly drained soils commonly have high Mo.

The total micronutrient content (zinc, iron, manganese, copper) is adequate in most of the Indian soils, but the micronutrients concentrations in soil solution is insufficient to meet the demand of growing crops (Singh, 1991).

Iron is the important micronutrient limiting plant growth. Due to the frequent change in its form (Fe^{2+} to Fe^{3+}), its availability to plants is greatly impaired with cropping sequences. Available (DTPA extractable) iron varies greatly from traces to 234 mg kg⁻¹ in Indian soils. Available zinc in Indian soils ranges from 0.08- 20.5 mg kg⁻¹ soil.

The available (DTPA extractable) copper content of the Indian soils ranges from traces to 32 mg kg⁻¹ soil. The highest copper deficiency was observed in soils of Kerala, followed by Uttar Pradesh. Available (DTPA extractable) manganese ranges from 0.6 to 164 mg kg⁻¹ soil in Indian soils with an average of 25 mg kg⁻¹. Hot water extractable boron ranged from traces to 8.2 mg kg⁻¹ in Indian soils (Gupta, 2005).

The total B contents in Indian soils varied from 7 to 630 mg kg⁻¹ of soil (Prasad *et al.*, 2014). The available (hot water soluble – HWS) B in Indian soils ranged from 0.75 to 8.0 mg B/kg (Das, 2000, Singh, 2001).

Total Zn content in the normal soils of world ranged between 10–300 ppm with an average of 50 mg kg⁻¹ (Mulligan *et al.*, 2001). In Indian soils total Zn ranges from a few ppm to about 1000 ppm. It ranges from 7 ppm in coarse textured alluvial soils (Entisols) to 284 ppm in fine textured vertisols (Ganjur *et al.*, 1973). Available Zn content in Indian soils ranges from 0.08 to 20.5 ppm.

2.3.1. Extent of micronutrient deficiencies in Indian soils

Smgh (2009) reported that Green revolution had significantly increased the food crop production in India, but continuous cultivation of high yielding crop varieties have led to depletion of native micronutrient soil fertility and now most of the soils are showing sign of fatigue for sustaining higher crop production. Besides, this hidden hunger of micronutrients is widely noticed leading to even entire failure of crops and reduced content of micronutrients in plant parts.

Takkar *et al.* (1989) reported that along with N, P and K the wide spread deficiencies of micronutrients are frequently seen in Indian soils. 43 per cent of Indian soils are deficient in Zn. High phosphate content of soils or high fertilization with phosphate may reduce the uptake of zinc and other nutrients (Dadhich and Somam, 2007, Kızılgoz and Sakın, 2010).

Deficiency of micronutrient has become a major constraint to the productivity, stability and sustainability of crops in many Indian soils and may further deteriorate due to global warming (Kumar *et al*, 2011) Till date, analysis of 3, 00,000 soil samples indicated that the average deficiency of Zn, Fe, Mn and Cu in Indian soils are 44, 15, 6 and 8 per cent respectively (Shukla and Behera, 2012) Similarly, the analysis of 50,000 soil samples showed B and Mo deficiencies of 33 and 13 per cent respectively B deficiency is reported more in acid soils than other parts of the country due to leaching of available B and continuous depletion of total soil reserved B

Zinc deficiency is prevalent in the soils having high pH, low organic matter and which are calcareous, sodic, sandy and limed acidic in nature (Rattan and Sharma, 2004) Much of the Zn associated with the solid phase is not available for plant uptake (Lake *et al*, 1984) It is estimated that 30 per cent of the world's cultivated soils are deficient in zinc (Suzuki *et al*, 2006)

B has emerged as an important micronutrient in Indian agriculture, next only to zinc in the context of the spread of its deficiency (Sathya *et al*, 2009) In India, boron deficiency was initially reported 2 per cent in the year 1980 (Katyal and Vlek, 1985), which has now increased to 52 per cent (Singh, 2012) Deficiencies of B in Indian soils ranged from 2 per cent in alluvial soils (Ustipsamments) of Gujarat, to 68 per cent in red soils (Calciorthhents, HaplustalFs) in Bihar, with a mean of 33 per cent for the whole country (Singh, 1999, Singh, 2006)

Mn deficiency is emerging very fast, particularly in wheat crops grown after rice in Haryana and Punjab due to continuous leaching of Mn from the surface layers of the coarse textured soils (Shukla *et al*, 2012)

Now, multi-nutrient deficiencies are emerging in many states of the country The Zn+Fe in swell-shrink soils, Zn+Mn or Zn+Fe+Mn in alluvial soils of Indo-Gangatic alluvial plains, Zn+Fe, Zn+B, Zn+Fe+B in highly calcareous soils of Bihar,

Saurashtra, Zn+B in acid leached alfisols, red and lateritic soils of India are leading to stagnation or a decline in productivity (Shukla and Behera, 2011)

Shukla *et al* (2012) confirmed that in acid soils of India, most of the soil samples indicated an adequate supply of Cu, Fe and Mn, low deficiencies of Zn (30 %) and higher deficiencies of B (46 %) and Mo (50 %)

Analysis of 20,000 plant samples for Zn, Cu, Fe and Mn indicated deficiencies of 44, 10, 6 and 4 per cent respectively, suggesting that the increasing multi-micronutrient deficiencies in soils and crops also affect animals and humans health along with crop productivity (Shukla *et al*, 2012)

Iron deficiency is a widespread agricultural problem that decreases plant growth and crop yields. Fe deficiency is a common nutritional disorder in many crop plants, causing chlorosis, poor yields and reduced nutritional quality. Increasing available Fe levels in major staple food crops is an important strategy to reduce Fe deficiency in people (Mori, 1999)

Manganese is a nutrient found in plant tissue at concentrations ranging from 10 to 500 mg kg⁻¹ or more. In most plants, it is deficient at less than 10 mg kg⁻¹ and toxic when the concentration exceeds about 300 mg kg⁻¹ (Sturgul, 2010)

The typical Cu content in plants ranged between 0.08–0.24 mmol g⁻¹ dry weight and Cu toxicity generally occurs when the plant tissue level exceeds 0.4 mmol g⁻¹ dry weight (Macnicol and Beckett, 1985)

2.4 STATUS OF MICRONUTRIENTS IN KERALA

Major soils of Kerala, derived from acid igneous rocks are deficient in Boron (SSO, 2007). Moreover, being highly mobile in the soil (Tisdale *et al*, 1985), leaching losses further aggravate B insufficiency in the high rainfall zones of Kerala, frequently leading to development of deficiency symptoms in crop plants.

Investigation on micronutrient status of cardamom- growing soils of Kerala conducted by Srinivasan *et al* (1993) indicated that available iron ranged from 14.6-65.8 ppm, available manganese from 1.3 to 44.8 ppm, available copper from 0.66 to 32.2 ppm, available zinc from 0.01 to 2.71 ppm, available boron from 0.05 to 3.7 ppm and available molybdenum from 0.01 to 11.1 ppm

Soils of Kerala are high in iron and manganese indicating high sesqui-oxide contents. Toxic levels of these elements are often observed in wetlands and deficiency of these elements has been very rarely reported, except from coastal sands and also the neutral to alkaline soils of Palakkad district. 31 per cent of Cu deficiency is observed in Kerala soils. In case of Zinc, deficiency was observed in Palakkad region. Widespread B deficiency has been observed in most of the districts with characteristic symptoms expressed in banana and cononut (DAO, 2013)

Srinivasan *et al* (1993) reported that 68 per cent of cardamom - growing areas of Kerala is deficient in zinc, 49 per cent deficient in boron, 28 per cent deficient in molybdenum and 9 per cent deficient in manganese

Evaluation of Soil fertility status of Kasargod district conducted by Suresh *et al* (2014) revealed that Kasarogod soils are 78 per cent deficient in B, 8 per cent deficient in Zn and 3 per cent deficient in Cu

Fe and Al toxicity is very wide spread disorder in acid sulphate soils of Kuttanad region and often leads to declining of crop yield of 50 to 70 per cent (Thampattil *et al*, 2005). The total Mn content of Kuttanad soils normally ranged between 28-350 mg kg⁻¹ (Rajendran and Aiyer, 1981, KAU, 1994) and total Fe content ranged from 2.75 to 7.72 per cent

2.5 FOLIAR APPLICATION OF MICRONUTRIENTS

Foliar spray of micronutrients is the most effective way to control nutritional deficiency problems than soil application (Torun *et al*, 2001). Foliar spray of

nutrients is preferred and gives quicker and better results than the soil application (Jamal *et al* , 2006) Generally, foliar application is very fast method for providing required elements in plants because nutrients are absorbing very quickly as compared with absorption that through plant roots (Hashemy *et al* , 1998)

Foliar application of nutrients has become an efficient way to improve yield and quality of crops (Roemheld and El-Fouly, 1999, Sarkar *et al* , 2007) In arid and semiarid regions, foliar application of nutrients is the most suitable option as compared with soil fertilization when the roots were not able to provide necessary nutrients to plants (Kaya *et al* , 2005, Kınacı and Gulmezoglu, 2007, Babaeian *et al* , 2011)

Other advantages are application of smaller quantity and thus, reducing toxicity arises from excessive accumulation of elements and preventing fixation of nutrients in soil (Malakouti and Tehrani, 1999) It may increase the efficacy of fertilizer use (Silberbush, 2002)

Application of fertilizers on leaves of growing plants with suitable concentrations is termed as foliar application Foliar feeding is a relatively new and controversial technique (Bernal *et al* , 2007 and Baloch *et al* , 2008) It is most economical way of fertilization, especially when sink competition for carbohydrates among plant organs take place, while nutrient uptake from the soil is restricted (Kannan, 1986 and Singh, M V , 2007)

In the early of 1950's the commercial production of mixed liquid fertilizers was started in United States and now it has been popularized in many countries (Wiley-VCH and Wiley-VCH, 2007)

For foliar fertilizers to be utilized by the plant for growth, the nutrient must first penetrate the leaf surface prior to entering the cytoplasm of a cell within the leaf

Penetration of foliar nutrients occurs through the cuticle the stomata, leaf hairs and other specialized epidermal cells (Fernandez and Eichert, 2009)

Foliar application of micronutrients resulted in quick absorption of nutrients by leaf epidermis and transported to other plant parts through xylem and phloem vessels (Hasslett *et al* , 2001, Nasiri *et al* , 2010)

Foliar feeding of nutrient promotes root absorption of the same nutrient or other nutrients by enhancing root growth and nutrients movement from terminal leaves to depth roots (El-Fouly and El-Sayed, 1997)

Recently foliar application of nutrients has become an important practice in crop production while soil application of fertilizers is the basic method (Alam *et al* , 2010)

Leaf feeding enhances overall nutrient level in the plant and sugar production during times of stress and it also increases biochemical activities in the leaf by increasing chlorophyll a, b and carotenoids, which presumably favour the photosynthesis Similarly, Shitole and Dhupal (2012) found that foliar application of micro-nutrients increased the primary metabolites like photosynthetic pigments and organic constituents of *Cassia angustifolia*

Foliar feeding was more efficient than soil application in such a way, N- 6 times more efficient, B- 4 times more efficient, Mn- 30 times more efficient, Zn- 20 times more efficient, P- 20 times more efficient and Mo- 14 times more efficient than soil application (Dixon, 2003) Likewise, Liew (1988) advocated that foliar application of micro-nutrients is 6-20 times more efficient than soil application, depending upon various soil types

Foliar spray of zinc, boron and copper has been reported to be equally or even more effective way to overcome micronutrients deficiency in subsoil as compared to soil application (Ali *et al* , 2009, Hussain *et al* , 2012)

Foliar application micronutrient leads to increase in grain yield components and protein percentage of wheat, maize, rice, barley and sorghum (Boorboori *et al* , 2012)

Foliar application of water soluble iron containing compounds may be applied in alkaline soil conditions to meet out the iron requirement of crops Foliar spray of Ferrous sulphate (0.5 per cent) solution is required repeatedly if chlorosis symptoms persist (Patricia *et al* , 2010)

Moosavi and Ronaghi (2011) found that both soil and foliar application of Fe and Mn notably increased the shoot concentration and uptake in soybean plant However, foliar feeding was found to be efficient than soil application

Copper has been used as a fungicide even before it was recognized as an essential plant nutrient (Graham and Webb, 1991)

Therefore, foliar feeding of nutrients has become an established procedure in crop production to increase yield and quality of crop products (Roemheld and El-Fouly, 1999) and it also minimizes environmental pollution and improves nutrient utilization through reducing the amounts of fertilizers added to the soil (Abou-El-nour, 2002)

2.6 MICRONUTRIENT MIXTURE ON GROWTH AND YIELD OF CROPS

Muthukrishnan and Velu (2011) developed and evaluated a micronutrient formulation for sugarcane and concluded that 92.5 kg MN formulation II as enriched farm yard manure containing 50 kg FeSO₄, 5kg MnSO₄, 30 kg ZnSO₄, 2.5 kg CuSO₄ and 5 kg of borax may be recommended for sugarcane for maximizing yield with higher B:C ratio Increased cane yield with higher levels of micronutrient mixture maybe attributed to the complementary and supplementary effect of fertilizers and FYM

A multi-micronutrient mixture having a composition of Zn (9.5 %) + B (2.6 %) + Cu (1.2%) + Mg (2.4 %) + N (0.46 %) was developed and found that 0.5 per cent of multi-micronutrient mixture was significantly improved the growth and yield of okra variety Varsha Uphar (Mini, 2015)

Micronutrients play a vital role in enhancing the productivity of crops. Optimum fertilization and manuring are prime need for the growth and yield of banana (Simmonds, 1966)

El-Magid *et al* (2000) found that application of NPK with foliar spraying of various micronutrient mixture containing Fe, Mn, Cu, Zn, B, Mg, S and Mo (in the brand names, Satrite SF, Borocol BSF-12 and Micnef MS-12) increased the grain yield of rice

Shueadshen (1991) reported that foliar application of 0.1 per cent B, Fe, Cu, Co, Zn and Mn at tillering stage has increased number of spikelets per panicle and grain yield and decreased the spikelet sterility

Durairaj (1993) reported that combined foliar application of 20ppm humic acid and 25ppm ZnSO₄ gave highest yield of paddy

Modaihsh (1997) found that combined application of micronutrients either in chelated or non-chelated forms resulted in higher biological and grain yields in wheat than the individual application of micronutrients

El-Magid *et al* (2000) reported that foliar spray of micronutrients containing Fe, Cu, Zn and Mn increased the grain and straw yields of wheat

Field experiment on the effect of various micronutrients such as Zn, Cu, Fe, Mn, B and a commercial micronutrient mixture (Brand name- Zarzameen) resulted in maximum grain yield, straw yield and dry matter of wheat (Asad and Rafique, 2002)

According to Patel *et al* (2008), foliar application of 1 per cent micronutrient mixture (containing Fe - 4 %, Mn -1 %, Zn - 6 %, Cu - 0.5 % and B - 0.5 %) at 30, 40, 50 days after sowing recorded highest plant height, number of tillers per plant, shoot weight, grain weight and increased uptake of micronutrients

Foliar spray of micronutrients (containing Fe - 1 %, Mn - 2 %, Zn - 2 %, Cu - 1 %, B - 1 %) on wheat increased the plants height, grams per spike, biological yield, harvest index, 1000-grain weight, straw and grain yield (Khan *et al* , 2010)

Combined foliar spraying of 0.5 per cent FeSO_4 and 0.5 per cent ZnSO_4 is most effective and increases seed yield by 43.1 per cent in cowpea cultivated oxisols of Kerala (Anitha *et al* , 2005)

Katkar (2005) concluded that 3 sprays of MgSO_4 (1 %) + ZnSO_4 (0.5 %) at squaring, flowering and boll development stage are recommended to obtain highest yield of cotton

According to Namdco (1992), combined application of recommended NPK rate ($120\ 60\ 60\ \text{kg ha}^{-1}$) + Foliar spray of one per cent Micnef MS-16 (consisting of Zn, Cu, Fe, Mn, Mo and B) at 30, 65 and 90 days after sowing resulted in highest seed cotton yield of $8.12\ \text{t ha}^{-1}$

Foliar fertilization of micronutrient mixture (Brand name- Agripower N-180,000 ppm, K- 1000 ppm, Mg- 100 ppm, Zn- 100 ppm, B-100 ppm, Fe- 1000 ppm, Cu- 20 ppm and Mn- 100 ppm) improved the growth and yield parameters of okra plants such as days to flowering, plant height, number of branches plant^{-1} , number of fruits plant^{-1} and fruit length (Abbasi *et al* , 2010)

Patel *et al* (2009) confirmed that green fruit yield of okra significantly improved with foliar application of multi-micronutrient mixture

According to Datir *et al* (2010), application of 2 per cent organically chelated micronutrient mixture containing Zn, Cu, B and Fe on okra increases plant height, leaf area and yield per plant

Suryanarayana and Rao (1981) investigated that foliar application of Fe with Zn, Cu, Mn, Mg, B and Mo in a chelated form (brand name- Agromm) resulted in maximum growth traits of okra

Application of Cu - 20 ppm, Mn - 100 ppm and Zn - 50 to 100 ppm significantly increased the yield parameters like fruit set percentage, number of seeds per pod and total dry bean yield of *Phaseolus vulgaris* cv Giza-3 (Gabal *et al* , 1985)

Yousefi and Zandi (2012) observed that foliar spray of Zn + Mn resulted in highest seed number, fruit yield, seed yield and oil content of pumpkin (*Cucurbita pepo* L.)

Yadav *et al* (2009a) reported with three times foliar application of multiplex (0.5 %) gave highest marketable head yield of cabbage with maximum net returns and maximum vegetative growth (number of leaves per plant, more head diameter, higher individual head weight as well as foliage yield)

Kanuja *et al* (2006) found that maximum plant spread, number of non-wrapper leaves, head diameter, head weight and head yield of cabbage var 'Golden Acre' were recorded with foliar application of micronutrient mixture (Zn, Fe, B, Mn, Cu and Mo) @ 100 ppm

Mishra *et al* (2012) reported that maximum number of fruits per plant and yield of tomato cv Utkal Urbasi was obtained with foliar spray of micronutrient mixture containing Zn-100 ppm, B- 100 ppm, Fe- 100 ppm, Mn- 100 ppm, Cu- 100 ppm and Mo- 50 ppm

Naruka *et al* (2000) found that foliar application of Zn and Mo at 0.2, 0.4 and 0.6 % and 30, 60 and 90 ppm respectively significantly increased the plant height, number of fruits, fruit diameter and fruit yield of okra cv Pusa sawani

Studies conducted by Sivaiah *et al* (2013) in tomato (*Lycopersicon esculentum* Mill) reported that foliar application of micronutrients either alone or in combination, enhances plant growth characteristics viz plant height, number of primary branches and compound leaves, fruit weight, seed yield and seed weight

Bhatt and Srivastava (2005) claimed that fresh and dry matter weight of tomato fruits, uptake of all the macro and micronutrients has remarkably influenced by the foliar spray of micronutrient mixture (Zn, Fe, B, Cu and Mn as 100 ppm and Mo - 50ppm)

Foliar application of micronutrients recorded higher fruit yields of brinjal than soil application (Selvi and Thiageshwari, 2002)

Al-Jobori and Al-Hadithy (2014) reported that application of full concentration of micronutrient mixture containing 330 g ZnSO₄ + 330 g MnSO₄ + 150 g FeSO₄ + 80 g CuSO₄ in 1 litre significantly increased the number of tuber / plant, mean tuber weight, total yield and dry matter percentage of potato (*Solanum tuberosum*)

According to Pal *et al* (2004), foliar application of Boron and Zinc @ 1000 ppm has significantly increases fruit base diameter, fruit weight and pericarp thickness as well as foliar application at 2000 ppm Boron increase quality traits like TSS, ascorbic acid, acidity and carotene content of bell pepper (*Capsicum annuum* L)

Singh and Rajput (1976) found that various combinations of foliar application of Zn (0.1, 0.2 and 0.4 %), Fe (0.1, 0.2 and 0.4 %) and B (0.1, 0.2 and 0.4 %) has

increased the length of terminal shoot, plant height, number of leaves and leaf area per shoot of mango tree

Studies conducted by Ghosh *et al* (1995) revealed that highest yield was obtained in 0.75 per cent brand name Multiplex- Trace elements formulation I with 62.7 kg/plant as compared to control (37.0 kg/plant) and fruit weight was highest in 0.70 per cent brand name Tracel- Trace elements formulation II in Mango cv Himsagar

According to Bhatt *et al* (2012), foliar spray of 0.5 per cent borax on mango trees showed highest fruit yield, fruit weight and fruit volume

Gurjar *et al* (2015) revealed that foliar application of 1 per cent $ZnSO_4$, 1 per cent $FeSO_4$ and 0.5 per cent borax on Alphonso mango (*Mangifera indica* L.) increased number of fruits per tree, fruit weight, yield tree⁻¹ and decreased the fruit drop

Modi *et al* (2012) revealed that foliar application of $ZnSO_4$ 0.5 per cent and borax 0.3 per cent at 30, 60 and 90 days after transplanting on papaya cv Madhu Bindu exerted great influence on plant height, stem girth, number of leaves, earlier initiation of flower bud, minimum days taken from fruit setting to first harvest, weight of fruit, number of fruit and yield of fruits per plant with maximum yield per hectare

Foliar sprays of Zn 0.5 per cent + B 0.1 per cent at 4th, 8th, 12th and 16th months after planting improved the total number of fruits per tree, fruit characters, fruit and latex yield of papaya (Kavitha *et al*, 2000a)

The monthly spray of $FeSO_4$, $ZnSO_4$ and borax @ 0.1 per cent with or without combination were effective in increasing the plant height and girth of papaya as compared to control (Veena and Lavania, 1998)

Singaram and Prabu (2001) reported that foliar application of ZnSO_4 @ 0.5 per cent + borax @ 0.2 per cent resulted in highest shoot length, number of internodes per shoot and number of leaves per shoot of grapes

Foliar application of Zinc (0.6 %), Copper (0.3 %) and Boron (0.3 %) enhanced the growth and vigour of Litchi plant (Babu and Singh, 2002)

Afria *et al* (1999) reported that foliar spray of ZnSO_4 + FeSO_4 + Borax in pomegranate resulted in maximum number of fruits plant⁻¹, fruit weight and yield plant⁻¹

Bhambota *et al* (1962) found that application of Zinc (0.6 %) + Iron (0.4 %) significantly increased the number of fruits, mean weight of fruit, diameter and volume of each fruits of citrus

Foliar spraying of Zn and Fe at 0.5 per cent (4 sprays) considerably increased the chlorophyll content of acidlime (Patel and Patel, 1985)

Muthukrishnan and Velu (2011) reported that micronutrient mixture formulation enriched with farmyard manure in sugarcane will largely helps to manage the micronutrient deficiency problems effectively and it clearly reveals that the positive improvement of growth and development due to conjoint use of NPK with organic and micronutrient fertilization

Manimaran *et al* (2009) reported that application recommended dose of fertilizer+ Acetobacter @ 10 kg ha⁻¹+ foliar spraying of 1 per cent micronutrient mixture at 45 and 75 DAR resulted in maximizing single cane weight, cane yield and sugar yield

