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(54) Title: MICROBIAL PROCESS AND COMPOSITION FOR AGRICULTURAL USE

(57) Abstract: Microbial compositions comprising at least two components are disclosed. The first component comprises HYTa which is a consortium of microbes derived from fertile soils and commercial sources. The second component comprises at least one of chitin, chitosan, glucosamine and amino acids. The various microbes in HYTa are capable of fixing nitrogen, digesting proteins and other biopolymers such as chitin and chitosan, providing protection against plant pathogens and supplementing the microbial flora of soil. Also disclosed are processes where the aforementioned microbial compositions are used to treat soil, seeds, seedlings and/or plant foliage alone or in combination with chitin, chitosan, glucosamine and/or amino acids.



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**MICROBIAL PROCESS AND COMPOSITION
FOR AGRICULTURAL USE**

This application claims the benefit under 35 U.S.C. §119
5 of U.S. Provisional Application Serial No. 61/355,447, filed
June 16, 2010.

FIELD OF THE INVENTION

Microbial processes and microbial compositions are
10 disclosed that enhance crop production, increase plant
defensive processes, decrease the level of plant pathogens
and reduce the amount of fertilizer used.

BACKGROUND OF THE INVENTION

15 Microbes have previously been used in agriculture.
Examples include those disclosed in US Patents 4,952,229;
6,232,270 and 5,266,096.

Chitin has also been used in agriculture either as a
protein complex (US Patent 4,536,207) or in combination with
20 various microbes (US Patents 6,524,998 and 6,060,429)

Chitosan in combination with other components has
been used in agricultural applications. See e.g. US Patents
6,649,566; 4,812,159; 6,407,040; 5,374,627 and 5,733,851. It
has also been used to treat cereal crop seeds. See US Patent
25 4,978,381. US Patent 6,524,998 also discloses that chitosan
can be used in combination with specific microbes for
agricultural use.

Notwithstanding the foregoing, there is a need to
provide improved microbial compositions and processes that
30 improve crop yield and reduce the amount of conventional
fungicides and insecticides used in agricultural and
horticultural applications.

SUMMARY OF THE INVENTION

Microbial compositions comprising at least two components are disclosed. The first component comprises HYTa which is a consortium of microbes derived from fertile soils and commercial sources. The second component comprises at least one of chitin, chitosan, glucosamine and amino acids. The various microbes in HYTa are capable of fixing nitrogen, digesting proteins and other biopolymers such as chitin and chitosan, providing protection against plant pathogens and supplementing the microbial flora of soil.

Also disclosed are processes where the aforementioned microbial compositions or their components are used to treat soil, seeds, seedlings and/or plant foliage.

In preferred embodiments HYTa is activated in an aqueous solution for 24-168 hours to allow the microbes to grow and reproduce before being used in the process. The conditions of the incubation influence the overall initial properties of HYTa.

In a preferred embodiment, HYTa is activated in the presence of chitin. Chitin responsive microbes in HYTa proliferate in this environment. This results in HYTa that has all of the properties of HYTa. However, it has enhanced capability against chitin containing plant pathogens.

In a preferred embodiment, the HYTa is activated in the presence of chitin, chitosan, glucosamine and amino acids. In this embodiment, after growth, the HYTa may contain residual chitin, chitosan, glucosamine and/or amino acids. Under such circumstances, the culture constitutes the disclosed microbial composition and can be applied directly to soil, seed, seedlings or plant foliage. Alternatively, one or more second components can be added to supplement the second components already in the composition or to change

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the components present in the thus formed microbial composition.

In some embodiments, the activated HYTa is combined with one or more second components and applied to the soil, seed, seedlings or plant foliage or the HYTa and the second component (s) are applied separately. Such second components include chitin, chitosan, glucosamine and amino acids.

The application of the disclosed microbial formulations allows for the elimination or significant reduction in the amount of fertilizer, fungicide and insecticide used in agricultural applications. In some embodiments, the use of the microbial formulations results in a decrease in the amount of green house gas emissions.

Also disclosed is treated soil composition comprising soil treated with HYTa.

Also disclosed is treated plant comprising plant treated with HYTa.

Also disclosed are treated seeds, seedlings and plants comprising seed, seedling or plant treated with HYTa.

In one aspect of the invention is a composition comprising HYTa and at least one of chitin, chitosan, glucosamine and amino acids.

In one aspect of the invention is a composition comprising HYTa and at least one of HYTb and HYTc.

In one aspect of the invention is a composition comprising HYTa, HYTb and HYTc.

In one aspect of the invention is a process comprising contacting soil, seed, seedling or plant foliage with HYTa to form treated soil, seed, seedling, or plant foliage.

In one