Foliar spray of micronutrient mixture of EDTA chelated Cu, Zn, Mn and Fe on capsicums cv Vinedale as 5 sprays at 15 days intervals resulted in increased in yield and fruit colour (Navrot and Levin, 1976)

Senthambizhselvi (2000) found that combined application of zinc sulphate - 4 g and ferrous sulphate - 8 g per plant through soil and 0.5 per cent of zinc sulphate and 1 per cent of ferrous sulphate through foliage exerted significant influence on plant height, number of shoots, plant spread and leaf area in *Jasminum sambac*

Katiyar *et al* (2012) reported that foliar application of Zn at 0.5 per cent was effective to increase leaf length, number of leaves, size of spike, thickness of spike and floret of gladiolus plants

Foliar application of micronutrients resulted in increased number of branches, number of leaves per plant, number of blind shoots, vase life of flower, number of flower per plants in rose under polyhouse condition (Jagtap *et al* , 2012)

Balakrishnan (2005) claimed that foliar application of zinc sulphate - 0.5 per cent + ferrous sulphate - 0.5 per cent sprayed at 30 and 45 days after planting in marigold significantly increased stem girth, plant spread, number of branches per plant and dry matter production

Foliar feeding of plants with ferrous sulphate + zinc sulphate + manganese sulphate @ 0.2 per cent each resulted in increasing flower yield per plant, flower diameter and plant height in gerbera (Balakrishnan, 2005)

Ahmad *et al* (2010) revealed that combined application B and Zn in rose plants resulted in increasing plant height, number of leaves branch¹, leaf area, number of flowers plant¹, flower stalk length and leaf Zn contents

2.6.1. Effect of micronutrient mixture on growth and yield of banana

Foliar application of micronutrient mixture (Mg, Fe, Zn, Cu and Mn) resulted in highest bunch weight, number of hands and fingers per bunch, higher numbers of green leaves per plant and decreased bunch shooting period of Hindi banana (*M cavendishi*) (Abdel-Kader *et al* , 1992)

Das and Mohan (1993) reported that the maximum plant height, leaf area and number of functional leaves were obtained with a combination of B, Zn, Cu and Mn applied at 3 and 5 months after planting of banana cvs 'Chenichampa' (AAB), 'Jahaji' (AAA) and 'Barjahaji' (AAA)

Ghanta and Mitra (1993) found that the combined application of Zn (0.3%), Cu (0.1%) and B (0.2%) showed the best response in plant growth at flowering in terms of height, girth of pseudostem and number of leaves per plant and also it increased the yield parameters such as number of hands per bunch, number of fingers per bunch, bunch weight and yield per hectare in banana cv Giant Governor

Suresh and Savithri (2001) observed that soil application of N, P, K and foliar spray of nutrients (1% DAP + 1% MOP + 0.5% ZnSO₄ + 0.3% CuSO₄ + 0.2% Borax) in addition to liming caused substantial increase in bunch yield of banana

Foliar application of micronutrient mixture (ZnSO₄ - 0.5%, FeSO₄ - 0.2%, CuSO₄ - 0.2% and H₃BO₃ - 0.1%) at 3, 5 and 7 months after planting of banana increased bunch weight and TSS in ripe fruits (Kumar and Jayakumar, 2001)

Phenotypic correlation analysis of 13 biometric characters of banana cv Nendran showed that bunch length, number of hands and number of fingers had highly significant correlation with yield. Plant height exerted the highest direct effect on bunch yield while the pseudostem girth, leaf area and leaf area duration contributed indirectly through their effect on plant height (Kumar, 2001)

Spraying of Zn (1%) + B (0.5%) produced the highest plant height, number of functional leaves, petiole length, number of suckers, number of hands per bunch, number of fingers per bunch, hand distance, hand weight, bunch length and bunch weight of dwarf cavendish banana (*Musa*, AAA, sub-group-cavendish, cv Giant Governor) (Mandal *et al*, 2002)

Studies conducted by Yadlod and Kadam (2008a) on the effect micronutrients (*Musa sp*) cv Grand Name revealed that early maturity (98.3 days) and minimum number of days required for ripening were observed with foliar spraying of micronutrients mixture 1 per cent (two sprays). Maximum length (21.9 cm) and girth (15.5 cm) of fingers, maximum weight of ripe finger (137.5 g) were also recorded.

Yadlod and Kadam (2008b) revealed that foliar spraying of micronutrients mixture 1 per cent (two sprays) resulted in maximum weight of bunch (23.8 kg), finger length (23.0 cm), finger girth (17.0 cm) and weight of mature finger (185.6 gm) and minimum number of days required for ripening of banana (*Musa sp*) cv Shrimanti (10.7 days).

Yadlod and Kadam (2008c) claimed that Micronutrients mixture 1 per cent two spray in banana (*Musa sp*) cv Ardhapuri enhanced the plant height (179.2 cm), pseudostem girth (67.3 cm) and number of leaves (14.2). Early maturity (106.0 days), Maximum finger length (22.0 cm) and girth (16.2 cm) and early ripening (9.8 days after harvesting) were also recorded with micronutrient mixture 1 per cent. Maximum increase in weight of mature finger was found in two sprays of 1 per cent micronutrients mixture (178.3 g) followed by micronutrients mixture 1 per cent one spray (171.3 g).

Jeyabaskaran and Pandey (2008) reported that under high soil pH condition micronutrient such as Fe, Zn and B applied through foliar spray which resulted in higher pseudostem height, girth, total leaf area and bunch weight, number of fingers per bunch of Karpuravalli banana.

Yadav *et al* (2010) reported that the higher level of chelated zinc (Zn-EDTA) with MnSO₄, CuSO₄, and Borox induced early and vigorous vegetative growth in terms of height and girth of the pseudostem, number of leaves and total leaf area consequently and also induces early inflorescence emergence in the plant and reduced the crop duration. Ultimately the treatment with micronutrients produced the plant

with higher bunch weight, number of hands and fingers per bunch, length and girth of finger and yield per hectare. It also produced favorable effect on fruit quality in terms of TSS, total sugars and reducing sugars, sugar/acid ratio and acidity percent.

Patel *et al* (2010) observed that the foliar application of $ZnSO_4$ (0.5 %) + $FeSO_4$ (0.5 %) of banana cv Basrai at 3rd, 5th and 7th months after planting resulted in highest bunch weight, bunch length, bunch girth, number of hands per bunch and fruit yield.

According to Pathak *et al* (2011), the combined application of Fe (0.5 %) and Zn (0.5 %) on banana cv Martaman (AAB, Silk) showed best response to plant height, basal girth of pseudostem, number of leaves, bunch weight, bunch length, bunch breadth, hands per bunch, fingers per bunch, days to shooting days to harvest, finger length, finger breadth, crop duration and days to ripening.

Integrated Nutrient Management involving supply of 50 per cent of nutrients by spray on leaf and bunch (0.5 % Urea, 0.5 % SOP, 0.2 % $ZnSO_4$ and 0.1 % Boric Acid) four sprays on leaf from 5th months to 8th months improve quality in banana (IIHR, 2013).

Anjali *et al* (2013) found that foliar spraying of micronutrient mixture containing $ZnSO_4$ (0.5 %) + $FeSO_4$ (0.5 %) + $CuSO_4$ (0.2 %) + H_3BO_3 (0.1 %) increased the growth and yield parameters of banana cv Grand Name such as plant height, pseudostem girth, number of leaves per plant, leaf area, bunch weight, number of hands per bunch, number of fingers per bunch and yield ($t\ ha^{-1}$).

2.7 MICRONUTRIENT MIXTURE ON QUALITY OF CROPS

Application of Zn, Fe and Mn at 0.1 and 0.2 per cent resulted in considerable improvement of TSS, sugars and ascorbic acid contents in mango fruit at harvest as well as on ripening as compared to control (Dutta and Dhua, 2002).

Foliar spraying of Zinc @ 0.1 per cent alone or in combination with B @ 0.4 per cent and Cu @ 0.05 per cent increases TSS content of mango fruit cv Langra (Panwar and Singh, 2007)

Foliar spraying of Boric acid 0.02 per cent on mango cv Alphonso significantly increased the total sugars, reducing sugars, non-reducing sugars, TSS, ascorbic acid, carotenoids and sugar acid ratio (Sankar *et al*, 2013)

Kanuja *et al* (2006) found that maximum shelf life of cabbage var 'Golden Acre' was recorded with foliar application of micronutrient mixture (Zn, Fe, B, Mn, Cu and Mo) @ 100 ppm

Foliar spray of Zn, B, and Mo at the rate of 0.2 per cent ZnSO₄, 0.4 per cent boric acid and 0.05 per cent ammonium molybdate, singly and in various combination on Guava revealed that maximum TSS, ascorbic acid, reducing and non-reducing sugars were obtained with combination of Zn, B and Mo (Singh and Chonkar, 1983)

Foliar spray of Zn @ 0.5 per cent + B @ 0.1 per cent at 4th, 8th, 12th and 16th months after planting in papaya increased total sugars, reducing sugars, non-reducing sugars, ascorbic acid and sugar acid ratio, in association with bio-chemical traits (Kavitha *et al*, 2000b)

Prabu and Singaram (2001) reported that the application of ZnSO₄ at 0.5 per cent + borax at 0.2 per cent through foliage increased the TSS, reducing sugars, total sugars and sugar acid ratio and reduced acidity of grapes

Sarkar *et al* (1984) studied that the foliar application of Zn, Cu, B and K on litchi increased the pulp weight, total sugars and TSS over control

Foliar spray of various concentrations of $ZnSO_4$ and borax increased the pulp weight, pulp peel ratio, TSS and sugar acid ratio and significantly decreased the acidity of litchi (Rani and Brahmachari, 2001)

Studies conducted by Dixit *et al* (1977) revealed that the sprays of $ZnSO_4$ and $FeSO_4$ on Kinnow mandarin, improved the juice content, TSS, total and reducing sugar, sugar acid ratio and ascorbic acid content

Haque *et al* (2000) found the maximum total sugar content, reducing sugar content, non-reducing sugar content and ascorbic acid content in fruit juice with spraying of $ZnSO_4$ @ 0.5 per cent in Mandarin orange

Modi *et al* (2012) revealed that foliar application of $ZnSO_4$ 0.5 per cent and borax 0.3 per cent at 30, 60 and 90 days after transplanting on papaya cv Madhu Bindu exerted great influence on ascorbic acid, total soluble solids, reducing sugar, non-reducing sugar and total sugar

Mishra *et al* (2012) reported that maximum TSS and Lycopene content of tomato cv Utkal Urbasi was obtained with foliar application of micronutrient mixture containing Zn-100 ppm, B- 100 ppm, Fe- 100 ppm, Mn- 100 ppm, Cu- 100 ppm and Mo- 50 ppm

2.7.1. Effect of micronutrient mixture on fruit quality of banana

Ghanta and Mitra (1993) observed that TSS, total sugars, reducing sugars, Pulp peel ratio, sugar acid and ascorbic acid content of banana cv Giant Governor were highest with foliar application of 0.3 per cent Zn + 0.1 per cent Cu + 0.2 per cent B at 3 and 5 months after planting

Das (1995) reported that the maximum TSS and total sugars were obtained with foliar spraying of B + Zn + Cu at 3 and 5 months after planting of banana cvs 'Chemchampa' (AAB), 'Jahajr' (AAA) and 'Barjahajr' (AAA)

Pertin and Das (1998) revealed that in addition to NPK fertilizers, Zn, Fe, Cu and B were applied separately at 0.5, 0.2, 0.2 and 0.1 per cent, respectively, or in a combination increased the plant growth of banana cv Barjahaj.

Foliar application of ZnSO₄ (0.5 %) + FeSO₄ (0.2 %) + CuSO₄ (0.2 %) + H₃BO₃ (0.1 %) at 3, 5 and 7 months after planting increased the pseudostem girth, number of leaves and total chlorophyll content, bunch weight and total soluble solid content of banana (*Musa AAA*) cv Robusta (Kumar and Jayakumar, 2001)

Suresh and Savithri (2001) found that the soil application of N, P and K and foliar spray on nutrients (1 % DAP + 1 % MDP + 0.05 % ZnSO₄ + 0.2 % CuSO₄ + 0.2 % Borax) in addition to liming had increased the TSS, sugar acid ratio and decreased the titrable acidity of banana

Jeyabaskaran and Pandey (2008) reported that under high soil pH condition micronutrient such as Fe, Zn and B applied through foliar spray which resulted in maximum total soluble solids, acidity and TSS/acid ratio of Karpuravalli banana

Micronutrient mixture 1 per cent at two times spraying in banana cv Grand Name significantly increases the total sugars, reducing sugars and non-reducing sugar content and minimizes per cent loss in weight during ripening (Yadlov and Kadam, 2008a)

Patel *et al* (2010) observed that the foliar application of ZnSO₄ (0.5 %) + FeSO₄ (0.5 %) of banana cv Basrai at 3rd, 5th and 7th months after planting resulted in highest ascorbic acid content, reducing sugars and total sugars

According to Pathak *et al* (2011), the combined application of Fe (0.5 %) and Zn (0.5 %) on banana cv Martaman (AAB, Silk) showed maximum sugar acid ratio (47.7), non-reducing sugar (10.0 %) also showed considerable improvement on total soluble solids (25.5 °Brix) and total sugar (17.24 %) content of pulp

Maeda *et al* (2011) confirmed that foliar spraying of zinc sulfate + citric acid + EDTA + iron sulfate + manganese sulfate + magnesium sulfate (9.4 % S, 5 % Zn, 1 % Fe, 1 % Mn, 1 % Mg, 1.5 % citric acid, and 4.3 % EDTA) has profound to increase TSS, Titratable acidity of pineapple

Yadav *et al* (2011) recorded maximum TSS and other quality parameters of banana cv Grand Naine with Recommended dose of fertilizers (200+90+200 NPK g/plant) + 40 g Zn EDTA + 20 g MnSO₄ + 5 g CuSO₄ + 10 g Borax/plant

According to Bhatt *et al* (2012), foliar spray of 0.5 per cent borax on mango trees showed highest TSS, reducing sugar, non reducing sugar and ascorbic acid content

Bauri *et al* (2014) reported that foliar spraying of Borax (0.1 %) at 5 and 20 days after last hand opening showed maximum total soluble solids (26.7 °Brix), total sugar (14.6 %) and sugar acid ratio (45.5) of banana cv Martaman (*Musa* AAB, Silk)

Ningavva *et al* (2014) found that the maximum pulp weight (147.2 g), peel weight (42.9 g), pulp to peel ratio (3.4), TSS (28.5 °Brix), reducing sugar (16.0 %), non reducing sugar (2.9 %), total sugars (19.0 %), Titrable acidity (0.3 %) and Sugar to acid ratio (65.4) of Ratoon Banana cv Grand Naine were recorded with 100 per cent RDF (100-108-200g NPK/plant) + foliar spray of ZnSO₄ (0.5 %) + boron (0.2 %) with double suckers per hill

Paul and Nair (2015) reported that foliar spray of ZnSO₄ 1.0 per cent + FeSO₄ 0.3 per cent + CuSO₄ 0.2 per cent + H₃BO₃ 0.2 per cent + (NH₄)₂MoO₄ 0.03 per cent recorded highest total sugar (19.4 %) and reducing sugar (16.9 %) and minimum non reducing sugar (2.5 per cent) of banana (*Musa* AAB) Nendran

2.8 EFFECT OF MICRONUTRIENTS ON NUTRIENTS CONTENT IN LEAF

Shueadshen (1991) reported that foliar application of 0.1 per cent B, Fe, Cu, Co, Zn and Mn at tillering stage increased N and P uptake over control in rice. Gupta and Gupta (1985) revealed that foliar application of chelated micronutrient mixture enhanced the uptake of Zn, Fe, Mn, Cu, Ca and Mg content in rice.

According to Patel *et al* (2008), foliar spray of multi-micronutrient mixture (Fe- 6 %, Mn- 1 %, Zn- 4 %, Cu- 0.3 % and B- 0.5 %) resulted in maximum micronutrient uptake by wheat grain and straw.

Spraying with different micronutrients (Zn 0.3 %, Cu 0.1 %, B 0.2 % and Mn 0.05 %) with or without combinations on banana cv Giant Governor significantly increased the leaf nitrogen, phosphorus and potassium content before flowering compared with control (Ghanta and Mitra, 1993).

According to Ray *et al* (2006), leaf content of 2.8, 0.52, 3.8 per cent NPK at shooting was a good indicator for satisfactory productivity in Robusta banana.

Foliar application of ZnSO₄ (0.5 %) + FeSO₄ (0.2 %) + CuSO₄ (0.2 %) + H₃BO₃ (0.1 %) at 3, 5 and 7 months after planting increased the leaf micronutrient status of banana (*Musa AAA*) cv Robusta (Kumar and Jayakumar, 2001).

Jeyabaskaran and Pandey (2008) reported that under high soil pH condition micronutrient such as Fe, Zn and B applied through foliar spray which resulted in higher K, Zn, B, Fe and Mn content of Karpuravalli banana leaves.

Foliar spray of micronutrients (Zn as ZnSO₄ at 0.3 %, Boron as boric acid 0.1 %, Cu at 0.1 % with 1 % SOP and 1 % Urea) four times at monthly intervals resulted in improving leaf Zn, B and Cu nutrient status of banana cv Ney Poovan (Muthaia and Raja, 2009).

Yadav *et al* (2010) recorded highest Zn content in banana cv Grand Name leaf with RDF (200+90+200 NPK g/plant) + 40 g Zn EDTA + 20 g MnSO₄ + 5 g CuSO₄ + 10 g Borax/plant While maximum Fe content in leaf noted from RDF + 25 g FeSO₄ + 2 g MnSO₄ + 5 g CuSO₄ + 10 g Borax and RDF + 25 g Fe EDTA + 20 g MnSO₄ + 5 g CuSO₄ + 10 g Borax/plant, respectively (Yadav *et al* , 2009b)

Dutta and Dhua (2002) reported that the leaf nutrient status of mango increased with application of Zn (0.1 and 0.2 %), Mn (0.1 and 0.2 %) and Fe (0.1 and 0.2 %) Application of Zn improved the leaf N and Zn content while Fe improved P and Fe and Mn improved the Mn and K contents

Foliar spraying of Boric acid 0.02 per cent on mango cv Alphonso significantly increased the N, P, K and B content of leaves (Sankar *et al* , 2013)

Aggarwal *et al* (1975) observed that different combination of ZnSO₄ (0.5 %), CuSO₄ (0.25 %) and FeSO₄ (0.25 %) increased the Zn, Cu and Fe contents in leaf tissues of Thompson seedless grapes over the control

Durgadevi *et al* (1997) reported that the application of ZnSO₄, FeSO₄ and MnSO₄ as foliar spray or soil application or combination of both, significantly increased the N, P, K, Ca and Mg contents in the Sathgudi orange leaves

Patel (2006) revealed that total iron content in leaves increased due to foliar application of iron which recovered iron chlorosis in citrus

Ram and Bose (2000) found that the foliar spray of MnSO₄ (2 %), CuSO₄ (0.4 %), ZnSO₄ (0.5 %), boric acid (0.1 %) and FeSO₄ (0.25 %) either singly or in various combinations significantly increased the N, P and K content of the Mandarin orange leaf

Afria *et al* (1999) reported that the foliar application of FeSO_4 (0.4 %), ZnSO_4 (0.25 %) and borax (0.2 %) on pomegranate increased the Fe, Zn and B content in leaves individually or in combination

Dalal *et al* (2011) found that foliar spraying of zinc sulphate and borax significantly increased the Zn and Mn uptake by ber leaves

Foliar spray of FeSO_4 (0.5 %) + ZnSO_4 (0.5 %) + Boric acid (0.2 %) resulted in maximum uptake of N and B in okra cv Phule Utkarsha (Satpute *et al*, 2013)

2.9 MICRONUTRIENTS ON PLANT DISEASE CONTROL

All the essential nutrients can affect the disease severity (Huber and Graham, 1999) and nutrients availability is the best way to control plant diseases in an integrated pest management system (Graham and Webb, 1991)

Micronutrient disorders of Zn, Mn, B, Cu and Fe are widespread in India and correction of these nutritional disorders resulted in resistance to plant diseases (Agrios, 2005). The effect of micronutrients on reducing severity of diseases can be attributed with physiology and biochemistry of plants (Marschner, 1995)

Micronutrient concentrations in plants are important in host ability to resist or tolerate infectious pathogens. Plant nutrition is an important component of disease control (Huber and Wilhelm, 1988)

Boron sufficiency in plants reduces the incidence and severity of diseases, while B deficiency enhances them (Gupta, 1993, Graham and Webb, 1991). Manganese concentration in host tissue commonly decreases as the incidence of disease increases. The severity of parasites on plants is influenced by many micronutrients (Engelhard, 1990, Graham and Webb, 1991, Huber, 1980). The lack of Zn, B, Mn, Mo, Ni, Cu and Fe in plant tissue can enhance various diseases on

plants (Engelhard, 1990; Graham and Webb, 1991, Huber, 1980, Fageria *et al* , 1997, Baligar *et al* , 1998)

Prophylactic treatment of micronutrient solutions as a foliar spray on the upper surface leaves of cucumber plants significantly inhibited powdery mildew development (Reuveni *et al* , 1997)

Simoglou and Dordas (2006) revealed that manganese fertilization decreased the pathogenic disease incidence such as powdery mildew and take-all of wheat. Soil application of Mn reduced common scab of potato (Kenath and Loria, 1996), *Fusarium* wilt of cotton and *Sclerotinia sclerotiorum* in squash (Graham and webb, 1991, Agrios, 2005)

Zinc nutrition is associated with important plant defense pathways against many fungal and bacterial pathogens. Grewal *et al* (1996) reported that root rot and *Fusarium graminearum* (Schwabe) diseases of wheat controlled with soil application of Zn

Proper Fe nutrition not only boosts plant vigour and health, it indirectly affects disease in the rhizosphere where its availability may limit the growth of pathogens. Iron reduced the disease severity of rust and smut of wheat, *Colletotrichum musae* of banana (Graham and Webb, 1991, Graham, 1983)

Graham (1983) observed that foliar application of Fe increased the disease resistance of apple and pear to *Sphaeropsis malorum* and cabbage to *Olipidium brassicae*. Rhizosphere microorganisms could be able to synthesize siderophores which lowers Fe levels for harmful microorganisms in soil. These iron chelators resulted in suppressing chlamydospores of *Fusarium oxysporum f.sp. cucumerinum*, crown-gall (*Agrobacterium tumefaciens*) and take-all diseases of wheat, soft rot of potato (*Erwinia caratovora*)

Copper fungicides are widely used for managing several plant diseases which having broad-spectrum activity, acting upon bacteria as well as fungi. Bacterial pustules successfully controlled by two sprays at 45 and 55 days after planting with a mixture of streptomycin (150 g/ha) + copper sulphate (1kg/ha) (Kanniyar and Prasad, 1979). Soil application of Cu reduced downy mildew (*Plasmopara viticola*) of grapes (Evans *et al* , 2007)

Patil (1981) reported that Molybdenum reduced Ascochyta blight (*Ascochyta* spp) in beans and peas and late blight (*Phytophthora infestans*) of potato. Symptoms of verticillium wilt of tomato reduced with Mo application (Kuti *et al* , 1989)

Foliar spray of micronutrients (Zn as ZnSO₄ at 0.3 %, Boron as boric acid 0.1 %, Cu at 0.1 % with 1 % SOP and 1 % Urea) four times at monthly intervals resulted in improving resistance to Panama wilt of banana cv Ney Poovan. The number of affected plants in control plot is 32 per cent whereas in micronutrients sprayed plot 5 per cent only (Muthaia and Raja, 2009)

According to Sanjeev and Eswaran (2008), the mycelial growth of *Fusarium oxysporum f. sp. cubense* causing Banana fusarium wilt was inhibited by *Trichoderma viride* applied with combination with Borax and Zinc sulphate amended medium and also found that Borax at higher concentration maximizes sporulation capacity

Reuveni *et al* (1998) reported that foliar spray of H₃BO₃, CuSO₄, MnCl₂ resulted in reduction of powdery mildew disease severity in cucumber

McMillan *et al* (2000) found that KeyPlex 350 (micronutrient preparation) treatment reduced the incidence of Sigatoka leaf spot of banana (*Musa acumata*) cv Dwarf Cavendish as compared to the untreated control

2.10 MICRONUTRIENT MIXTURE ON ENHANCING ECONOMICS OF CULTIVATION OF CROPS

Yadav and Patel (2013) reported that application of micronutrients (ZnSO_4 40 g + MnSO_4 20 g + CuSO_4 5 g + Borax 10 g) on Grand Naine banana gave the maximum additional net income of Rs 42812 per hectare with the cost benefit ratio of 1.717.

Patel *et al* (2010) observed that the foliar application of ZnSO_4 (0.5 %) + FeSO_4 (0.5 %) of banana cv Basrai at 3rd, 5th and 7th months after planting resulted in maximum gross income and net profit with higher cost benefit ratio (1.194).

According to Pathak *et al* (2011), the combined application of Fe (0.5 %) and Zn (0.5 %) at 3, 5 and 7 months after planting of banana cv Martaman (AAB, Silk) gave better B:C ratio.

Bauri *et al* (2014) found that foliar sprang of Borax (0.1 %) and 2,4-dichlorophenoxyacetic acid (30 ppm) at 5 and 20 days after last hand opening showed maximum B:C ratio of banana cv Martaman (*Musa* AAB, Silk).

3. MATERIALS AND METHODS

An investigation entitled “Formulation and evaluation of micronutrient mixture for foliar application in TC banana (*Musa* sp) var Nendran” was carried out at College of Agriculture, Padannakkad and Regional Agricultural Research Station (RARS) farm, Nileshwar during November 2014 to March 2016. The objectives of the study were to prepare micronutrient mixture as a liquid formulation and to study the effect of application of this mixture on hardening stage and in the field condition of tissue culture banana var Nendran. The whole study was conducted in three steps:

Part I Preparation of micronutrient mixture formulation

Part II studies on hardening of TC banana plants using foliar sprays of micronutrients

Part III Field experiment

3.1 PREPARATION OF MICRONUTRIENT MIXTURE FORMULATION

Micronutrient mixture containing zinc, boron, iron, copper, manganese and molybdenum was prepared at different concentrations and evaluated for stability and keeping quality in solution form. The different micronutrient salts used were Zinc sulphate, Boric acid, Copper sulphate, Manganous sulphate, Ferrous sulphate and Ammonium molybdate. The technique was standardized after trying a series of combinations. Finally the micronutrient mixture was formulated and it contains two separate mixtures: solution A and solution B. Each series of the nutrient solutions were evaluated for their phyto suitability by spraying on leaves and observing the leaves for one week. The solutions which produced toxic/ necrotic spots were discarded. The prepared nutrient mixture was then tested with tissue culture Nendran banana as a foliar application in both pot culture and field experiment. This work was done as a laboratory trial with the salts containing these nutrients.

3.2 POT CULTURE EXPERIMENT

The secondary hardening studies were conducted in mist chamber of Department of Plant Biotechnology, COA, Padannakkad with tissue culture plants of uniform stage multiplied from same genotype. The experiment was conducted in completely randomized design with four different levels and three different time of application of micronutrient mixture as a foliar spray. There were 12 treatment combinations and one control with three replications each. Four plants were maintained in each replication, so that the total number of plants was 156. The potting mixture was prepared uniformly with 1:1:1 ratio of sand, soil and cowdung and the plants were raised in polybags for a period of 20 days in mist chamber (primary hardening stage). The plants were transferred to bigger polybags and then secondary hardening was carried out in partial shade condition for a period of one month and this stage, different sprayings were given to plants. Plate 1 shows the secondary hardening of tissue culture banana var Nendran.

3.2.1. Design and Layout

Crop	Tissue culture banana
Variety	Nendran
Design	CRD
Treatments	13
Replication	3

3.2.2. Levels of nutrients

The standardized micronutrient mixture formulation was sprayed at 0.25 per cent, 0.5 per cent, 0.75 per cent and 1.0 per cent foliar sprays. The volume of spray solution was 150ml per plant.

Concentration	One spray	Two sprays	Three sprays
0.25 %	T ₁	T ₅	T ₉
0.5 %	T ₂	T ₆	T ₁₀
0.75 %	T ₃	T ₇	T ₁₁
1.0 %	T ₄	T ₈	T ₁₂

T₁₃ control

3.2.3. Time of application

One spray at immediately after planting, two sprays at immediately and two weeks after planting and three sprays at immediately, two and three weeks after planting during secondary hardening stage

3.2.4. Biometric observations

The important biometric observations like plant height, number of leaves, leaf length and leaf breadth were recorded one month after treatment imposition

3.2.4.1. Plant height

The plant height was measured from the base of the pseudostem to the youngest leaf

3.2.4.2. Number of leaves

The number of photosynthetically active leaves was counted

3.2.4.3. Leaf length

Leaf length was measured from petiole to tip of the leaf

3.2.4.4. Leaf breadth

Leaf breadth was measured from the widest portion of the leaf

3.3 FIELD EXPERIMENT

The field experiment was conducted at Regional Agricultural Research Station (RARS) farm, Nileshwar to evaluate the effect of micronutrient mixture as a foliar spray in tissue culture banana var Nendran. The experiment was conducted in randomized block design with thirteen treatments and three replications such that four plants were maintained in each replication. Nitrogen, phosphorus and potassium application and other cultural practices were followed as per Package of Practices recommendations, KAU (2011) uniformly for all the treatments. Plate 2 shows the field view of the experimental plot at RARS farm, Nileshwar.

3.3.1. Design and layout

Crop	Tissue culture banana
Variety	Nendran
Design	RBD
Treatments	13
Replication	3

3.3.2. Levels of nutrients

The standardized micronutrient mixture formulation was sprayed at 1 per cent, 2 per cent, 3 per cent and 4 per cent foliar sprays. The volume of spray solution was 750ml per plant. Package of Practices recommendations were followed for all treatments.

Concentration	One spray	Two sprays	Three sprays
1 %	T ₁	T ₅	T ₉
2 %	T ₂	T ₆	T ₁₀
3 %	T ₃	T ₇	T ₁₁
4 %	T ₄	T ₈	T ₁₂

Control-T₁₃ was PoP (Package of Practices recommendations)

3.3.3. Time of application

One spray at two months after planting, two sprays at two and four months after planting and three sprays at two, four and six months after planting

3.3.4. Biometric observations

Biometric observations like pseudostem height, pseudostem girth and number of leaves were taken four months after planting and at the time of shooting. Days to bunch emergence, incidence of pest and diseases were recorded. Days to harvest, number of suckers at harvest, bunch weight, number of hands per bunch, number of fingers per hand, average weight of finger, days to ripening, finger length and finger breadth were recorded at the time of harvesting.

3.3.4.1. Pseudostem height

The pseudostem height from the ground level of pseudostem upto the base of youngest leaf at 4 months after planting and upto the base of boot leaf at the time of shooting was measured and it is expressed in meter (m)

3.3.4.2. Pseudostem girth

Girth of pseudostem at 90cm of height from the ground level was measured at four months after planting and at the time of shooting and it is expressed in cm

3.3.4.3 Number of leaves

Total number of leaves was recorded at four months after planting and at the time of shooting

Figure 1 Layout of Field experiment

Block I	Block II	Block III
T ₅	T ₇	T ₂
T ₁₃	T ₁₁	T ₉
T ₂	T ₁	T ₁₂
T ₁₀	T ₉	T ₅
T ₈	T ₂	T ₇
T ₄	T ₅	T ₄
T ₇	T ₁₃	T ₁
T ₁₂	T ₆	T ₁₀
T ₃	T ₁₂	T ₆
T ₉	T ₁₀	T ₁₁
T ₆	T ₂	T ₁₃
T ₁₁	T ₄	T ₈
T ₁	T ₈	T ₃

3.3.4.4. Days to bunch emergence

Number of days to bunch emergence of individual treatments from the date of planting was recorded

3.3.4.5. Incidence of pest and diseases

Incidence of pest and diseases were monitored and necessary control measures also were taken from hardening stage to harvesting

3.3.4.6. Bunch maturity period

Number of days taken from day of bunch emergence to harvest of banana were recorded for individual treatment

3.3.4.7. Days to harvest

Number of days taken from the date of planting to the date of harvest was recorded for individual treatments

3.3.4.8. Number of suckers at harvest

The total number of suckers produced by the plant at the time of harvest was recorded.

3.3.4.9. Weight of male bud

Fresh weight of male bud one month after bunch emergence for individual treatments was recorded

3.3.4.10. Bunch weight

Bunch weight of individual treatment was recorded at the time of harvest

3.3.4.11. Number of hands per bunch

Number of hands in each bunch were counted at the time of harvest

3.3.4.12. Number of fingers per bunch

Total number of fingers in each bunch of all individual plants were counted at the time of harvest

3.3.4.13. Average weight of fingers

Average weight of fingers was calculated for individual bunch by taking weight of fingers from top, middle and bottom of bunch and averaged

3.3.4.14. Days to ripening

Number of days to ripening for individual treatments was recorded from the time of harvest to ripened fruit

3.3.4.15. Finger length

Average length of fingers was calculated by measuring length of finger in topmost, middle and bottom of the bunch and their average was taken

3.3.4.16. Finger breadth

Average breadth of fingers was calculated by measuring breadth of finger in topmost, middle and bottom of the bunch and their average was taken

3 4 FRUIT ANALYSIS

Quality parameters of banana such as Total Soluble Solids, titrable acidity, vitamin C content, total sugar and pulp to peel ratio were analyzed using standard

methods Loss in weight during ripening and keeping quality of fruits in ambient condition was also recorded

3.4.1. Total soluble solids (TSS)

Total soluble solids of the fully ripened banana fruit were analyzed using handheld portable refractometer and it is expressed in terms of °brix

3.4.2. Titrable Acidity

For analyzing titrable acidity of fully ripened banana fruits, 50 g of fruit was ground from which 25 g of pulped material was transferred to a 250 ml beaker. In that 100 ml of distilled water was added and boiled for 30 minutes. Once it is cooled, transferred to 250 ml standard flask and volume was made up and from that 50ml was taken which was diluted with equal quantity of hot water. Titrated against 0.1N NaOH using phenolphthalein indicator till the appearance of light pink colour which was the end point and titrable acidity was worked out and the results are expressed as percentage of maleic acid.

3.4.3. Reducing sugar content

Reducing sugar content was determined by titrating fruit juice with Fehling solution A and B as per procedure given by Ranganna (1986) and the results are expressed in percentage.

3.4.4. Total sugar content

Total sugar content was determined by titrating fruit juice with Fehling solution A and B and the results are expressed in percentage.

Materials and Methods

3.4.5. Non-reducing sugars

Non-reducing sugar content was determined by subtracting reducing sugars from total sugars content and the results are expressed in percentage

3.4.6. Pulp to peel ratio

Weight of pulp and peel was recorded separately and the relative proportion of pulp to peel was recorded (i.e. pulp weight divided by the peel weight)

3.4.7. Sugar / acid ratio

Total sugars (%) divided by acidity (%) gave the sugar / acid ratio of ripened fruit

3.4.8. TSS / acid ratio

TSS (°brix) divided by acidity (%) gave the TSS / acid ratio of ripened fruit

3.4.9. Keeping quality of fruits at ambient condition

Keeping quality of fruits was calculated by counting number of days from day of commencement of ripening of fruits to end of saleable life or edible life

3.4.10. Per cent loss in weight during ripening

Per cent loss in weight was calculated from weight of fruits at the time of harvest to weight of ripened fruit and it is expressed in percentage

3.5 SOIL ANALYSIS

3.5.1. Initial soil analysis

Soil samples for initial analysis were collected from the prepared field. Soil samples were drawn from surface 15 cm from 10 different places of the field which

was then pooled, reduced to required quantity and air dried. The air dried soil samples were ground and passed through 2 mm sieve and stored in air tight containers.

The samples were analyzed for bulk density, particle density, porosity, texture, pH, EC, CEC, organic carbon, available nutrients such as N, P, K, Ca, Mg, S, Fe, Cu, Zn, Mn, B and Mo following standard procedures given in table 1. The initial soil properties were given in table 2.

3.5.2. Experimental soil analysis

Soil samples for laboratory analysis were collected from all the treatments at the time of harvest. The samples were air dried, ground, sieved with 2mm sieve and stored in air tight container. They were analyzed for available nutrients such as N, P, K, Ca, Mg, S, Fe, Cu, Zn, Mn, B and Mo as per the standard procedures as given in the table 1.

3.6 PLANT ANALYSIS

Plant samples were collected from the index leaf (third fully opened leaf from top) at the time of bunching and at harvest and analyzed for various macro and micronutrient status in it using standard procedures as given in the table 3.

3.7 BENEFIT COST RATIO

Cost of cultivation for all the treatments was worked out on the basis of prevailing input cost and market price of fruits, male bud and suckers at the time of experimentation. The net income was calculated by deducting the cost of cultivation from the gross return. The benefit cost ratio (BCR) was worked out as follows:

$$BCR = \frac{\text{Gross return ha}^{-1} (\text{Rs})}{\text{Cost of cultivation ha}^{-1} (\text{Rs})}$$

3 8 STATISTICAL ANALYSIS

The data obtained from pot culture and field experiment was analyzed statistically and tested for its significance using WASP 2.0 software given by ICARGOA

Table 1 Analytical methods followed for soil analysis

S No	Parameters	Method	Reference
1	pH	pH meter	Jackson (1958)
2	EC	Conductivity meter	Jackson (1958)
3	Organic carbon	Chromic acid wet digestion method	Walkley and Black (1934)
4	Bulk density	Undisturbed core sample	Black <i>et al</i> (1965)
5	Particle density	Pycnometer method	Black <i>et al</i> (1965)
6	Porosity	-	Black <i>et al</i> (1965)
7	Textural analysis	International pipette method	Robinson (1922)
8	Available N	Alkaline permanganate method	Subbiah and Asija (1956)
9	Available P	Bray extraction and photoelectric colorimetry	Jackson (1958)
10	Available K	Flame photometry	Pratt (1965)
11	Available Ca	Atomic absorption spectroscopy	Jackson (1958)
12	Available Mg	Atomic absorption spectroscopy	Jackson (1958)

13	Available S	Photoelectric colorimetry	Massouni and Comfield (1963)
14	Available Zn	Atomic absorption spectroscopy	Emmel <i>et al</i> (1977)
15	Available B	Photoelectric colorimetry	Bingham (1982)
16	Available Fe	Atomic absorption spectroscopy	Sims and Johnson (1991)
17	Available Cu	Atomic absorption spectroscopy	Emmel <i>et al</i> (1977)
18	Available Mn	Atomic absorption spectroscopy	Sims and Johnson (1991)
19	Available Mo	Inductively coupled plasma optical emission spectrometry (ICP OES)	Soltanpour and Schwab (1977)

Table 2 Properties of the initial soil sample

S No	Parameter	Value
I Physical properties		
1	Bulk density (g cm^{-3})	1.34
2	Particle density (g cm^{-3})	2.35
3	Pore space (%)	43%
II Mechanical composition		
1	Sand (%)	77.25
2	Silt (%)	19.00

Table 2 Continued

3	Clay (%)	3 75
4	Textural class	Loamy sand
III Chemical properties		
1	pH	5 31
2	EC (dS/m)	0 08
3	Organic carbon (%)	0 40
4	Organic matter (%)	0 69
5	CEC (meq/100g)	7 50
6	Available N (kg/ha)	213 20
7	Available P (kg/ha)	82 69
8	Available K (kg/ha)	184 73
9	Available Ca (mg/kg)	148 25
10	Available Mg (mg/kg)	23 65
11	Available S (mg/kg)	6 25
12	Available Zn (mg/kg)	0 91
13	Available B (mg/kg)	0 26
14	Available Fe (mg/kg)	26 8
15	Available Cu (mg/kg)	2 63
16	Available Mn (mg/kg)	22 10
17	Available Mo (mg/kg)	0 002

Table 3 Analytical methods followed for plant analysis

S.No	Parameter	Method	Reference
1	Total N	Modified kjeldhal digestion method	Jackson (1958)
2	Total P	Vanadomolybdate yellow colour method	Piper (1966)
3	Total K	Flame photometry	Jackson (1958)
4	Total Ca	Atomic absorption spectroscopy	Issac and Kerber (1971)
5	Total Mg	Atomic absorption spectroscopy	Issac and Kerber (1971)
6	Total S	Turbidimetric method	Bhargava and Ragupathi (1995)
7	Total Zn	Atomic absorption spectroscopy	Emmel <i>et al</i> (1977)
8	Total B	Azomethane - H colorimetric method	Bingham (1982)
9	Total Fe	Atomic absorption spectroscopy	Piper (1966)
10	Total Cu	Atomic absorption spectroscopy	Emmel <i>et al</i> (1977)
11	Total Mn	Atomic absorption spectroscopy	Piper (1966)
12	Total Mo	Inductively coupled plasma optical emission spectrometry (ICP OES)	Soltanpour and Schwab (1977)



Plate 1a Secondary hardening of TC plants (using micronutrient mixture formulation)



Plate 1b Treatment vs Control

Plate 1 Secondary hardening of tissue culture banana var. Nendin (using micronutrient mixture formulation)



Plate 2 a Field view - 2 MAP



Plate 2 b Field view - 4 MAP



Plate 2 c Field view at bunching stage

Plate 2 Field view of the experimental plot at RARS farm, Nilleshwar

Results

4. RESULTS

An investigation was carried out to formulate micronutrient mixture in liquid formulation and to examine the effect of micronutrient mixture in tissue culture banana var Nendran during hardening stage as well as in field. The results of biometric observations (taken after hardening stage and in field experiment) and the various soil, leaf and fruit characteristics were statistically analyzed and the results are presented below.

4.1 PREPARATION OF MICRONUTRIENT MIXTURE AS A LIQUID FORMULATION

Liquid micronutrient mixture for foliar application was formulated using different micronutrient salts. Different combinations were tried and its keeping quality, stability and compatibility for foliar application was evaluated, finally it is standardized in 2 parts solution A and solution B. Solution A consisting of Zinc sulphate, Copper sulphate, Ferrous sulphate, Boric acid, Manganous sulphate and Ammonium molybdate in different proportion. Solution B contains 1 per cent humic acid which acts as a natural chelating agent for improving micronutrient use efficiency. These two solutions are to be diluted and blended before sprays. Plate 3 shows preparation of micronutrient mixture as a liquid formulation.

4.1.1. Composition of Solution A (1 litre)

ZnSO ₄ 7H ₂ O	- 100 g
CuSO ₄ 5H ₂ O	- 40 g
FeSO ₄ 7H ₂ O	- 20 g
H ₃ BO ₃	20 g
MnSO ₄ H ₂ O	- 1 g
(NH ₄) ₆ Mo ₇ O ₂₄ 4H ₂ O	- 0.5 g

This micronutrient mixture was prepared using these salts but it did not have desired pH for foliar spraying and the its keeping quality and stability were poor. The pH of the mixture was 3.4 and there were precipitates in formulation and it was visually not a clear solution.

4.1.2. Stability and keeping quality of the formulation

For improving the stability and keeping quality of the micronutrient mixture, various chemicals were tried with solution A and the keeping quality of the mixture was evaluated. The different chemicals used were Ammonia (5%), nitric acid (5%), tartaric acid (10%) and citric acid (20%). Except citric acid other chemicals were forming precipitate with micronutrient mixture which is presented in Plate 3 c. When 30 ml of citric acid (20%) was added to 1 litre of nutrient mixture it became clear liquid, there was no precipitation of nutrient compounds and its keeping quality was excellent. It was further monitored at specific intervals to evaluate its stability and keeping quality. Solution A with 30ml/litre of citric acid (20%) can be kept as such at ambient conditions upto 1 year whereas with nitric acid and ammonia it forms precipitate within a day and with tartaric acid it is stable upto a maximum of one week.

Solution B contains 1 per cent humic acid, from solution B 30 ml/litre was uniformly added to final diluted solution of A. Both solutions should be mixed thoroughly just before spraying.

4.1.3. Properties of micronutrient mixture

The pH of micronutrient mixture was noted just before spraying and it was observed as 5.56 at the desired dilution. Colour of Solution A is Green and Solution B is Brown. Based on the properties like pH, keeping quality and trial on standing crop, formulated micronutrient mixture was found suitable for foliar application on banana.



Plate 3 a Micronutrient mixture with and without citric acid

Plate 3 b Micronutrient mixture solution A and solution B

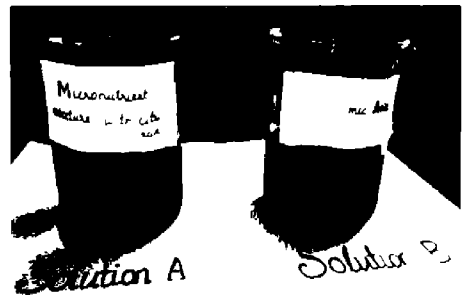


Plate 3 c Keeping quality of solution A with different chemicals

Plate 3 Preparation of Micronutrient mixture as a liquid formulation

4.2 POT CULTURE EXPERIMENT

Tissue culture banana plants of uniform stage multiplied from the same genotype were obtained from Department of Plant Biotechnology, College of Agriculture, Padannakkad. The secondary hardening studies were done in the mist chamber with various treatment combinations of micronutrient mixture as foliar spray. The treatments consisted of 4 levels of micronutrient mixture sprayed at 3 different intervals. The different treatment combinations were T₁, T₅, T₉ at 0.25 per cent (one, two and three sprays), T₂, T₆, T₁₀ at 0.50 per cent (one, two and three sprays), T₃, T₇, T₁₁ at 0.75 per cent (one, two and three sprays), T₄, T₈, T₁₂ at 1 per cent (one, two, three sprays) and T₁₃- control. One spray was given immediately after planting, two sprays at immediately and two weeks after planting and three sprays at immediately, two and three weeks after planting. The experiment was intended to evaluate the role of this micronutrient mixture formulation on performance of plants in hardening stage. Periodic biometric observations were also made. The biometric observations of hardening studies were taken at the end of one month after treatment imposition.

4.2.1. Plant height

The results of the effect of micronutrient mixture on biometric characters of TC banana are given in Table 4. All the treatments were found to be superior to the control. Among the treatments, T₁₀ (0.5% as three time spraying) recorded highest plant height (20.2 cm) which was on statistically par with treatments T₉ and T₅ whereas the control T₁₃ recorded lowest plant height (16.6 cm) which was on par with T₄, T₁₂, T₈ and T₇.

Table 4 Foliar application of micronutrient mixture on biometric characters of TC banana

Treatments	Plant height (cm)	Number of leaves	Leaf length (cm)	Leaf breadth (cm)
T ₁	18 63	6 58	17 24	6 43
T ₂	19 08	6 58	17 95	6 21
T ₃	18 81	6 08	18 07	6 83
T ₄	16 88	5 83	15 63	5 73
T ₅	19 93	6 08	19 58	7 03
T ₆	18 68	6 33	18 24	6 49
T ₇	17 38	5 42	16 29	6 32
T ₈	17 15	5 50	16 66	5 94
T ₉	20 04	6 75	18 32	7 33
T ₁₀	20 21	6 67	19 33	6 94
T ₁₁	17 46	5 75	17 11	6 51
T ₁₂	16 98	5 33	16 47	6 10
T ₁₃	16 62	5 08	16 68	5 93
SEm (±)	0 369	0 103	0 800	0 199
CD (0 05)	1 019	0 538	1 501	0 749

4.2.2. Number of leaves

Number of leaves was counted and is presented in Table 4. Maximum leaf number was recorded with T₉ (6.8) which was statistically on par with T₁₀, T₁, T₂ and T₆. T₁₃ (control) recorded minimum leaf number (5.1).

4.2.3. Leaf length

There was significant difference between treatments with respect to leaf length (Table 4). Leaf length was found to be high in T₅ (0.25% - two sprays) with 19.6 cm followed by T₁₀, T₉ and T₆ and lowest was T₄ (15.6 cm) which was on par with T₇.

4.2.4. Leaf breadth

There was notable difference in leaf breadth between the treatments (Table 4). Leaf breadth found to be maximum in T₉ (0.25% - three sprays) with 7.3 cm which was statistically on par with T₅ and minimum leaf breadth was found in T₄ (5.7 cm) which was on par with T₁₃ and T₈.

4.3 FIELD EXPERIMENT

The field experiment was conducted at Regional Agricultural Research Station farm, Nileshtar to study the effect of foliar application of micronutrient mixture in tissue culture banana var Nendran. The experiment was conducted in RBD with four different levels and three different time of application of micronutrient mixture as a foliar spray. There were 12 treatment combinations and one control with three replications each. Four plants were maintained in each replication, so that the total number of plants was 156. The different treatment combinations were T₁, T₅, T₉ at 1 per cent (one, two and three sprays), T₂, T₆, T₁₀ at 2 per cent (one, two and three sprays), T₃, T₇, T₁₁ at 3 per cent (one, two and three

sprays), T₄, T₈, T₁₂ at 4 per cent (one, two and three sprays) and T₁₃- control One spray was given at two months after planting, two sprays at two and four months after planting and three sprays at two, four and six months after planting Periodic growth and yield parameters were recorded, detailed soil, fruit and leaf analysis were carried out and statistically analyzed The salient finding of field experiments are presented below

4.3.1. Effect on vegetative characters

4.3.1.1. Pseudostem height

The pseudostem height was recorded at 4 months after planting and at the time of shooting and the results are presented in table 5 The height of the pseudostem was significantly influenced by the nutrient treatments at 4 months after planting T₁₂ recorded maximum plant height of 1.5 m while T₅ recorded lowest height (1.2 m) whereas control T₁₃ was recorded with 1.3 m T₁₂ was statistically on par with T₁₀ (1.4 m), T₁ (1.4 m), and T₈ (1.4 m)

At the time of shooting, there was no notable difference between the treatments on pseudostem height (Table 5) Even though not significant, T₆ recorded maximum pseudostem height (2.9 m) followed by T₁₀ and T₁₁ (2.8 m) whereas T₄ recorded minimum height of 2.7 m whereas control T₁₃ was recorded with 2.7 m

4.3.1.2. Pseudostem girth

Pseudostem girth at 90 cm height was measured at 4 months after planting and at the time of shooting (Table 5) T₁₂ (29.6 cm) recorded maximum pseudostem girth at 4 months after planting followed by T₁₀ (29.0 cm) and minimum girth was found in T₅ (23.8 cm) However, there was no significant difference between the treatments and control

There was notable difference in pseudostem girth between the treatments at the time of shooting (Table 5) T₁₀ recorded maximum girth of 52.1 m which was

statistically on par with T₆ (51.5 cm), T₇ (51.1 cm), T₅ (50.7 cm) and T₁₁ (50.0 cm) and minimum pseudostem girth was found in T₃ (47.7 cm) which was on par with control T₁₃ (48.0 cm)

Table 5 Foliar application of micronutrient mixture on pseudostem height and girth of banana

Treatments	Pseudostem height (m)		Pseudostem girth (cm)	
	4 months after planting	At the time of shooting	4 months after planting	At the time of shooting
T ₁	1.36	2.75	28.28	51.12
T ₂	1.30	2.79	24.70	49.17
T ₃	1.24	2.70	24.23	47.67
T ₄	1.21	2.68	24.08	48.75
T ₅	1.18	2.77	23.75	50.71
T ₆	1.24	2.85	26.01	51.54
T ₇	1.26	2.69	26.15	48.46
T ₈	1.35	2.83	28.23	48.42
T ₉	1.25	2.73	26.84	48.89
T ₁₀	1.39	2.84	28.96	52.08
T ₁₁	1.25	2.84	26.87	49.96
T ₁₂	1.47	2.76	29.64	48.50
T ₁₃	1.32	2.69	25.07	47.96
SEm (±)	0.006	0.011	8.054	2.579
CD (0.05)	0.130	NS	NS	2.706

4.3.1.3. Number of leaves

Number of leaves at both stages of observation exhibited significant differences among the treatments (Table 6). At 4 months after planting T₆ recorded highest number of leaves (11.7) which was statistically on par with T₁ (11.6), T₉ (11.2), T₃ (11.2), T₁₀ (11.0), T₂ (11.0), T₈ (10.9) and the lowest number of leaves was produced in control T₁₃ (9.6).

At the time of shooting, T₆ recorded highest number of leaves (12.8) which was on par with T₁₁ (12.3), T₁ (11.9), T₄ (11.8) while T₁₃ (10.3) recorded lowest number of leaves.

4.3.1.4 Number of suckers at harvest

The data on number of suckers at the time of harvest is presented in Table 6. There was no considerable difference between the treatments. Even though not significant, T₁₀ and T₁₂ recorded maximum number of suckers (7.3) followed by T₃ and T₇ (6.8) while T₄ and control T₁₃ recorded minimum number of suckers at the time of harvest (5.8).

4.3.2. Effect of yield attributes

4.3.2.1. Weight of male bud

Weight of male bud at the time of detachment was noted and compared (Table 7). There was a significant difference among the treatments with respect to weight of male bud. T₅ recorded highest male bud weight (1.5 kg) which was statistically on par with T₁₁ (1.5 kg), T₂ (1.4 kg), T₈ (1.4 kg) and lowest male bud weight was found in T₁ (1.1 kg) whereas control T₁₃ recorded 1.2 kg.

Table 6 Foliar application of micronutrient mixture on number of leaves and suckers at harvest of banana

Treatments	Number of leaves		Number of suckers at harvest
	4 months after planting	At the time of shooting	
T ₁	11 58	11 92	6 00
T ₂	11 00	11 08	6 08
T ₃	11 17	11 33	6 83
T ₄	10 58	11 83	5 83
T ₅	10 83	11 50	6 33
T ₆	11 67	12 75	6 25
T ₇	10 83	11 50	6 83
T ₈	10 92	11 17	6 42
T ₉	11 17	11 42	6 75
T ₁₀	11 00	11 25	7 33
T ₁₁	10 50	12 25	6 25
T ₁₂	10 67	11 08	7 33
T ₁₃	9 583	10 25	5 83
SEm (±)	0 216	0 458	0 678
CD (0 05)	0 784	1 140	NS

4.3.2.2. Bunch weight

Bunch weight of plant is the major economic factor considered in banana cultivation. There was a significant influence of treatments on bunch weight and the results are presented in Table 7. All the treatments were found to be superior over control on which T₁₁ recorded maximum bunch weight (12.8 kg) and control T₁₃ recorded minimum bunch weight (8.3 kg) which was on par with T₃. T₁₁ was statistically on par with T₁₀ (12.7 kg), T₉ (12.2 kg), T₆ (12.2 kg), T₇ (12.1 kg), T₈ (11.9 kg) and T₅ (11.5 kg). Plate 4 shows comparison of bunch weight of selected treatments and control.

4.3.2.3. Number of hands per bunch

There was no significant difference between the treatments with respect to number of hands per bunch (Table 7). However T₃, T₆ and T₇ were recorded with 5.4 hands per bunch while T₂ recorded minimum number of hands per bunch (4.8) while control T₁₃ was having 5.0 hands per bunch.

4.3.2.4. Number of fingers per bunch

The data on total number of fingers per bunch was recorded at the time of harvest and it is presented in Table 8. There was significant difference between the treatments. T₉ recorded higher number of fingers per bunch (54.4) and T₁₃ recorded lowest number of fingers per bunch (46.7) followed by T₁₂ (50.4). All other treatments were on par with T₉. Plate 4 f shows the highest number of fingers per bunch in best treatment.

4.3.2.5. Average weight of fingers

The average weight of fingers was calculated by taking the best finger in each hand per bunch and averaged. There was notable difference in average weight of fingers between the treatments (Table 8). T₁₁ produced maximum average weight of

fingers of 282.6 g whereas minimum was found in T₁ with 222.5 g which was statistically on par with T₅ and control T₁₃. T₁₁ was on par with T₁₀, T₉, T₆, T₇, T₈, T₄ and T₁₂. Plate 5 shows the finger characteristics of treatments and control.

Table 7 Foliar application of micronutrient mixture on yield parameters of banana

Treatments	Weight of male bud (kg)	Bunch weight (kg)	No of hands per bunch
T ₁	1.12	10.60	5.33
T ₂	1.40	10.22	4.83
T ₃	1.25	9.55	5.44
T ₄	1.35	10.48	4.89
T ₅	1.54	11.45	5.17
T ₆	1.24	12.19	5.44
T ₇	1.35	12.07	5.44
T ₈	1.38	11.90	5.33
T ₉	1.24	12.23	4.97
T ₁₀	1.30	12.73	5.00
T ₁₁	1.48	12.76	5.11
T ₁₂	1.25	11.02	5.00
T ₁₃	1.18	8.32	5.00
SEm (±)	0.007	0.766	0.127
CD (0.05)	0.177	1.475	NS

Table 8 Foliar application of micronutrient mixture on finger characteristics of banana

Treatments	No of fingers per bunch	Average weight of fingers (g)	Average Finger length (cm)	Average finger breadth (cm)
T ₁	51 33	222 50	21 15	13 26
T ₂	51 08	233 34	20 78	14 68
T ₃	53 33	237 38	21 28	14 00
T ₄	50 89	247 98	21 11	14 48
T ₅	53 89	225 88	20 62	14 38
T ₆	53 00	271 65	20 55	14 60
T ₇	53 67	268 80	22 38	14 94
T ₈	51 67	256 13	22 15	14 15
T ₉	54 39	271 75	22 50	13 99
T ₁₀	53 94	275 00	22 69	15 25
T ₁₁	54 06	282 64	23 05	15 25
T ₁₂	50 44	245 06	21 75	14 16
T ₁₃	46 68	230 00	17 94	13 02
SEm (±)	4 345	306 077	0 633	0 653
CD (0 05)	3 513	38 122	1 734	NS

4.3.2.6. Average length of fingers

The effect of treatments on average length of fingers is presented in Table 8 T₁₁ was recorded with maximum finger length of 23.1 cm while control T₁₃ recorded minimum length of 17.9 cm T₁₁ was on par with T₁₀ (22.7 cm), T₉ (22.5 cm), T₇ (22.4 cm), T₈ (22.2 cm) and T₁₂ (21.8 cm)

4.3.2.7. Average breadth of fingers

The effect of treatment application on finger girth is presented in Table 8 Treatment effects on finger breadth at the middle portion was found to be non significant Maximum finger breadth was noticed in T₁₁ and T₁₀ with 15.3 cm whereas control T₁₃ recorded minimum finger breadth of 13.0 cm

4.3.2.8. Days to bunch emergence

Effect of treatment application on the number of days taken from planting to bunch emergence was recorded and presented in Table 9 There was no appreciable difference between the treatments T₁₀ produced earliest bunch in 186 days after planting while control T₁₃ produced bunch in 198 days after planting which was the late one An average bunch emergence happened between 185-197 days

4.3.2.9. Days to harvest

There was a remarkable difference between the treatments with respect to days to harvest Effect of treatment application on the number of days taken from planting to harvest was recorded and presented in Table 9 T₁₃ (control) showed maximum days from planting to harvest (303 days) while T₁₁ took minimum of 271 days T₁₁ was statistically on par with T₁₀ (272 days) and T₉ (274 days)

4.3.2.10. Bunch maturity period

Effect of treatment application on bunch maturity period (from the day of bunch emergence to harvest) was recorded and presented in Table 9. There was a remarkable difference between the treatments. T₁₁ took minimum maturity period of 82 days which was on par with T₁₀ (86 days) and T₉ (88 days) while control T₁₃ took maximum days to maturity with 106 days.

4.3.2.11. Days to ripening

Effect of treatment application on the number of days taken from harvest to ripening stage of fruit was recorded and presented in Table 9. Treatments found to be non significant in influencing the days to ripening. T₄ was recorded with minimum of 4 days to ripening while T₁₃ took 6 days.

4.3.2.12. Incidence of pest and disease

The banana-leaf caterpillar (*Spodoptera litura*) was prevailed at 2 and 3 months after planting. The major disease noticed was sigatoka leaf spot at 4 months after planting caused by *Mycosphaerella musicola*. Necessary control measures were taken uniformly for all the plants.

Table 9 Foliar application of micronutrient mixture on days to bunch emergence, days to harvest, Bunch maturity period and days to ripening of banana

Treatments	Days to bunch emergence	Days to harvest	Bunch maturity period (days)	Days to ripening
T ₁	190 83	283 67	92 83	5 50
T ₂	190 67	293 33	102 67	4 83
T ₃	194 78	286 67	91 89	5 50
T ₄	195 83	289 00	93 17	4 33
T ₅	195 67	290 33	94 67	5 83
T ₆	191 11	282 67	91 56	6 00
T ₇	191 00	283 33	92 33	5 00
T ₈	190 67	283 67	93 00	6 07
T ₉	185 83	274 00	88 17	4 67
T ₁₀	185 72	271 67	85 94	5 65
T ₁₁	188 67	270 67	82 00	5 42
T ₁₂	191 08	283 33	92 25	5 67
T ₁₃	197 67	303 17	105 50	6 00
SEm (±)	22 870	25 982	25 918	1 176
CD (0 05)	NS	8 590	8 580	NS



Plate 4 a Best treatment T₁₁ (3 % as sprays)



Plate 4 b 2nd best treatment T₁₀ (2 % as 3 sprays)

Plate 4 c Control T₁



Plate 4 d Highest number of fingers per bunch

Plate 4 Bunches from best treatments vs control



Plate 5 a Highest average weight of fingers T₁ (3 % as 3 sprays)



Plate 5 b Control T

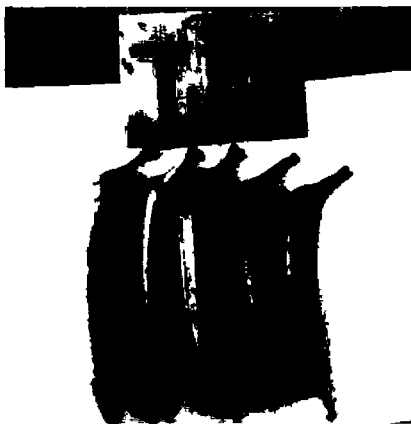


Plate 5 c 2 - best treatment - T₁ (2 % as 3 sprays)

Plate 5 d Control - T₀



Plate 5 e Finger of best treatment - T₁ (3 - as - sprays)

Plate 5 f Finger characteristics of treatments and control

4.4 FRUIT CHARACTERISTICS

The effect of foliar application of micronutrient mixture on fruit characteristics of banana was examined. Various quality parameters like titrable acidity, vitamin C, TSS, reducing sugar content, total sugars, per cent loss in weight during ripening, keeping quality of the fruits at ambient condition and pulp to peel ratio were analyzed and the results are presented below.

4.4.1 Titrable acidity

The effect of treatment application on titrable acidity (%) of ripened banana fruits is presented in Table 10. The treatments had notable influence on titrable acidity as compared to control. T₁₃ was recorded highest acidity of 0.53 % which was on par with T₁ (0.50 %), T₂ (0.45 %) and T₃ (0.43 %) while T₁₁ was recorded with minimum acidity of 0.26 % followed by T₁₀ (0.27 %) and T₉ (0.28 %).

4.4.2. Total Soluble Solids (TSS)

There was notable difference among the treatments with respect to total soluble solids (Table 10). T₁₁ recorded maximum TSS content (30.0 °brix) which was on par with T₁₀ (29.9 °brix), T₉ (29.7 °brix) and T₄ (29.5 °brix). Control T₁₃ recorded minimum TSS of 26.3 °brix which was statistically on par with T₁ (26.6 °brix).

4.4.3. Pulp to peel ratio

The effect of treatment application on pulp to peel ratio is presented in Table 10. There was significant difference among the treatments. T₁₂ recorded highest pulp to peel ratio of 3.7 followed by T₁₁ (3.6). Lowest ratio was found in T₂ (2.8) which were on par with control T₁₃ (2.9) and T₁ (2.9).

Table 10 Foliar application of micronutrient mixture on titrable acidity, TSS and pulp to peel ratio of ripened banana fruit

Treatments	Titrable acidity (%)	TSS (°brix)	Pulp to peel ratio
T ₁	0.50	26.60	2.91
T ₂	0.45	27.30	2.79
T ₃	0.43	27.70	3.13
T ₄	0.38	29.50	3.10
T ₅	0.36	28.20	3.41
T ₆	0.36	28.70	3.37
T ₇	0.29	28.40	3.06
T ₈	0.39	28.80	3.61
T ₉	0.28	29.70	3.52
T ₁₀	0.27	29.90	3.48
T ₁₁	0.26	30.00	3.64
T ₁₂	0.31	27.70	3.70
T ₁₃	0.53	26.30	2.88
SEm (±)	0.003	0.121	0.050
CD (0.05)	0.112	0.756	0.488

4.4.4. Total sugars

The influence of treatment application on total sugars percentage of ripened banana fruits are presented in Table 11. Higher sugar content and lower acidity indicates better fruit quality. The effect of treatments was found to be statistically significant. Maximum total sugars content was recorded in T₁₀ with 21.3 % which was statistically on par with T₉ (21.0 %), T₄ (19.7 %) and T₁₁ (19.4 %). Control T₁₃ recorded minimum of 13.3 % followed by T₁ (14.1 %).

4.4.5. Reducing sugars

The effect of treatment application was highly evident in case of reducing sugar content of fruits and the treatments showed superior and significant differences. Among the treatments, T₁₀ exhibited highest reducing sugar content of 19.0 per cent which was on par with T₁₁ (17.5 %) whereas T₁ recorded lowest reducing sugar content of 9.3 % followed by control T₁₃ (9.7 %). The results of treatment application on reducing sugar content of the ripened banana fruits are presented in Table 11.

4.4.6. Non-reducing sugars

There was notable difference among the treatments with respect to non-reducing sugar content of ripened banana fruits (Table 11). The highest non-reducing sugar content was found in T₅ with 4.9 per cent while T₁₁ recorded lowest non-reducing sugar content (1.9 %). Except T₁₁ and T₁₀ all other treatments were statistically on par with T₅.

4.4.7. Sugar / acid ratio

The influence of treatment application on sugar / acid ratio of ripened banana fruits are presented in Table 11. The effect of treatments was found to be statistically significant. Highest sugar / acid ratio were found in T₁₀ with 78.7 which were

statistically on par with T₁₁, T₉ and T₇ Control T₁₃ recorded lowest sugar / acid ratio of 25.3 which was on par with T₁, T₂ and T₃

4.4.8. TSS / acid ratio

The effect of treatment application was highly evident in case of TSS / acid ratio of fruits and the treatments showed superior and significant differences. Highest TSS/ acid ratio indicates better fruit quality. Among the treatments, T₁₁ exhibited highest TSS / acid ratio of 116.1 whereas control T₁₃ reported lowest TSS / acid ratio of 50.2. T₁₁ was on par with T₁₀, T₉ and T₇. The results of treatment application on the TSS / acid ratio of the ripened banana fruits are presented in Table 12.

4.4.9. Per cent loss in weight during ripening

There was significant difference between treatments with respect to per cent loss in weight during ripening of fruits (Table 12). Per cent loss in weight from day of harvest to ripened fruit was recorded and statistically analyzed. Maximum per cent loss in weight was found in T₁₃ with 16.7 while minimum loss was recorded in T₁₁ (11.9%) which was on par with T₁₀, T₄, T₁₂, T₃, T₆ and T₂.

4.4.10. Keeping quality of fruits at ambient condition

Treatment application on keeping quality of fruits was recorded from the day of ripening to the end of saleable or edible life of fruits. Keeping quality of fruits showed notable difference among the treatments which is presented in Table 12. Maximum keeping quality was recorded in T₁₀ with 4.40 days which was on par with T₁₁ (4.37 days) and T₉ (4.08 days). T₃ was found to be the minimum with 3.30 days while control T₁₃ was recorded with 3.43 days.

Table 11 Foliar application of micronutrient mixture on total sugars, reducing sugars, non-reducing sugars and sugar / acid ratio of ripened banana fruit

Treatments	Total sugars (%)	Reducing sugars (%)	Non-reducing sugars (%)	Sugar / acid ratio
T ₁	14.07	9.33	4.75	28.31
T ₂	16.38	12.13	3.88	36.84
T ₃	17.04	12.25	4.79	40.12
T ₄	19.70	15.45	4.25	52.63
T ₅	17.86	12.98	4.89	52.92
T ₆	17.80	14.05	3.75	50.12
T ₇	17.46	13.29	4.17	62.80
T ₈	18.36	13.51	4.85	50.15
T ₉	20.95	17.25	3.70	74.12
T ₁₀	21.25	19.00	2.25	78.70
T ₁₁	19.40	17.50	1.90	75.43
T ₁₂	16.41	11.98	4.44	53.40
T ₁₃	13.26	9.68	3.58	25.34
SEM (±)	0.798	0.494	0.508	68.512
CD (0.05)	1.946	1.531	1.553	18.036

Table 12 Foliar application of micronutrient mixture on TSS / acid ratio, per cent loss in weight during ripening and keeping quality of fruits at ambient condition

Treatments	TSS / acid ratio	Per cent loss in weight during ripening (%)	Keeping quality of fruits at ambient condition (days)
T ₁	53.70	15.00	3.46
T ₂	61.33	14.15	3.38
T ₃	65.33	13.78	3.30
T ₄	78.93	12.50	3.83
T ₅	83.56	14.96	3.43
T ₆	80.85	13.88	3.75
T ₇	102.31	15.96	3.73
T ₈	78.90	16.69	3.43
T ₉	105.20	16.33	4.08
T ₁₀	110.74	12.25	4.40
T ₁₁	116.07	11.87	4.37
T ₁₂	90.17	13.65	3.75
T ₁₃	50.20	16.70	3.43
SEm (±)	133.859	1.892	1.130
CD (0.05)	25.210	2.997	2.316

4.5 SOIL CHARACTERISTICS

4.5.1. pH

There was no considerable difference among the treatments with respect to soil pH (Table 13). T₉ recorded the lowest soil pH value of 4.83 and T₈ had the highest pH 5.52 while control T₁₃ recorded pH value of 5.42.

4.5.2. EC

Electrical conductivity of the soils was not influenced by the foliar application of micronutrient mixture (Table 13). Even though not significant, highest EC value was found in T₆ (0.27 dS/m) followed by T₁₁ (0.25 dS/m) and the lowest EC value was found in T₈ (0.16 dS/m).

4.5.3. Organic carbon content

Treatment application had significant effect on the organic carbon content of soils (Table 13). Highest OC was recorded in T₉ (0.7%) which was on par with T₁₂ (0.7%), T₁₁ (0.6%), T₈ (0.6%), T₁₀ (0.6%), T₃ (0.6%) and minimum OC was found in T₁₃ (0.4%) which was statistically on par with T₄ and T₁.

4.5.4. Available N

The results of available N content of soils are presented in Table 14. There was significant difference among the treatments with respect to available N content of soils. T₁₂ recorded highest available nitrogen content of soil (338.6 kg/ha) followed by T₁₁ (263.4 kg/ha) and the lowest available nitrogen content was found in T₃ (175.5 kg/ha) while control T₁₃ recorded 219.4 kg/ha.

Table 13 Foliar application of micronutrient mixture on soil pH, EC and Organic carbon

Treatments	pH	EC (dS/m)	Organic carbon (%)
T ₁	5.04	0.19	0.43
T ₂	5.18	0.20	0.51
T ₃	5.13	0.20	0.60
T ₄	5.05	0.18	0.43
T ₅	5.25	0.23	0.51
T ₆	5.13	0.27	0.44
T ₇	5.12	0.17	0.57
T ₈	5.52	0.16	0.62
T ₉	4.83	0.24	0.73
T ₁₀	5.14	0.18	0.61
T ₁₁	5.03	0.25	0.63
T ₁₂	4.94	0.24	0.66
T ₁₃	5.42	0.17	0.40
SEm (±)	0.277	0.004	0.009
CD (0.05)	NS	NS	0.163

4.5.5. Available P

The effect of treatment application was found to be significant in case of available P content of soils (Table 14). Highest P content was recorded in T₁₂ with 110.4 kg/ha whereas T₁₁ found to have lowest P content of 80.7 kg/ha. T₁₂ was

statistically on par with T₅, T₁₃, T₁₀ and T₆ Control T₁₃ was recorded with 102.8 kg/ha

Table 14. Foliar application of micronutrient mixture on available N, P and K content of soil

Treatments	Available N (kg/ha)	Available P (kg/ha)	Available K (kg/ha)
T ₁	238.20	93.04	412.67
T ₂	225.70	88.70	272.33
T ₃	175.50	92.31	272.00
T ₄	231.90	85.93	373.33
T ₅	250.80	110.00	425.00
T ₆	250.80	101.29	253.33
T ₇	231.90	96.91	348.00
T ₈	200.60	90.18	428.67
T ₉	231.90	85.26	349.33
T ₁₀	238.20	102.52	322.00
T ₁₁	263.40	80.65	308.67
T ₁₂	338.60	110.39	292.00
T ₁₃	219.40	102.84	417.67
SEm (±)	543.185	52.829	214.735
CD (0.05)	50.784	12.249	24.695

4.5.6. Available K

The effect of treatment application was significant in case of available K content of soil which is presented in Table 14. The highest K content was recorded in T₈ with 428.7 kg/ha which was statistically on par with T₅, T₁₃ and T₁ while lowest K

content was found in T₆ with 253.3 kg/ha. Control T₁₃ was found to have 417.7 kg/ha.

4.5.7. Available Ca

The effect of treatment application on the available Ca status of soil is presented in Table 15. Available Ca content in the soil was found to be non significant. However, maximum Ca content was found in T₅ (533.3 mg/kg) followed by T₄ (501.7 mg/kg) and minimum was found in T₈ with 323.3 mg/kg while control T₁₃ was with 456.7 mg/kg.

4.5.8. Available Mg

There was a significant influence of treatments on Mg content in soil as shown in Table 15. The highest Mg content was recorded in T₉ (40.7 mg/kg) which was on par with T₅, T₁₁, T₈, T₁₀ and T₁₃. T₇ recorded the lowest Mg content of 29.5 mg/kg.

4.5.9. Available S

The effect of treatment application on the available S content of soil is presented in Table 15. Available S content in the soil were statistically non significant, even then the available S content in treatment plots was higher than control plot. T₁₁ recorded with highest S content (20.8 mg/kg) while control T₁₃ recorded lowest S content of 8.7 mg/kg.

4.5.10. Available Zn

The effect of treatment application on available Zn content of soil is presented in Table 16. There was remarkable difference between the treatments. T₅ recorded the maximum Zn content of 8.9 mg/kg followed by T₆ (6.0 mg/kg) whereas control T₁₃ recorded with minimum Zn content of 0.4 mg/kg.

Table 15 Foliar application of micronutrient mixture on available secondary nutrients content of soil

Treatments	Available Ca (mg/kg)	Available Mg (mg/kg)	Available S (mg/kg)
T ₁	394 17	29 68	17 29
T ₂	351 67	33 98	9 79
T ₃	396 67	33 30	7 29
T ₄	501 67	32 78	10 54
T ₅	533 33	40 30	16 75
T ₆	419 50	34 42	7 50
T ₇	334 17	29 53	14 58
T ₈	323 33	38 63	12 50
T ₉	353 33	40 72	19 13
T ₁₀	346 67	37 13	11 88
T ₁₁	480 00	38 67	20 75
T ₁₂	375 00	32 28	10 63
T ₁₃	456 67	36 85	8 67
SEm (±)	14690 042	10 079	41 490
CD (0 05)	NS	5 350	NS

4.5.11. Available Fe

The treatments had notable influence on available Fe content of soil (Table 16). T₄ was recorded with highest Fe content of 43 3 mg/kg which was on par with T₃ (39 0 mg/kg) and T₅ (37 2 mg/kg) while control (T₁₃) recorded with lowest Fe content (23 2 mg/kg).

4.5.12. Available Mn

The data on available Mn content of soil is presented in Table 16. The highest Mn content was found in T₁₀ (11.4 mg/kg) which was statistically on par with T₁, T₅ and T₃ while T₄ was registered with lowest Mn content of 2.3 mg/kg. Control T₁₃ was with 4.5 mg/kg.

4.5.13. Available Cu

There was no considerable difference in available Cu content of soil between the treatments (Table 16). T₅ was recorded highest Cu content of 3.9 mg/kg while lowest was found in T₂ (2.8 mg/kg).

4.5.14. Available B

The data on available B content of soil is presented in Table 16. Significant difference was found between the treatments. The highest B content was recorded in T₄ (0.7 mg/kg) which was statistically on par with T₁₁, T₂ and T₇. All other treatments were on par with control. T₁₃ was found to have lowest B content of 0.2 mg/kg.

4.5.15. Available Mo

The data on available Mo content of soil is presented in Table 16. Significant difference was found between the treatments. The highest Mo content was recorded in T₁₂ (0.01 mg/kg) which was statistically on par with T₁₁ and T₇ whereas control T₁₃ was found to have the lowest Mo content of 0.002 mg/kg. T₁₃ was statistically on par with T₁, T₅, T₂, T₄ and T₃.

Table 16 Foliar application of micronutrient mixture on available micronutrients content of soil

Treatments	Zn (mg/kg)	Fe (mg/kg)	Mn (mg/kg)	Cu (mg/kg)	B (mg/kg)	Mo (mg/kg)
T ₁	2 61	32 47	10 05	3 16	0 29	0 002
T ₂	3 46	30 47	6 57	2 76	0 47	0 003
T ₃	4 59	39 03	8 56	3 15	0 37	0 004
T ₄	2 30	43 33	2 28	3 19	0 69	0 003
T ₅	8 91	37 17	8 89	3 87	0 31	0 002
T ₆	5 98	34 43	5 56	2 84	0 29	0 004
T ₇	1 91	30 80	3 07	3 20	0 47	0 009
T ₈	1 67	29 60	5 56	3 22	0 22	0 007
T ₉	0 64	28 47	3 94	3 06	0 24	0 008
T ₁₀	1 30	31 47	11 42	3 17	0 40	0 008
T ₁₁	2 60	32 47	6 69	3 19	0 52	0 010
T ₁₂	1 18	30 13	7 49	3 59	0 27	0 010
T ₁₃	0 38	23 23	4 45	3 49	0 19	0 002
SEm (±)	1 394	19 041	5 188	0 194	0 026	0 001
CD (0 05)	1 990	7 354	3 839	NS	0 272	0 002

4.6 NUTRIENT CONCENTRATIONS IN LEAF

Various chemical analysis of banana leaf samples was conducted in order to examine the effect of foliar spraying of micronutrient mixture on leaf nutrient status. Leaf samples were collected at bunching stage and at the time of harvest, dried in hot air oven, powdered and were analyzed using standard analytical procedures as described in materials and methods.

4.6.1. Nitrogen

The effect of treatment application on nitrogen content of banana leaf at bunching stage and at the time of harvest are presented in Table 17. At bunching stage, the treatments found to be non significant with respect to leaf N content. However, T₇ recorded highest N content of 4.2 per cent and T₁₂ recorded lowest N content (2.5 %) while control T₁₃ was found to have 2.8 per cent.

Leaf N content of banana at the time of harvest found to be non significant. However, T₁ and T₇ recorded highest N content of 3.2 per cent whereas T₁₂ and T₁₃ recorded lowest N content in leaves (2.1 %).

4.6.2. Phosphorus

The effect of treatment application on phosphorus content of banana leaf at bunching stage and at harvest is presented in Table 17. At bunching stage, the treatments found to be non significant with respect to leaf P content. T₁₁ recorded highest P content of 0.28 per cent whereas control T₁₃ recorded lowest P content in leaves (0.15 %).

There was notable difference among the treatments with respect to P content of leaf at the time harvest. T₃ and T₇ recorded highest P content of 0.31 per cent which was on par with T₁₁, T₁₀ and T₈. Lowest P content was found in T₁ (0.12 %) followed by T₁₃ (0.13 %).

Table 17 Foliar application of micronutrient mixture on N, P and K content of banana leaf

Treatments	N (%)		P (%)		K (%)	
	Bunching stage	At harvest	Bunching stage	At harvest	Bunching stage	At harvest
T ₁	3.15	3.15	0.25	0.12	3.70	2.50
T ₂	3.85	2.80	0.25	0.17	3.65	3.38
T ₃	3.50	2.45	0.23	0.31	3.85	3.25
T ₄	3.85	2.80	0.23	0.22	3.90	3.50
T ₅	3.15	2.45	0.17	0.17	3.75	3.13
T ₆	3.50	2.45	0.19	0.18	3.55	3.38
T ₇	4.20	3.15	0.21	0.31	3.84	3.38
T ₈	3.15	2.45	0.16	0.26	3.25	2.63
T ₉	3.15	2.45	0.18	0.18	3.75	3.75
T ₁₀	3.50	2.80	0.18	0.26	3.95	3.85
T ₁₁	3.85	2.80	0.28	0.28	4.50	3.88
T ₁₂	2.45	2.10	0.19	0.20	3.33	3.63
T ₁₃	2.80	2.10	0.15	0.13	2.04	1.90
SEM (±)	0.402	0.129	0.001	0.001	0.111	0.150
CD (0.05)	NS	NS	NS	0.076	0.727	0.843

4.6.3. Potassium

The effect of treatment application on potassium content of banana leaf at bunching stage and at harvest is presented in Table 17. At bunching stage, the

treatments had notable difference with respect to K content. T₁₁ recorded highest K content of 4.5 per cent which was on par with T₁₀, T₄, T₃ and T₇ whereas control T₁₃ was found to have lowest K content (2.1 %)

There was significant difference between the treatments with respect to K content of leaf at the time of harvest. T₁₁ recorded highest K content (3.9 %) whereas control T₁₃ recorded the lowest K content of 1.9 per cent. Except T₁₃, T₁ and T₈ all other treatments were on par with T₁₁.

4.6.4. Calcium

Leaf Ca content at bunching stage and at harvest showed notable difference among the treatments (Table 18). At bunching stage, T₈ recorded highest Ca content of 1.5 per cent which was on par with T₆, T₇ and T₅. The lowest Ca content was found in T₁ (0.7 %) while control T₁₃ was with 1.0 per cent.

T₁₁ recorded highest Ca content of 6.55 per cent at harvest whereas control T₁₃ recorded lowest Ca content of 2.2 per cent. Except T₁₁ and T₇ all other treatments were statistically on par with T₁₃. T₁₁ was on par with T₇ (6.4 %).

4.6.5. Magnesium

The effect of treatment application on magnesium content of banana leaf at bunching stage and at harvest is presented in Table 18. At bunching stage, significant difference was noticed among the treatments. The highest Mg content was observed in T₁₀ (0.25 %) which was on par with T₉, T₈, T₁₁, T₁₃ and T₁₂ except these treatments all other treatments were statistically on par with T₃. T₃ was found to have lowest Mg content of 0.12 per cent.

At the time of harvest, there was no appreciable difference found among the treatments with respect to Mg content of banana leaf. However, the highest Mg

content was registered in T₇ (0.36%) while control T₁₃ was found to have lowest Mg content of 0.10 per cent

Table 18 Foliar application of micronutrient mixture on Ca, Mg and S content of banana leaf

Treatments	Ca (%)		Mg (%)		S (%)	
	Bunching stage	At harvest	Bunching stage	At harvest	Bunching stage	At harvest
T ₁	0.70	2.84	0.13	0.21	0.14	0.10
T ₂	0.94	3.24	0.13	0.21	0.14	0.06
T ₃	1.01	3.58	0.12	0.22	0.09	0.06
T ₄	1.11	3.10	0.13	0.25	0.08	0.08
T ₅	1.20	3.35	0.13	0.21	0.15	0.08
T ₆	1.43	2.98	0.13	0.19	0.08	0.09
T ₇	1.35	6.44	0.19	0.36	0.12	0.25
T ₈	1.53	3.45	0.24	0.19	0.11	0.07
T ₉	1.15	3.16	0.24	0.17	0.18	0.14
T ₁₀	0.73	3.66	0.25	0.15	0.13	0.18
T ₁₁	0.77	6.55	0.24	0.15	0.20	0.29
T ₁₂	0.98	3.19	0.22	0.19	0.22	0.13
T ₁₃	1.00	2.24	0.23	0.10	0.07	0.05
SEm (±)	0.022	0.654	0.001	0.004	0.002	0.03
CD (0.05)	0.325	1.762	0.052	NS	NS	0.121

4.6.6. Sulphur

The effect of treatment S application on sulphur content of banana leaf at bunching stage and at harvest are presented in Table 18. At bunching stage, there were no significant differences found among the treatments. T₁₂ recorded highest S content of 0.22 per cent while control T₁₃ recorded lowest S content in leaves (0.07 %).

Leaf S content showed appreciable differences among the treatments at the time of harvest. The highest S content of leaf was found in T₁₁ with 0.29 per cent whereas lowest was recorded in control T₁₃ (0.05 %). T₁₁ was statistically on par with T₇ (0.25 %) and T₁₀ (0.18 %).

4.6.7. Zinc

The effect of foliar spraying of micronutrient mixture on the Zn content of leaf at bunching stage and at harvest are presented in Table 19. There was significant difference among the treatments in both stages of observation. At bunching stage, T₁₂ was recorded highest Zn content of 101.0 ppm followed by T₁₁, T₁₀ and T₉. The minimum was found in T₁₃ (9.0 ppm) which was statistically on par with T₁, T₂, T₃ and T₄.

At the time of harvest, T₁₁ found to have highest Zn content of 95.7 ppm which was on par with T₁₀ (83.3 ppm) whereas control (T₁₃) was found to have lowest Zn content of 3.8 ppm which was on par with T₁, T₂, T₃ and T₄.

4.6.8. Iron

The effect of foliar spraying of micronutrient mixture on Fe content of banana leaf at bunching stage and at harvest are presented in Table 19. Fe content of the leaves was significantly influenced by treatment application at bunching stage. The

highest Fe was recorded in T₁₂ with 415 0 ppm which was statistically on par with T₇ (383 8 ppm) whereas control T₁₃ recorded lowest Fe content of 75 0 ppm

Leaf Fe content showed notable differences among the treatments at the time of harvest. Highest Fe content was found in T₁₂ (945 0 ppm) which was statistically on par with T₄ and T₉. Lowest Fe content was found in T₁₃ (398 5 ppm)

Table 19 Foliar application of micronutrient mixture on Zn, Fe and Mn content of banana leaf

Treatments	Zn (ppm)		Fe (ppm)		Mn (ppm)	
	Bunching stage	At harvest	Bunching stage	At harvest	Bunching stage	At harvest
T ₁	11 40	5 79	115 00	762 50	2425 00	2175 00
T ₂	11 55	11 25	112 50	625 00	2525 00	2550 00
T ₃	16 00	14 00	104 75	575 00	2675 00	2525 00
T ₄	22 25	14 10	175 60	915 00	3075 00	3250 00
T ₅	37 23	36 50	147 50	500 00	2675 00	3575 00
T ₆	32 45	27 48	307 25	745 00	3240 00	2050 00
T ₇	25 75	28 25	383 75	765 00	3200 00	3560 00
T ₈	40 65	32 75	175 00	645 00	3625 00	3775 00
T ₉	62 75	70 03	115 00	887 50	3850 00	3375 00
T ₁₀	80 13	83 30	227 75	837 50	3825 00	3800 00
T ₁₁	82 75	95 74	227 00	695 00	3925 00	3145 00
T ₁₂	101 00	48 05	415 00	945 00	4550 00	3300 00
T ₁₃	9 00	3 78	75 00	398 50	2450 00	2275 00
SEM (±)	45 436	33 889				
CD (0 05)	14 688	12 685	47 101	79 646	188 271	174 834

4.6.9. Manganese

The effect of foliar spraying of micronutrient mixture on Mn content of banana leaf at bunching stage and at harvest are presented in Table 19. Mn content of the leaves was significantly influenced by treatment application at bunching stage. The highest Mn was found in T₁₂ with 4550 ppm followed by T₁₁, T₉ and T₁₀ whereas T₁ was found to have lowest Mn content of 2425 ppm which was statistically on par with control T₁₃ (2450 ppm) and T₂ (2525 ppm).

At the time of harvest also treatment application resulted in appreciable difference among the treatments with respect to Mn content of leaves. Highest Mn content was found in T₁₀ with 3800 ppm which was on par with T₈ (3775 ppm) whereas T₆ recorded lowest Mn content of 2050 ppm.

4.6.10. Copper

The effect of foliar spraying of micronutrient mixture on Cu content of banana leaf at bunching stage and at harvest are presented in Table 20. There was significant difference among the treatments with respect to Cu content of leaves at bunching stage. Highest Cu content was found in T₁₁ with 129.7 ppm followed by T₁₂ and T₁₀ whereas T₁ recorded lowest Cu content of 22.2 ppm. Except T₁₁, T₁₂, T₁₀ and T₉, all other treatments were statistically on par with T₁.

At the time of harvest also significant difference found among the treatments with respect to Cu content of banana leaf. T₁₁ recorded highest Cu content of 105.3 ppm and lowest was found in T₂ with 18.2 ppm. T₁₁ was statistically on par with T₁₂, T₄ and T₁₀.

4.6.11. Boron

The effect of foliar spraying of micronutrient mixture on B content of banana leaf at bunching stage and at harvest are presented in Table 20. At bunching stage,

significant difference was found between the treatments with respect to B content of leaf. Highest B content recorded in T₃ with 60.0 ppm which was on par with T₆, T₄, T₇ and T₅ except these treatments all others were on par with T₁₂. Lowest B content was found in T₁₂ (12.0 ppm) while control T₁₃ was with 20.0 ppm.

Leaf B content showed significant difference among the treatments at harvest. Highest B content was found in T₁₂ with 90 ppm which was on par with T₁ (82.5 ppm) whereas T₈ recorded lowest B content of 30 ppm. Control T₁₃ was found to have 47.5 ppm.

4.6.12. Molybdenum

The effect of foliar spraying of micronutrient mixture on Mo content of banana leaf at bunching stage and at harvest are presented in Table 20. At bunching stage, significant difference was found between the treatments with respect to Mo content of leaf. Highest Mo content was found in T₁₁ and T₁₂ with 2.2 ppm which was on par with T₁₀ whereas control T₁₃ was found to have the lowest Mo content of 0.5 ppm. T₁₃ was statistically on par with T₁, T₂ and T₃.

Leaf Mo content showed significant difference among the treatments at the time of harvest. Highest Mo content was found in T₁₁ with 2.1 ppm which was on par with T₁₂ and T₁₀ whereas control T₁₃ was found to have the lowest Mo content of 0.4 ppm. T₁₃ was statistically on par with T₁, T₂, T₃ and T₄.

4.7 ECONOMIC ANALYSIS

The data on economics of banana cultivation are given in Table 21. The treatment T₁₁ registered the maximum net returns of Rs 6,94,964 ha⁻¹ and it was on par with T₁₀, T₉, T₆ and T₇ whereas control T₁₃ recorded the minimum net returns of Rs 3,53,251 ha⁻¹.

With regard to benefit-cost ratio, the treatment T₁₁ (3 sprays of 3 % micronutrient mixture) recorded the maximum value of 2.3 and was on par with T₁₀, T₉, T₆ and T₇. The lowest Benefit / Cost ratio of 1.6 was recorded by control T₁₃.

Table 20 Foliar application of micronutrient mixture on Cu, B and Mo content of banana leaf

Treatments	Cu (ppm)		B (ppm)		Mo (ppm)	
	Bunching stage	At harvest	Bunching stage	At harvest	Bunching stage	At harvest
T ₁	22.20	21.75	15.50	82.50	0.65	0.60
T ₂	24.50	18.15	29.50	60.00	0.70	0.60
T ₃	29.00	23.65	60.00	60.00	0.75	0.68
T ₄	26.60	82.65	49.50	42.50	0.93	0.88
T ₅	32.70	61.85	47.00	55.00	1.35	1.28
T ₆	34.00	31.13	52.50	32.50	1.45	1.40
T ₇	45.28	41.15	49.50	47.50	1.40	1.34
T ₈	54.12	47.89	18.34	30.00	1.25	1.18
T ₉	61.44	50.38	28.50	32.50	1.65	1.47
T ₁₀	68.60	75.59	35.00	55.00	1.75	1.68
T ₁₁	129.71	105.25	12.50	77.50	2.15	2.08
T ₁₂	68.93	86.03	12.00	90.00	2.15	2.02
T ₁₃	23.60	22.75	20.00	47.50	0.45	0.40
SEm (±)	252.111	278.782	206.010	31.545	0.038	0.052
CD (0.05)	34.598	36.382	31.275	12.238	0.426	0.0499

Table 21 Foliar application of micronutrient mixture on economics of banana cultivation

Treatments	Cost of cultivation (Rs ha ⁻¹)	Gross returns (Rs ha ⁻¹)	Net returns (Rs ha ⁻¹)	BCR
T ₁	537180	1032396	493295 913	1 770
T ₂	537255	1002917	467770 667	1 870
T ₃	537930	956319 4	414357 593	1 918
T ₄	538305	1023528	485357 515	1 901
T ₅	537615	1115083	588485 481	2 094
T ₆	538365	1177194	637435 920	2 184
T ₇	539115	1176042	693185 667	2 286
T ₈	539865	1154583	616097 512	2 141
T ₉	537877 5	1184653	647935 324	2 190
T ₁₀	538815	1216944	641210 987	2 204
T ₁₁	539752 5	1230375	694963 583	2 287
T ₁₂	530690	1087458	546210 765	2 010
T ₁₃	536775	834041 7	353250 657	1 558
SEm (±)			-	0 022
CD (0 05)			128028 716	0 252

Cost of inputs

Nitrogen – Rs 12 kg⁻¹

Phosphorus – Rs 34 kg⁻¹

Potassium – Rs 16 kg⁻¹

FYM – Rs 800 t⁻¹

Lime – Rs 10 kg⁻¹

Calcium	–Rs 9 55 kg ¹
Magnesium	–Rs 70 kg ¹
Cost of planting material	–Rs 20 plant ¹
Solution A	–Rs 150 / litre
Solution B	–Rs 4 / litre
Labour cost	–Rs 600/day

Price of produce

Fruits	–	Rs 35 kg ¹
Malbud	–	Rs 10 kg ¹
Suckers	–	Rs 5 of each

Discussion

5. DISCUSSION

Discussion of the results on investigation carried out at College of Agriculture, Padannakkad and Regional Agricultural Research Station farm, Nileshwar to formulate micronutrient mixture for foliar spray and to study the effect of micronutrient mixture on growth, yield and quality of fruits of tissue culture banana cv Nendran are presented in this chapter. The whole investigation comprised of preparation of micronutrient mixture as liquid formulation, hardening studies of tissue culture banana plants using micronutrient mixture and studying the effect of micronutrient mixture on growth, yield and quality of banana.

5.1 MICRONUTRIENT MIXTURE PREPARATION

Preparation of micronutrient mixture as a liquid formulation was tried with the use of different micronutrient salts containing Zn, B, Fe, Mn, Cu and Mo. The formulation technique was standardized and it contains two liquids: Solution A and Solution B. Solution A consisting of $ZnSO_4 \cdot 7H_2O$, $CuSO_4 \cdot 5H_2O$, $FeSO_4 \cdot 7H_2O$, H_3BO_3 , $MnSO_4 \cdot H_2O$ and $(NH_4)_6Mo_7O_{24} \cdot 4H_2O$ in desired proportion. For maintaining stability and keeping quality of solution A, various chemicals were tried. Citric acid (20 %) was found best for maintaining keeping quality of solution A which acts as an excellent chelating agent. Its stability was periodically monitored at specific intervals and it was found that solution A with 30 ml/litre of citric acid (20 %) can be kept as such at ambient conditions upto 1 year. Solution B is prepared as 1 per cent humic acid. Humic acid might benefit plant growth by chelating unavailable nutrients and buffering pH. Required quantity of this has to be blended with diluted solution A just before spraying. The pH of micronutrient mixture was noted just before spraying and it was observed 5.56 at the desired dilution. Humic acid plays a key role both in enhancing the bioavailability of nutrients to plants and in reducing the adverse effect of some of the free ions due to its chelate forming capability (Bloom *et al*, 1979).

Muthukrishnan and Velu (2011) developed and evaluated a micronutrient formulation for sugarcane and concluded that 92.5 kg micronutrient mixture formulation II as enriched farm yard manure containing 50 kg FeSO₄, 5 kg MnSO₄, 30 kg ZnSO₄, 2.5 kg CuSO₄ and 5 kg of borax may be recommended for sugarcane for maximizing yield with higher B:C ratio. Increased cane yield with higher levels of micronutrient mixture may be attributed to the complementary and supplementary effect of fertilizers and FYM.

A multi-micronutrient mixture having a composition of Zn (9.5 %) + B (2.6 %) + Cu (1.2%) + Mg (2.4 %) + N (0.46 %) was developed and found that 0.5 per cent of multi-micronutrient mixture enhanced the growth and yield of okra variety Varsha Uphar (Mim, 2015).

5.2 HARDENING STUDIES

Foliar application of micronutrient mixture significantly increased the plant growth characteristics of banana such as plant height, number of leaves, leaf length and leaf breadth (Table 4). This might be due to the beneficial effect of micronutrient on secondary hardening of tissue culture banana since their widespread deficiencies and their relatively low content in soil had been reported (KSPB, 2013).

Increase in growth with foliar fertilization might also be due to the fact that, foliar application of nutrients is readily absorbed by leaves and it enhances the physiological processes. Foliar spraying of 0.25 per cent and 0.5 per cent of micronutrient mixture as three sprays at immediately after planting, two weeks and three weeks after planting significantly enhanced all the biometric characters of tissue culture banana cv Nendran.

Potarzycki and Grzebiusz (2009) reported that zinc exerts a great influence on basic plant life processes, such as (i) nitrogen metabolism- uptake of nitrogen and protein quality (ii) photosynthesis- chlorophyll synthesis which ultimately improved

the number of leaves. Boron influences cell development and elongation. It is involved in the transport of sugars across cell membranes and in the synthesis of cell wall material. Boron also regulates the carbohydrate metabolism in plants and plays a role in amino acid formation and synthesis of proteins. Boorboori *et al* (2012) stated that, Iron (Fe) is another micronutrient that is a co-factor for approximately 140 enzymes that catalyze unique biochemical reactions. Hence, iron has many essential roles in plant growth and development including chlorophyll synthesis, thylakoid synthesis and chloroplast development. Manganese plays an important role in chlorophyll production and its presence is essential in Photo-system II, also involved in cell division and plant growth. Copper helps in the utilization of iron during chlorophyll synthesis. Mo increased the metabolic pools required for the synthesis of saccharides, along with the enhanced photosynthetic capacity.

Earlier, Vasane and Kothari (2006 & 2008) also reported enhanced growth traits such as plant height, girth, leaf length, leaf width and number of leaves were obtained with foliar application of multiplex (commercial micronutrient formulation containing Zn - 3 %, Fe - 2.5 %, Cu - 1 %, Mn - 1 %, B - 0.5 % and Mo - 0.1 %) during secondary hardening in banana cv. Grand Naine.

Increased plant growth characters by the application of micronutrients may be due to their involvement in chlorophyll formation, which might have helped to favour cell division, synthesis of hormone and proteins, enzymes activity, meristematic activity in apical tissue, expansion of cell and formation of new cell wall (Singh and Maurya, 1979, Havlin *et al*, 2014).

5.3 FIELD EXPERIMENT

Tissue culture hardened plants were planted out in the field and various levels of micronutrient mixture were sprayed on foliage at 2, 4 and 6 months after planting and evaluated. It was found that foliar application of micronutrient mixture at concentrations 1, 2, 3 and 4 per cent had significantly enhanced the vegetative

characters like pseudostem height, pseudostem girth and number of leaves at 4 months after planting and at the time of shooting and yield characteristics like days to harvest, bunch weight, number of fingers, average weight of fingers and finger length produced. Fruit quality parameters like acidity, TSS, total sugars, reducing sugars, non-reducing sugars, pulp to peel ratio, sugar / acid ratio, TSS / acid ratio, per cent loss in weight during ripening, keeping quality of the fruits at ambient condition also exhibited significant difference among the treatments. The quality parameters of fruits were better in plants receiving the micronutrient mixture as foliar application. This might be due to the beneficial effect of micronutrients on growth, yield and quality of banana.

5.3.1. Effect of treatments on plant growth parameters

The effect of micronutrient mixture on pseudostem height, pseudostem girth and number of leaves were found to be significant. It shows that micronutrients could make substantial effect on the above growth parameters of banana cv Nendran. Improvement in growth of banana plant might be due to enhancement of photosynthetic and other metabolic activities which lead to an increase in various plant metabolites responsible for cell division and cell elongation.

Foliar application of 4 per cent micronutrient mixture as three sprays increased the pseudostem height and girth of banana at 4 months after planting. At the time of shooting, three sprays of 2 per cent micronutrient mixture as enhanced the pseudostem girth. Ghanta and Mitra (1993) found that foliar spraying of 0.5 per cent $ZnSO_4$, 0.1 per cent $CuSO_4$, 0.2 per cent H_3BO_3 and 0.05 per cent $(NH_4)_2 MoO_4$ at 3 and 5 months after planting enhanced the plant height of banana. Anonymous (2005) reported the beneficial effect of micronutrients on height of pseudostem at shooting. Mandal *et al* (2002) observed that the beneficial effect of micronutrients and their combination on pseudostem girth over control in banana cv Giant Governor.

The beneficial effect of micronutrients on height and girth might be due to micronutrients especially Zn which is the activator of the enzymes involved in protein synthesis and had direct effect on the level of IAA in plants (Ram and Bose, 2000), Copper activates several enzymes in plants and helps chlorophyll synthesis and involve in carbohydrate and protein metabolism (Ram and Bose, 2000) and Boron increases photosynthetic activity and respiration in plants and thus improves the growth (Lal and Rao, 1954)

Foliar spraying of 2 per cent micronutrient mixture as two sprays significantly increased the number of leaves of banana at 4 months after planting and at the time of shooting. The synthesis and transport of plant assimilates to the developing banana fruit is greatly affected by the retention of green leaves after the flowering stage, especially when assimilate flow from other plant parts becomes limiting (Kumar and Kumar, 2007). Increase in number of leaves may be due to Zn stimulates photosynthetic activity and its presence is important for protein synthesis which in turn can enhance the rate of leaf production.

These results are in harmony with those obtained by Ghanta and Mitra (1993) in banana cv Giant Governor, Haque *et al* (2000) in mandarin orange, Babu and Singh (2002) in Itchi, Das and Mohan (1993) in banana cvs Chenichampa, Jahaji and Barjahaji, Anjali *et al* (2013) in banana cv Grand Name, Yadav and Patel (2013) in banana cv Grand Name, Yadlod and Kadam (2008c) in banana cv Ardhapuri, Pathak *et al* (2011) in banana cv Martaman (AAB, Silk), Jeyabaskaran and Pandey (2008) in Karpuravalli banana, Mandal *et al* (2002) in banana cv Giant Governor, Subramanian and Pillai (1997) and Kumar and Jayakumar (2001) in banana.

In the present study, foliar spraying of micronutrient mixture in banana showed no significant influence on number of suckers at harvest. Even though, the higher number of suckers was produced in plants treated with three times foliar

spraying of 2 per cent and 4 per cent micronutrient mixture which was superior than control. Similar result was observed in banana cv Grand Name (Yadav *et al* , 2010)

Increased in vegetative growth of banana due to foliar application of 2 and 4 per cent of micronutrient mixture sprays might be attributed to their stimulatory effect on plant metabolism. These results corroborate the findings of Hada *et al* (2014) in guava

5.3.2. Effect of treatments on yield characteristics

Yield of banana is a function of bunch weight and number of plants per hectare. Hence, any nutrient management study should aim at producing maximum bunch weight, so that, the productivity could be enhanced reasonably (Kumar and Kumar, 2008)

In the present study, foliar application of micronutrient mixture in tissue culture banana cv Nendran has shown positive and significant influence in the yield characteristics of the crop. Increase in yield may be attributed to important role of micronutrients in enhancing cell elongation and division, photosynthetic activity and increased production and accumulation of carbohydrates (Abdel-Kader *et al* , 1992). Treatment application has significantly increased the weight of male bud, bunch weight, number of fingers, average weight of fingers and finger length, which are the traits of paramount importance from the economic point of view.

Bunch weight in treated plants was found to be superior to that of control. Figure 2 shows the effect of different treatment levels and time of application on bunch weight of banana. It was also found that with increasing level of micronutrient mixture and time of application from one sprays to two or three sprays resulted in improvement of bunch weight because of their specific role in improving bunch weight. At lower level and time of application the bunch weight of fruits were decreased because of its lower concentration it might not be sufficient for enhancing

yield attributes of banana. Three sprays of 3 per cent micronutrient mixture enhanced the bunch weight of banana. This may be due to the provision of micronutrients at latter stages of crop, which might have enhanced accumulation of assimilates in fruits and thus resulting in heavier bunch weight. The possible reason for increase in bunch weight by the micronutrients might be due to faster loading and mobilization of photo-assimilates to fruits and involvement in cell division and cell expansion which ultimately reflected into more weight of fruit in treated plants.

There was no difference between the treatments with respect to number of hands produced per bunch. Similar finding was reported by Anjali *et al* (2013) in banana cv Grand Name with foliar spraying of micronutrients. The maximum hands were noted with 2 per cent and 3 per cent micronutrient mixture as foliar sprayed at two and three times, which were on par.

Number of fingers per bunch was increased with three sprays of 1 per cent micronutrient mixture. Finger length was significantly influenced by treatment application as compared to that of control and the maximum finger length was recorded in 3 sprays of 3 per cent micronutrient mixture. Average weight of finger was found to be enhanced with three sprays of 3 per cent micronutrient mixture. These results are in line with those obtained by Yadlod and Kadam (2008a) in banana cv Grand Name, Pathak *et al* (2011) in banana cv Martaman, Suresh and Savithri (2001) in Nendran banana. Decrease in average fruit weight may be due to increasing finger number and yield which means the partition of assimilates into a high number of fruit and consequently reduction in individual finger weight and other related characters.

Two or three time foliar spraying of micronutrient mixture as 2 per cent and 3 per cent might have helped the crop to increase in chlorophyll content of leaves, photosynthetic efficiency, translocation of metabolites from the source to sink as and

when needed by the crop and it may be responsible for retaining more number of fruits, increase in weight of fruit and productivity as compared to control

The most outstanding effect of micronutrients on yield was reported by Ghanta and Mitra (1993) in banana cv Giant Governor, Suresh and Savithri (2001) in Nendran banana, Yadlod and Kadam (2008b) in banana cv Shrimanti, Haque *et al* (2000) in mandarin, Modi *et al* (2012) in papaya cv Madhu Bindu, Yadav and Patel (2013), Ghanta and Dwivedi (1993), Subramanian and Pillai (1997), Kumar and Jayakumar (2001) in banana, Dutta and Dhua (2002), Dutta (2004) in mango, Gurjar *et al* (2015) in Alphonso mango, Pathak *et al* (2011) in banana cv Martaman (AAB, Silk), Jeyabaskaran and Pandey (2008) in Karpuravalli banana, Yadav *et al* (2010) and Anjali *et al.* (2013) in banana cv Grand Naine

There was significant difference among different treatments regarding total crop duration. Total crop duration was significantly altered by the application of micronutrient mixture 1 per cent, 2 per cent and 3 per cent as three sprays. Minimum days required from planting to harvesting observed in three times spraying of 3 per cent micronutrient mixture as compared to rest of the treatments. Bunch maturity period was found to be minimum in 3 sprays of 3 per cent micronutrient mixture and maximum in control. This might be due to reduced flowering and maturity duration which could be attributed to enhancing effect of zinc in enzymatic reaction, cell division as well as in growth (Supriya and Bhattacharyya, 1993, Yadav and Patel, 2013). Similar findings were noted by Ghanta and Mitra (1993) in banana cv Giant Governor, Yadav *et al* (2010) banana cv Grand Naine, Babu and Singh (2002) in mandarin orange, Yadlod and Kadam (2008c) in banana cv Ardhapuri

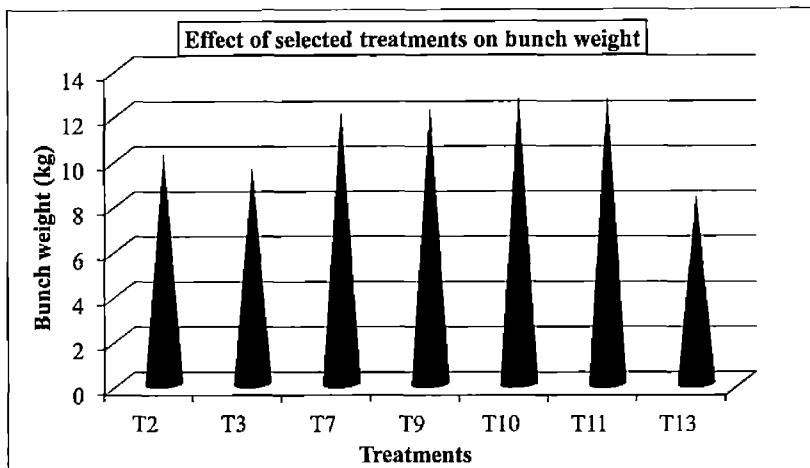


Figure 2 Foliar application of micronutrient mixture on bunch weight of banana

5.3.3. Effect of treatments on fruit characteristics

Quality standards have become most important factor for determining monetary yield as well as farmer's income in case of high value crops like banana. Any management system should aim to produce quality fruits, besides maximizing productivity (Kumar and Kumar, 2007). Hence, the present investigation reveals that all the quality parameters like titrable acidity, TSS, total sugars, pulp to peel ratio, reducing sugars, non-reducing sugars, sugar / acid ratio, TSS / acid ratio, per cent loss in weight during ripening and keeping quality of the fruits at ambient condition were significantly improved with the foliar application of micronutrients mostly at 1, 2 and 3 per cent in three sprays. Increasing the micronutrient mixture levels and time of application might stimulate and activate the biochemical processes in plant, which might improve the fruit quality.

Zn and Mn elements have main role in synthesis of proteins, enzyme activation, oxidation and revival reactions, improved the auxin content and

metabolism of carbohydrates. Zinc helped other enzymatic reaction like transformation of carbohydrates, activity of hexokinase and formation of cellulose and change in sugar considered due to its action on zymohexose (Dutta and Dhua, 2002). By utilizing of fertilizers contain above elements, performance on quality of crops is increasing and with shortage of this elements due to decline in plant photosynthesis and destroy RNA, amount of solution carbohydrates and synthesis of protein decreased and then performance and quality of crop will be decreased.

Treatment application of micronutrients exhibited significant differences in the titrable acidity percentage of the fruits. Figure 3 shows the effect of treatment application on titrable acidity of ripened banana fruits. Highest acidity percentage was found in control plants and lowest acidity was found in 3 sprays of 1, 2 and 3 per cent of micronutrient mixture. A reduction in acidity is usually preferred and here it is found that the treatments have reduced the acidity percentage of fruits as compared to that of control. With increasing level and time of application of micronutrient mixture the titrable acidity per cent has been reduced and in case of control it has been increased greatly. The reduction of acidity in micronutrient treated fruit juice might be due to their utilization in respiration and rapid metabolic transformation of organic acids into sugars (Brahmachari *et al*, 1997, Ningavva *et al*, 2014). Similar results were also reported by Deolankar and Frake (2001) in banana, Singh *et al* (2003) in pomegranate, Patil and Hiwarale (2004) in acid lime.

Total soluble solids (TSS) of fruits exhibited significant differences among the treatments. Figure 4 shows the effect of different levels and times of application of nutrients on TSS content of fruits. At lower concentration the TSS content was less, whereas with increasing level and time of application of micronutrient mixture, TSS of fruits also increased. This is because of the micronutrients were needed in higher quantity for enhancing fruit quality parameters. Three sprays of 2 or 3 per cent micronutrient mixture have shown to improve the TSS of ripened fruits. The higher TSS was due to the enhanced total sugar content, owing to the increased

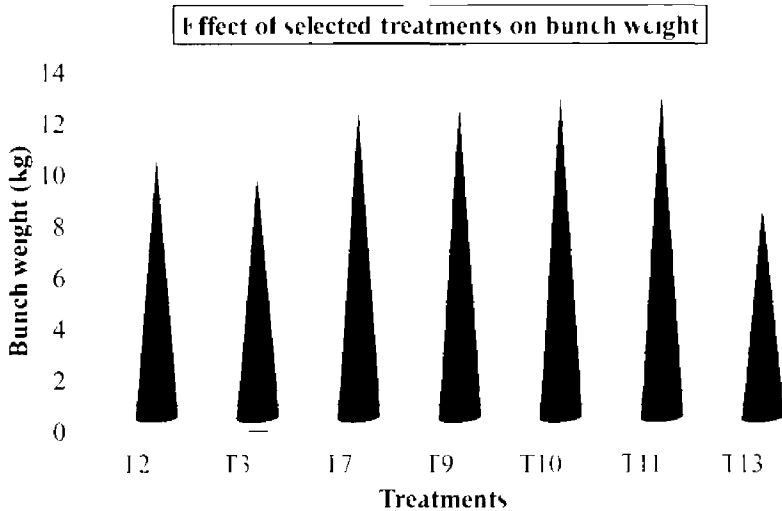


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photosynthesis activity, which ultimately leads to efficient translocation of available photosynthates to fruit pulp rather than to other parts (Kulkarni, 2004). Borate ion may be associated with the cell membrane where it could be complex with sugar molecules and facilitates its passage across the membrane that might be the reason of increased total soluble solids (Meena *et al*, 2006) which confirmed the present findings.

Figure 5 shows the effect of different levels and times of application of nutrients on pulp to peel ratio. The maximum amount of pulp content was recorded in 3 sprays of 1, 3 or 4 per cent micronutrient mixture which might be due to foliar application of micronutrients especially B. This might have made rapid synthesis of metabolites particularly carbohydrates and their translocation to the fruits causing relatively greater pulp content. Higher pulp weight and lower peel weight might be due to increasing levels and times of application of micronutrient mixture. This can also be possible as the complexes of polyhydroxy compounds with B can facilitate transport of carbohydrates within phloem tissue (Hewitt, 1963). These are in agreement with those findings reported by Pathak *et al* (2011) in banana cv Martaman, Yadlod and Kadam (2008a) and Ningavva *et al* (2014) in banana cv Grand Naine.

Application of micronutrients found to bring beneficial effect on total sugars, reducing sugars and non-reducing sugar content of ripened banana fruit, sugar / acid ratio, TSS / acid ratio as compared to that of control. Figure 6 shows that effect of treatment application on total sugar content of fruits. With increasing levels and times of application of micronutrient mixture the total sugar content also increased but it decreased in 3 sprays of 4 per cent micronutrient mixture because of their toxicity effect. Total sugar content of ripened banana fruits was found to be maximum in 3 sprays of 2 per cent micronutrient mixture. This is due to its action on converting complex substances into simple ones, which enhances the metabolic activity in fruits and it results in increased total sugar of fruit. Similar findings were

reported by Paul and Nair (2015) in banana cv Nendran banana, Pathak *et al* (2011) in banana cv Martaman, Yadlod and Kadam (2008b) in banana cv Shrumanti. The highest sugar content in juice of sweet orange fruits was observed with foliar application of ZnSO_4 (0.5 %) + FeSO_4 (0.4 %) + Borax (0.2 %) as reported by Kulkarni (2004).

Increased in reducing sugars with 3 sprays of 2 per cent micronutrient mixture might be due to that formation and translocation of carbohydrate, which improves the fruit quality as reported by Pathak and Mitra (2008). Figure 7 shows the effect of treatment application on reducing sugar content of ripened banana fruits. With increasing levels and times of application of micronutrient mixture, the reducing sugar content of ripened fruits also increased because of their necessity in fruit quality attributes. At lower level and time of application the reducing sugar content of fruits decreased because its lower concentration may not be enough for enhancing fruit quality attributes. The multi-micronutrient mixture comprised of Zn and B and other micronutrients have an important role in sugar metabolism. Zinc promotes hydrolysis of starch into sugars and B expedites sugar transport across the membrane by a temporary formation of sugar borate complex (Gauch and Dugger, 1953). Ghanta and Dwivedi (1993) reported the significant effect of micronutrient on reducing sugar content of fruit and the best result (6.99 %) in combined spraying of Zn+Cu+B. Similar results were also reported by Aziz and Wahab (1970), Sharma (1976), Paul and Nair (2015) in banana cv Nendran, Yadlod and Kadam (2008a) in banana cv Grand Name.

The highest non-reducing sugar content was observed in 2 sprays of 1 per cent micronutrient mixture and lowest was found in 3 sprays of 3 per cent micronutrient mixture. Paul and Nair (2015) also found that foliar spraying of micronutrient mixture (ZnSO_4 - 1.0 % + FeSO_4 - 0.3 % + CuSO_4 - 0.2 % + H_3BO_3 - 0.2 % + $(\text{NH}_4)_2\text{MoO}_4$ - 0.03 %) reduced the non-reducing sugar content of banana cv Nendran.

Foliar application of micronutrient mixture 2 and 3 per cent as three sprays significantly increased the sugar / acid ratio and TSS / acid ratio of ripened banana fruit. Figure 8 and Figure 9 shows the effect of treatment application on sugar / acid ratio and TSS / acid ratio of ripened banana fruits. It might be due to increase in sugar content and decrease in acidity level of fruits by these treatments. With increasing levels and times of application of micronutrient mixture the sugar content of ripened fruits has improved greatly and acidity percent has been decreased. But in case of lower levels the sugar content decreased and acidity per cent increased which led to reduction in sugar / acid ratio and TSS / acid ratio of ripened banana fruit. Similar beneficial effect of foliar application of Zn, Mn, B, Cu and Fe on mango, orange, banana and pineapple fruits have been reported by Singh and Rajput (1976) and Nehete *et al* (2011) in mango, Patel *et al* (2010) in banana cv Basrai, Pathak *et al* (2011) in banana cv Martaman, Kavitha *et al* (2000a) in papaya, Yadav and Patel (2013) in banana cv Grand Naine.

Per cent loss in weight during ripening and keeping quality of fruits at ambient condition exhibited significant differences among the treatments. Figure 10 shows the effect of foliar application of micronutrient mixture on per cent loss in weight of fruits during ripening at ambient condition. Foliar spraying of 2 or 3 per cent micronutrient mixture as 3 sprays shown to improve shelf life of ripened fruit as well as minimum loss in weight during ripening. With increasing levels and times of application of micronutrient mixture loss in weight during ripening of fruits were reduced. Reduction in weight loss might be due to decreased rate of respiration and transpiration, restricting ethylene accumulation and production in fruits during ripening. Enhanced shelf life of banana fruits with the application of micronutrient mixture due to their ability to improve the quality of harvested produce. These results are in harmony with those obtained by Paul and Nair (2015) in banana cv Nendran, Yadlod and Kadam (2008 b,c) in banana cv Shrimanti and Ardhapuri.

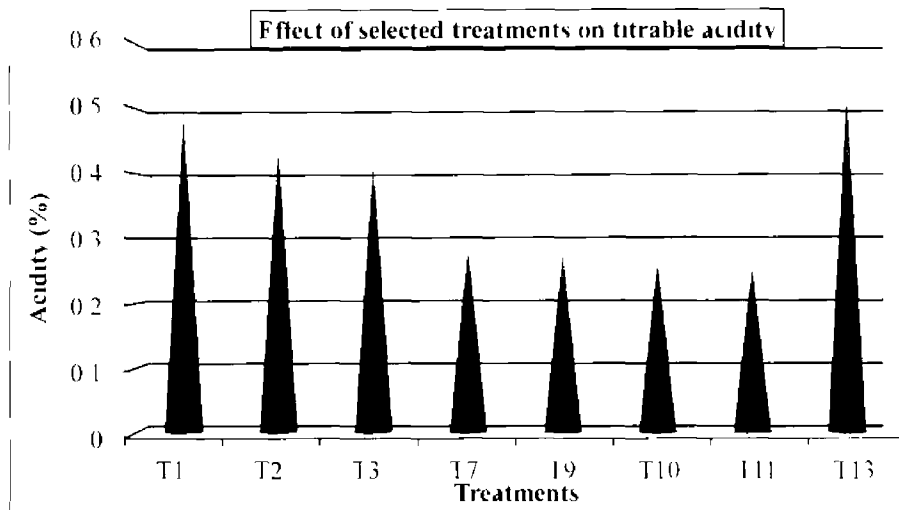


Figure 3 Foliar application of micronutrient mixture on titrable acidity of ripened banana fruits

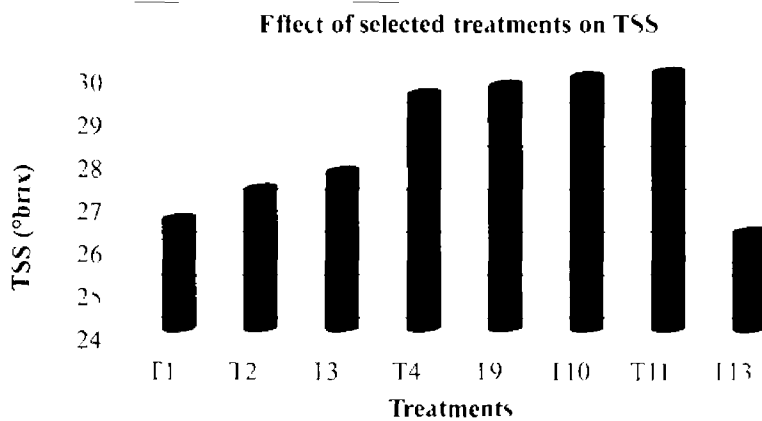


Figure 4 Foliar application of micronutrient mixture on TSS of ripened banana fruits

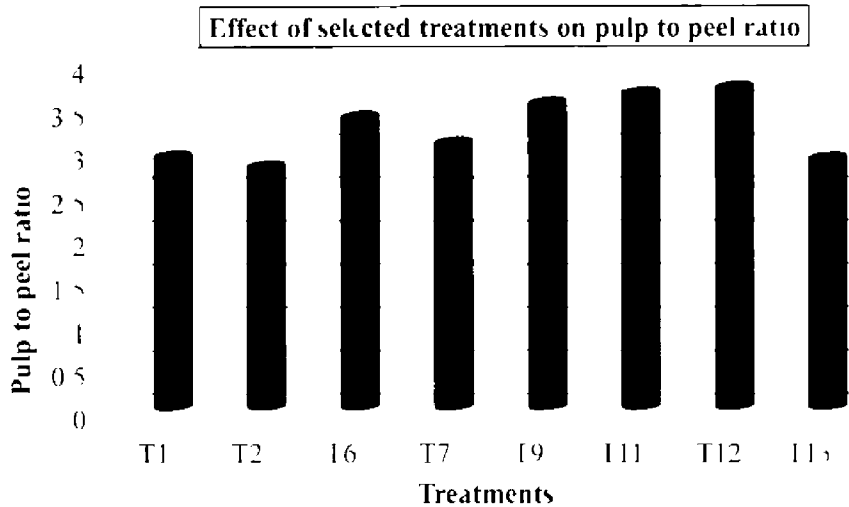


Figure 5 Foliar application of micronutrient mixture on pulp to peel ratio ratio of ripened banana fruits

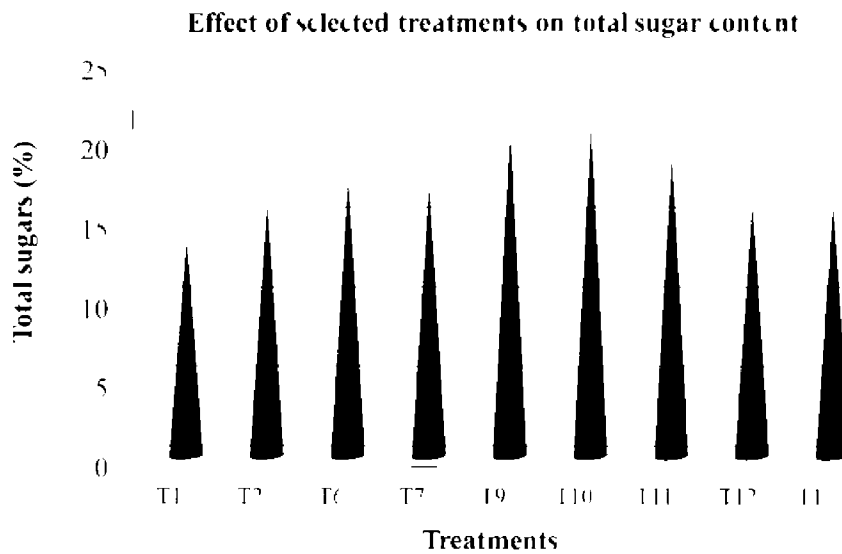


Figure 6 Foliar application of micronutrient mixture on total sugar content of ripened banana fruits

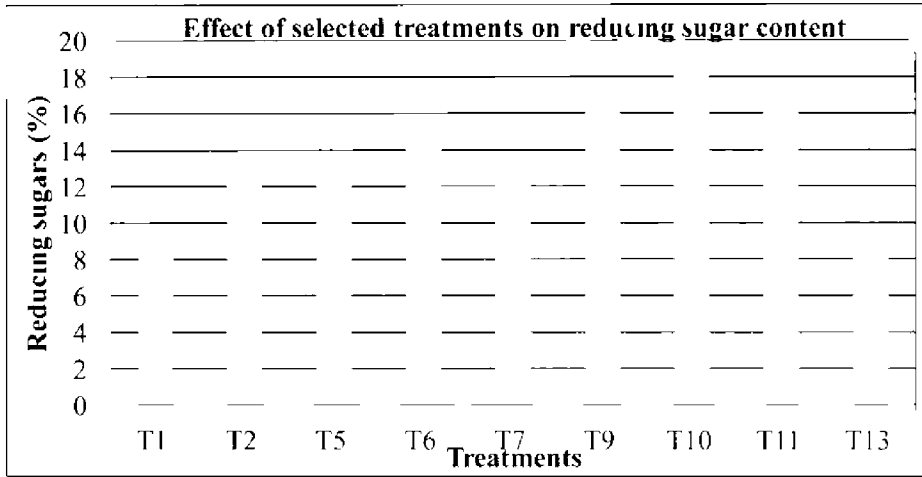


Figure 7 Foliar application of micronutrient mixture on reducing sugar content of ripened banana fruits

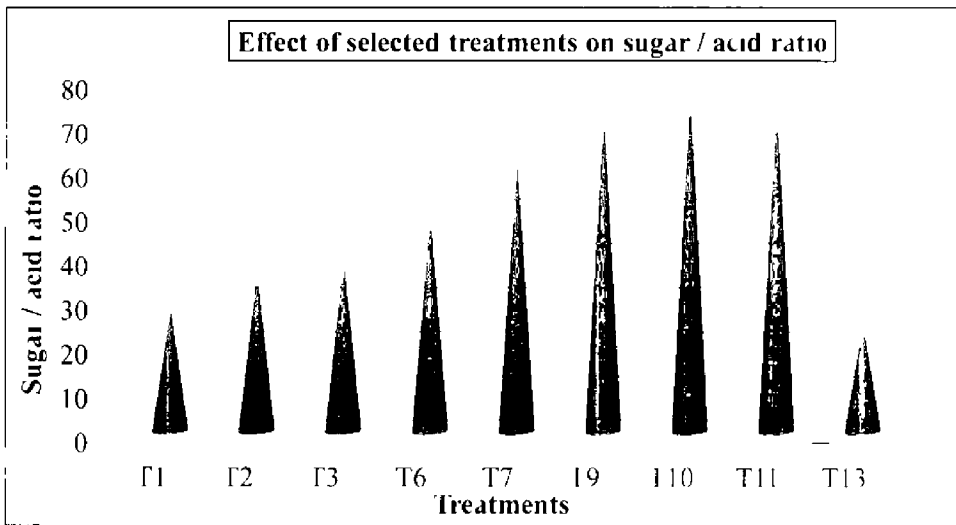


Figure 8 Foliar application of micronutrient mixture on sugar / acid ratio of ripened banana fruits

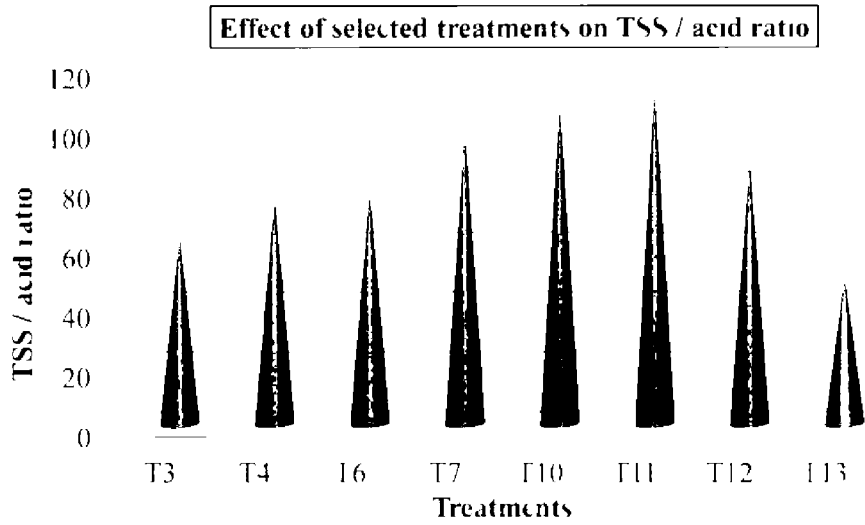


Figure 9 Foliar application of micronutrient mixture on TSS / acid ratio of ripened banana fruits

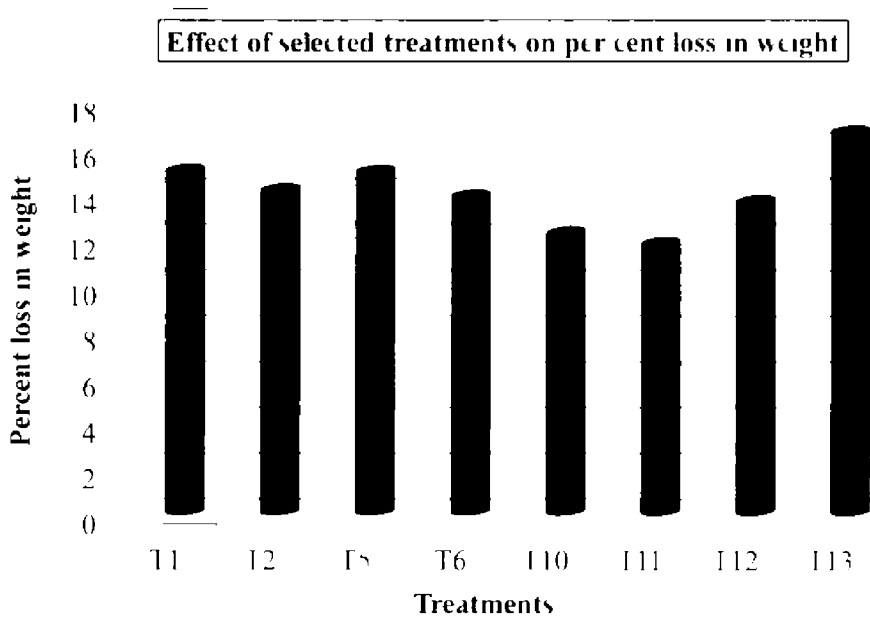


Figure 10 Foliar application of micronutrient mixture on per cent loss in weight during ripening of banana fruits

5.3.4. Effect of treatments on soil nutrient status

The effect of foliar application of micronutrients on soil nutrient status were studied and found that OC, N, P, K, Mg, Zn, B, Fe, B and Mo was found to be significant with treatment application while pH, EC, Ca, S and Cu content soil was found to be non significant

There was no significant effect of treatment application on soil pH and electrical conductivity. However average pH of the treatment plots were lower than that of control plot, which in turn enhance the nutrient availability and can bring about a better soil environment for plant growth. 3 sprays of 1 per cent micronutrient mixture treated plots were found to have lesser pH of 4.8. Even though non significant, average EC of treatment plots were higher than that of control plot. Highest EC value was found in 2 sprays of 2 per cent micronutrient mixture treated plots and lowest EC value was recorded 2 sprays of 4 per cent micronutrient mixture treated plots. It might be due to increase in the soluble salt content of the soils.

Organic carbon content of the soils was significantly influenced by treatment application. 3 sprays of 1 per cent micronutrient mixture treated plots have increased organic carbon content of soil whereas control plot was found to have lowest organic carbon content.

Available Nitrogen, phosphorus and potassium content of soil were significantly influenced by treatment application. With increasing level of micronutrient mixture the available nitrogen content of soil also increased. Highest available N was recorded in 3 sprays of 4 per cent micronutrient mixture treated plots whereas lowest was recorded in one spray of 3 per cent micronutrient mixture treated plots. Available P content of soil increased with treatment application as compared to that of control. Maximum available P content was found in 3 sprays of 4 per cent micronutrient mixture treated plots and control plot recorded minimum available P content. Available K content of soil also improved with treatment application as

compared to that of control. Highest available K content of soil was recorded in 2 sprays of 4 per cent micronutrient mixture treated plots while control plot was found to have lowest available K content.

Ca and S content of soil was not influenced by treatment application. Non significance of nutrients might be due to increased plant uptake and leaching losses. Under humid tropical climatic conditions, soils dominated by kaolinitic clay readily leaches out Ca^{2+} (Sims and Ellis, 1983). Mg content of treated plots was increased with increased time of treatment application. Highest Mg content was recorded in 3 sprays of 1 per cent micronutrient mixture treated plot.

Application of micronutrient mixture brought out significant changes in availability of iron, manganese, zinc, molybdenum and boron in soil. Zinc content of the soils exhibited significant difference among the treatments. Two sprays of 1 per cent micronutrient mixture treated plot recorded maximum Zn content of soils and control plot recorded the minimum. The maximum soil Zn concentration is mainly due to humus-Zn chelating process. The mobility and availability of trace elements is controlled by chemical and biochemical processes including precipitation, dissolution, adsorption-desorption, complexation-dissolution and redox processes (Hursthouse, 2001).

Iron content of soil was also significantly influenced by treatment application. With increasing level of micronutrient mixture the Fe content of soil has been decreased. Maximum Fe content was recorded in one spray of 4 per cent micronutrient mixture treated plot whereas control plot recorded the minimum content.

Mn content of soil was significantly influenced by treatment application. Highest Mn content was recorded with 3 sprays of 2 per cent micronutrient mixture treated plots. When Cu status is considered, treatment application did not influence the Cu content of soils. However, highest Cu content was found in 2 sprays of 1 per

cent micronutrient mixture treated plots and lowest was recorded in in one spray of 2 per cent micronutrient mixture treated plots

B content of soil was found to higher in all treated plots as compared to control plot One spray of 4 per cent micronutrient mixture treated plots was found to have highest B content in soil The contribution of foliar fertilization in increasing nutrient content of soils might be due to washing and by draining fertilizer solution from leaves to soil surface (Nomura *et al* , 2011)

Mo content of soil was found to higher in all treated plots as compared to control plot Three sprays of 4 per cent micronutrient mixture treated plots were found to have highest Mo content in soil

5.3.5. Effect of treatments on leaf nutrient contents

The effect of foliar spraying of micronutrient mixture showed significant influences in leaf nutrient contents of banana at bunching stage and at harvest Except nitrogen all other elements like P, K, Ca, Mg, S, Zn, B, Fe, Mn, Mo and Cu content of leaves increased at one or both stage of observation The increased in macro and micronutrient content of leaf might be attributed to fact that the physiological processes of leaves led to rapid absorption and utilization of nutrients for primary metabolic processes Arunachalam *et al* (1976) showed that adequate levels of nutrient in banana leaf ranged from 3.18-3.43, 0.46-0.54, 3.36-3.76, 2.3-2.4 and 0.25-0.28 per cent for N, P, K, Ca and Mg, respectively The results obtained in the present investigation are in line with Arunachalam *et al* (1976) The increased uptake of P, K, Mn and Zn by various crops were reported by Ghanta and Mitra (1993) in banana cv Giant Governor, Durgadevi *et al* (1997) in citrus, Lal *et al* (2000) in guava, Aggarwal *et al* (1975) in grapes, Afria *et al* (1999) in pomegranate, Nehete *et al* (2011) in mango and Yadav *et al* (2009b) in banana cv Grand Naine

Nitrogen content of banana leaves at bunching stage as well as at harvest was not influenced by treatment application. Even then, in both stages the N content ranged between 2.10- 4.20 per cent which was sufficient for crop growth. Two times foliar spraying of 3 per cent micronutrient mixture was found to have highest N content in leaves and control plants was recorded lowest N content in leaves. Hewitt (1955) reported that 2.6 per cent N in the leaf is adequate for banana, while Murray (1962) showed that <1.5 per cent nitrogen is designated as deficient for banana. There was gradual decrease in N content from bunching to harvest, because N is the primary element which may be mostly utilized by plants. Similar finding was reported by Jeyabaskaran and Pandey (2008) in Karpuravalli banana.

Significant differences were noticed in the P content of leaves among the treatments at the time of harvest of banana. Highest P content was recorded in 1, 2 and 3 sprays of 3 per cent micronutrient mixture treated plants. Control and one spray of 1 per cent micronutrient mixture were found to have lowest P content. This might be due to rapid phosphorus uptake by plants, their metabolism and translocation to plant parts. Similar findings were reported by Yadav *et al* (2009b) in banana cv Grand Name, Sankar *et al* (2013) in mango cv Alphonso.

Potassium content of leaves was found to be significant at bunching stage as well as at harvest of crop. 3 sprays of 3 per cent micronutrient mixture were found to have highest K content of leaves. Control plants had lowest K content in leaves. This might be due to role of micronutrients especially B in encouragement of potassium absorption from soil rather than utilization in plant tissues. Similar report was also obtained by Yadav *et al* (2009b) in banana cv Grand Name, Sankar *et al* (2013) in mango cv Alphonso, Jeyabaskaran and Pandey (2008) in Karpuravalli banana and Paul and Nair (2015) in banana cv Nendran. The application of micronutrients favoured the accumulation of macronutrients due to their role in activation of enzymes, involved in metabolic processes (Muthukrishnan *et al*, 2014).

Leaf Ca content at bunching stage and at harvest showed notable difference among the treatments. At bunching stage 2 sprays of 4 per cent micronutrient mixture and at harvest 3 sprays of 3 per cent micronutrient mixture recorded highest Ca content in leaves. Mg content was found highest with 3 sprays of 2 per cent micronutrient mixture at bunching stage. Mg content of leaves increased with increasing levels and time of treatment application. Sulphur content of leaves also increased with increasing level and time of treatment application. Highest S content was found in 3 sprays of 3 per cent micronutrient mixture at the time of harvest.

The addition of micronutrients either alone or through mixture, must have exerted its direct effect on its own composition. Application of micronutrient mixture brought out significant changes in leaf Zn, Fe, Mn, Cu, B and Mo content at bunching stage and at harvest of banana. Similar findings were also observed by Thiyyageshwari and Ramanathan (2001), Bhatt and Srivastava (2005), Kumbhar and Deshmukh (1993), Prabha and Singaram (1996).

Treatment application significantly influenced the Zn content of leaves at both stages of observation. With increasing level of micronutrient mixture Zn content of leaves also increased. At bunching stage, the highest Zn content was recorded in 3 sprays of 4 per cent micronutrient mixture treated plants and at harvest, 3 sprays of 3 per cent micronutrient mixture treated plants recorded the maximum Zn content. In both stages, control plants found to have the lowest Zn content. This increase in leaf Zn content was due to its maximum absorption from Zn source through foliage and less translocation to other parts of the plant. Similar finding was also reported by Yadav *et al* (2009b) in banana cv Grand Name, Dalal *et al* (2011) in Ber, Hafeezur-Rahman and Izhar-ul-Haq (2014) in sweet orange and Lalithya *et al* (2014) in sapota.

Application of micronutrients increased the concentration of iron as it was supplied through micronutrient spray. Fe concentration of leaves in treated plants

was found to be higher than that of control plants. 3 sprays of 4 per cent micronutrient mixture found to have highest Fe content in leaves at both stages whereas control plants recorded the lowest Fe content. The acidic environment in the root vicinity leads the reduction of Fe^{3+} to Fe^{2+} form which is water soluble and available form to plants. This results in increase in the Fe content of leaves and might have led to higher Fe uptake. The results are in conformity with Lalithya *et al* (2014) in sapota, Yadav *et al.* (2009b) in banana cv Grand Naine, Dalal *et al* (2011) in Ber, Jeyabaskaran and Pandey (2008) in Karpuravalli banana.

Significant differences were noticed in the Mn uptake among the treatments. However, very high levels of Mn were observed in banana leaves. At bunching stage, 3 sprays of 4 per cent micronutrient mixture and at harvest 3 sprays of 2 per cent micronutrient mixture were recorded with highest Mn content of leaves. Turner and Barkus (1983) also reported the high Mn concentration in the leaves as a special character of banana. Average Mn content of the treated plants was higher than the control plants. A concentration upto 2200 ppm is considered as optimum for banana whereas above which it is high under subtropical conditions as reported by Turner and Barkus (1983). Similar findings were obtained by Dalal *et al* (2011) in Ber, Hafeez-ur-Rahman and Izhar-ul-Haq (2014) in sweet orange, Jeyabaskaran and Pandey (2008) in Karpuravalli banana and Hasan *et al* (2012) in pomegranate leaves.

Average Cu content of the leaves in treated plants was higher than that of control at bunching stage as well as at harvest of banana. 3 sprays of 3 per cent micronutrient mixture found to have highest Cu content. Application of micronutrients increased the concentration of copper as it was supplied through foliar spray. With increasing level and time of application of micronutrient mixture the Cu content of the leaves also increased. Similar findings were also reported by Paul and Nair (2015) in banana cv Nendran, Lalithya *et al* (2014) in sapota and Yadav *et al* (2009b) in banana cv Grand Naine.

Significant differences in B concentration of the leaves were noticed at both stages of observation with foliar application of micronutrient mixture. This might be due to the presence of boron binding compounds in the cell which might have increased the mechanism of boron uptake, which is thought to be a non-metabolic process determined in plant by the formation of non-exchangeable boron complexes within the cytoplasm and cell wall. Similar findings were reported by Brown and Hu (1994) in sunflower, Saran and Kumar (2011) in mango, Asgharzade *et al* (2012) in apple, Sankar *et al* (2013) in mango cv Alphonso.

Average Mo content of the leaves in treated plants was higher than that of control at both of stages. Three sprays of 3 per cent micronutrient mixture was found to make highest Mo content in leaves. Application of micronutrients increased the concentration of molybdenum as it was supplied through foliar spray. With increasing level and time of application of micronutrient mixture the Mo content of the leaves also increased.

The results of net return and BCR (Table 21) indicated that there exists a significant difference between the treatments. The treatment T₁₁ (3 sprays of 3 % micronutrient mixture) registered the maximum net return and BCR of Rs 6,94,964 ha⁻¹ and 2.29. Foliar spraying of micronutrient mixture enhanced the growth and yield parameters, which in turn gave higher net returns and BCR. This was in accordance with the results of Yadav and Patel (2013) in banana cv Grand Name, Patel *et al* (2010) in banana cv Basrai, Pathak *et al* (2011) and Bauri *et al* (2014) in banana cv Martaman (AAB, Silk).

Summary

6. SUMMARY

The salient findings of the present study entitled “Formulation and evaluation of micronutrient mixture for foliar application in TC banana (*Musa* sp) var Nendran” are summarized in this chapter

Investigations were carried out at College of Agriculture, Padannakkad and Regional Agricultural Research Station farm, Nileshwar during November 2014 to March 2016, with the objectives of preparation of a micronutrient mixture formulation containing Zinc, Iron, Boron, Copper, Manganese and Molybdenum for foliar spray, to evaluate stability of formulation and its keeping quality and to investigate the effect of this formulation on growth, yield and quality of fruits in banana (*Musa* AAB) Nendran The whole study consisted of three parts – preparation of micronutrient mixture formulation, hardening studies and field experiment

Preparation of micronutrient mixture as a liquid formulation was tried with the use of different micronutrient salts containing Zn, B, Fe, Mn, Cu and Mo The formulation technique was standardized and it contains two liquids Solution A and Solution B Solution A consisting of $ZnSO_4 \cdot 7H_2O$, $CuSO_4 \cdot 5H_2O$, $FeSO_4 \cdot 7H_2O$, H_3BO_3 , $MnSO_4 \cdot H_2O$ and $(NH_4)_6Mo_7O_{24} \cdot 4H_2O$ in desired proportion For maintaining stability and keeping quality of solution A, various chemicals were tried Citric acid (20 %) was found best for maintaining keeping quality of solution A Its stability was periodically monitored at specific intervals Solution B is prepared as 1 per cent humic acid Required quantity of this has to be blended with diluted solution A just before spraying The second and third part of the experiment was evaluation of this mixture This micronutrient mixture was diluted to different concentrations and applied as foliar spray in tissue culture banana cv Nendran to study the effect of this nutrient mixture on hardening and field experiment

The hardening studies were conducted in the mist chamber, Department of Plant Biotechnology, College of Agriculture, Padannakkad, Kerala with TC plants of uniform size multiplied from same genotype. The experiment was carried out in completely randomized design with 12 treatments and one absolute control maintaining 3 replications each and there were 4 plants in each replication. The treatments consisted of 4 levels of micronutrient mixture (0.25 %, 0.5 %, 0.75 % and 1 %) sprayed at 3 different intervals. The 3 different sprays were one spray was given immediately after planting, two sprays at immediately and two weeks after planting and three sprays at immediately, two and three weeks after planting during secondary hardening stage. Thus there were 12 treatment combinations and one control which were T₁, T₅, T₉ at 0.25 per cent (one, two and three sprays), T₂, T₆, T₁₀ at 0.50 per cent (one, two and three sprays), T₃, T₇, T₁₁ at 0.75 per cent (one, two and three sprays), T₄, T₈, T₁₂ at 1 per cent (one, two and three sprays) and T₁₃- control. The important biometric observations such as plant height, number of leaves, leaf length and leaf breadth were recorded at one month after keeping it for hardening. The results showed that the application of micronutrient mixture have positive effect on biometric characters of the crop. Maximum plant height (20.2 cm) was recorded in T₁₀ (0.5% micronutrient mixture as 3 sprays). Highest number of leaves (6.8) and leaf breadth (7.3 cm) was recorded in T₉ (0.25% micronutrient mixture as 3 sprays) and T₅ recorded highest leaf length (19.6 cm).

The field experiment was carried out at Regional Agricultural Research Station farm, Nileshwar to study the effect of foliar application of micronutrient mixture in tissue culture banana cv Nendran. One fifty six hardened tissue culture banana plantlets were planted in randomized block design with thirteen treatments and three replications such that four plants were maintained in each replication. Nitrogen, Phosphorus and Potassium application and other cultural practices were uniformly followed for all the plants as per Package of Practices, KAU (2011).

The treatments consisted of 4 levels of micronutrient mixture (1 %, 2 %, 3 % and 4 %) sprayed at 3 different intervals. The 3 different sprays were one spray was given at two months after planting, two sprays at two and four months after planting and three sprays at two, four and six months after planting. Thus there were 12 treatment combinations and one control which were T₁, T₅, T₉ at 1 per cent (one, two and three sprays), T₂, T₆, T₁₀ at 2 per cent (one, two and three sprays), T₃, T₇, T₁₁ at 3 per cent (one, two and three sprays), T₄, T₈, T₁₂ at 4 per cent (one, two, three sprays) and T₁₃- control

The results of the field experiment showed that treatment application significantly enhanced the vegetative characters like pseudostem height, pseudostem girth and number of leaves and yield parameters like bunch weight, number of fingers, average weight of fingers, finger length and minimum days to harvest. Maximum pseudostem height was recorded in T₁₂ (4 % micronutrient mixture as 3 sprays) at 4 months after planting, pseudostem girth at 90 cm was found to highest in T₁₀ (2 % micronutrient mixture as 3 sprays) at the time of shooting, highest number of leaves was recorded in T₆ (2 % micronutrient mixture as 2 sprays) at 4 months after planting as well as at the time of shooting and number of suckers produced at harvest were highest in T₁₀ (2 % micronutrient mixture as 3 sprays) and T₁₂ (4 % micronutrient mixture as 3 sprays). The highest male bud weight was found in T₅ (1 % micronutrient mixture as 2 sprays). Bunch weight was found to be highest in T₁₁ (3 % micronutrient mixture as 3 sprays) with 12.76 kg whereas control T₁₃ recorded the lowest bunch weight of 8.32 kg with 53.4 % increases in yield for treatment over control. T₉ (1 % micronutrient mixture as 3 sprays) found to have maximum number of fingers, highest average weight of fingers and finger length was found in T₁₁ (3 % micronutrient mixture as 3 sprays). Highest breadth was recorded with T₁₁ (3 % micronutrient mixture as 3 sprays) and T₁₀ (2 % micronutrient mixture as 3 sprays). T₁₀ produced the bunch in shortest time of 186 days whereas T₁₁ (3 % micronutrient mixture as 3 sprays) took minimum days (271 days) for harvesting as well as

minimum bunch maturity period (82 days) and T₄ (4 % micronutrient mixture as 1 spray) took minimum days for harvesting to ripening of fruits (4 days)

Fruit characteristics like titrable acidity, TSS, pulp to peel ratio, total sugars, reducing sugars, non-reducing sugar content, sugar / acid ratio, TSS / acid ratio, per cent loss in weight during ripening and keeping quality of the fruits at ambient condition were studied. All the fruit quality parameters showed significant differences among the treatments. Lesser acidity indicates better quality and the lowest titrable acidity was recorded in T₁₁ (3 % micronutrient mixture as 3 sprays) whereas control T₁₃ recorded the maximum acidity content of fruits. Maximum TSS content (30.0 °brix) was found in T₁₁ (3 % micronutrient mixture as 3 sprays) whereas T₁₂ (4 % micronutrient mixture as 3 sprays) recorded highest pulp to peel ratio of 3.70. The highest total sugar content (21.3 %) and reducing sugar content (19.0 %) was recorded with T₁₀ (2 % micronutrient mixture as 3 sprays) and the highest non-reducing sugar content was recorded in T₅ (1 % micronutrient mixture as 2 sprays) whereas T₁₁ (3 % micronutrient mixture as 3 sprays) recorded the lowest non-reducing sugar content. Highest sugar / acid ratio and keeping quality of the fruits at ambient condition were found in T₁₀ (2 % micronutrient mixture as 3 sprays). T₁₁ (3 % micronutrient mixture as 3 sprays) was found to have highest TSS / acid ratio and minimum loss in weight of fruits during ripening.

The effect of foliar application of micronutrients on soil nutrient status were studied and found that OC, N, P, K, Mg, Zn, B, Fe, B and Mo status were found to be significant with treatment application while pH, EC, Ca, S and Cu content soil was found to be non significant.

Lowest pH (4.8) was found in T₉ which receiving 1 per cent micronutrient mixture as 3 sprays, highest EC value was found in T₆ (2 % micronutrient mixture as 2 sprays) treated plots and lowest EC value was recorded T₈ (4 % micronutrient mixture as 2 sprays). T₉ (1 % micronutrient mixture as 3 sprays) found to have

highest organic carbon content of soil whereas T₁₃ (control) recorded the lowest organic carbon content

Available N content of the soils was significantly influenced by treatment application. With increasing levels and times of application of micronutrient mixture, the available nitrogen content of soil also increased. Highest available N and available P content were recorded in T₁₂ (4 % micronutrient mixture as 3 sprays), whereas control plot recorded the minimum. Available K content of soil also improved with treatment application as compared to that of control. Highest available K content of soil was recorded in T₈ (4 % micronutrient mixture as 2 sprays), while control plot was found to have the lowest content.

Ca and S content of soil was not influenced by treatment application. However, maximum Ca content was recorded in T₅ (1 % micronutrient mixture as 2 sprays) and S content in T₁₁ (3 % micronutrient mixture as 3 sprays). Mg content of treated plots was increased with increasing time of application. Highest Mg content was recorded in T₉ (1 % micronutrient mixture as 3 sprays).

Application of micronutrient mixture brought out significant changes in available Zn, B, Fe, Mo and Mn. Highest Zn and Cu content was found in T₅ (1 % micronutrient mixture as 2 sprays) and control plot recorded the minimum. With increasing level of micronutrient mixture the Fe content of soil has been decreased. Maximum Fe content was recorded in T₄ (4 % micronutrient mixture as one spray) and Mn content was in T₁₀ (2 % micronutrient mixture 3 sprays). B content of soil was found to be higher in all treated plots as compared to control plot. T₄ (4 % micronutrient mixture as one spray) was found to have highest B content in soil. Highest Mo content was recorded with T₁₂ (4 % micronutrient mixture 3 sprays).

As leaf nutrient concentration of banana is considered, significant influence of treatment application of micronutrient mixture were found in P, K, Ca, Mg, S, Zn, B, Fe, Mn, Mo and Cu content at one or both stage of observation (bunching and at

harvest) Highest N content was found in T₇ (3 % micronutrient mixture 2 sprays) at bunching stage as well as at the time of harvest of banana Highest P content was found in T₃ (3 % micronutrient mixture 1 spray) and T₇ (3 % micronutrient mixture 2 sprays) at harvest Highest K content of leaves was recorded in T₁₁ (3 % micronutrient mixture as 3 sprays) at bunching stage and at harvest

At bunching stage T₈ (4 % micronutrient mixture as 2 sprays) and at harvest T₁₁ (3 % micronutrient mixture as 3 sprays) recorded highest Ca content in leaves Mg content of leaves increased with increasing levels and time of application Mg content was found highest with T₁₀ (2 % micronutrient mixture as 3 sprays) at bunching stage and the highest S content was found in T₁₁ (3 % micronutrient mixture as 3 sprays) at harvest

Regarding micronutrient content of leaves there was significant differences between the treatments and the control with respect to Zn, Fe, Mn, Cu, Mo and B content at bunching and at harvest of banana The highest Zn content was recorded in T₁₂ (4 % micronutrient mixture as 3 sprays) at bunching stage and T₁₁ (3 % micronutrient mixture as 3 sprays) at harvest whereas control plants found to have the lowest Zn content With increasing level of micronutrient mixture Zn content of leaves also increased Fe concentration of leaves in treated plants was found to be higher than that of control plants

At both stages of observation, T₁₂ (4 % micronutrient mixture as 3 sprays) found to have highest Fe content in leaves Average Mn content of the treated plants was higher than the control plants Highest Mn content was found in T₁₂ (4 % micronutrient mixture as 3 sprays) at bunching stage and T₁₀ (2 % micronutrient mixture as three sprays) at harvest Average Cu content of the leaves in treated plants was higher than that of control at bunching stage and at harvest In both stages, T₁₁ (3 % micronutrient mixture as 3 sprays) found to have highest Cu content

Highest B content was recorded in T₃ (3 % micronutrient mixture as one spray) at bunching and T₁₂ (4 % micronutrient mixture as 3 sprays) at harvest. Mo content was found to be highest with T₁₁ (3 % micronutrient mixture as 3 sprays) in bunching and at harvest.

The Highest net return (Rs 6,94,964) and benefit cost ratio (2.3) was registered with foliar application of 3 % micronutrient mixture as three sprays (T₁₁). The quantity of spray solution required for best treatment T₁₁ on projection to one ha is 56 litres, with an additional cost of Rs 160/ litre for the mixture plus Rs 600 / day as cost of labour for application.

The positive effects of the micronutrient foliar spray solution indicate the need of these nutrients on banana. This effect can be further evaluated on vegetables and other crops. On the basis of the results thus obtained, recommendation can be made on an ad hoc basis.

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7. REFERENCES

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var. NENDRAN**

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ABSTRACT

The experiment entitled "Formulation and evaluation of micronutrient mixture for foliar application in TC banana (*Musa sp*) var Nendran" was carried out with the objectives of preparing a micronutrient mixture formulation for foliar spray, to evaluate stability of formulation and its keeping quality and to investigate the effect of this formulation on growth, yield and quality of fruits in banana (*Musa sp*) Nendran. The whole study consisted of three parts – preparation of micronutrient mixture formulation, hardening studies and field experiment. The studies were carried out at College of Agriculture, Padannakkad and RARS farm, Nilesiwar during 2014–2016.

Micronutrient mixture formulation was prepared with the use of different micronutrient salts containing Zn, B, Fe, Mn, Cu and Mo. The formulation technique was standardized and it was prepared as two liquids. Solution A and Solution B. Solution-A consisted of ZnSO₄, CuSO₄, FeSO₄, H₃BO₃, MnSO₄ and (NH₄)₂MoO₄ in desired proportion. For maintaining stability and keeping quality of solution A, addition of citric acid was found to be the best. Solution B was prepared as 1 per cent humic acid. Required quantity of solution A and B was diluted and blended just before spraying.

The hardening studies were conducted in the poly house with TC plants of uniform stage multiplied from same genotype. The experiment was carried out in CRD with 12 treatments and one control with 3 replications each and there were 4 plants in each unit. The treatments consisted of 4 levels of micronutrient mixture (0.25%, 0.5%, 0.75% and 1%) sprayed at 3 different intervals. The 3 different spray scheduled were one spray- immediately after planting and 2 subsequent sprays given at 2nd and 3rd weeks after planting during secondary hardening stage. Thus there were 12 treatment combinations and one control. The treatments were T₁, T₂, T₃ and T₄ - one spray, T₅, T₆, T₇ and T₈ - two sprays and T₉, T₁₀, T₁₁ and T₁₂ - three sprays of 0.25, 0.5, 0.75 and 1 per cent concentrations respectively. T₁₃ was control.

Significant differences among the treatments were observed in the biometric characters of the plants. Plant height (20.2 cm) was found to be highest in T₁₀ (0.5% as 3 sprays). Highest number of leaves (6.8) and leaf breadth (7.3 cm) was recorded in T₉ (0.25% as 3 sprays) and T₅ recorded highest leaf length (19.6 cm).

The field experiment was carried out at RARS farm, Nileshwar in RBD with 12 treatments and one control with 3 replications each and there were 4 plants in each unit. Major nutrients *viz* N, P, K and other cultural practices were uniformly followed for all plants as per PoP, KAU (2011). The treatments consisted of 4 levels of micronutrient mixture (1%, 2%, 3% and 4%) sprayed at 3 different intervals. The 3 different spray schedules were one spray was given at 2 MAP, two sprays at 2 and 4 MAP and three sprays at 2, 4 and 6 MAP. T₁, T₂, T₃ and T₄ - one spray, T₅, T₆, T₇ and T₈ - two sprays and T₉, T₁₀, T₁₁ and T₁₂ - three sprays of 1, 2, 3 and 4 per cent concentration respectively and T₁₃ was control.

The results of the field experiment showed that treatment application significantly enhanced the vegetative characters like pseudostem height, pseudostem girth and number of leaves. Among the yield characteristics, bunch weight, number of fingers, average weight of fingers, finger length and days to harvest showed significant difference. Bunch weight was found to be highest in T₁₁ (12.8 kg) with 53.4% increase in yield over control (8.3 kg). The highest number of fingers was found in T₉, while highest average weight of fingers, finger length was found in T₁₁. Minimum days from planting to bunch emergence (186 days) was noticed in T₁₀ whereas T₁₁ took minimum days from planting to harvest (271 days) as well as minimum bunch maturity period of 82 days.

Characteristics of fruits like titrable acidity, TSS, pulp to peel ratio, total sugars, reducing sugars, non-reducing sugar content, sugar / acid ratio, TSS / acid ratio, per cent loss in weight during ripening and keeping quality of the fruits at ambient conditions were also found to be influenced by treatments.

After harvest, the effect of these treatments on soil nutrient availability was studied. The results showed that soil organic carbon status and available nutrient status of N, P, K, Mg, Zn, B, Fe, B and Mo were found to be significant. Similarly, leaf nutrient analysis at bunching and at the time of harvest revealed that P, K, Ca, Mg, S, Zn, B, Fe, Mn, Mo and Cu content at one or both stages of observation showed significant difference among the treatments.

The results of both the hardening studies and field experiment indicated the beneficial effect of micronutrient mixture on TC banana. For hardening studies T_9 - 0.25 per cent and T_{10} - 0.5 per cent given as 3 sprays were noticed better. On mature plants T_{10} - 2 per cent and T_{11} - 3 per cent both as 3 sprays were found promising. Yield increase was noticed as 53.4 per cent over control with an enhancement of 4.4 kg in bunch weight. The benefit cost ratio calculated on projection to one ha is 2.3 with net returns of Rs 6,94,964 ha⁻¹ in T_{11} (3% micronutrient mixture as 3 sprays). The quantity of spray solution required for best treatment T_{11} on projection to one ha is 56 litres, with an additional cost of Rs 160 / litre for the mixture plus Rs 600 / day as cost of labour for application.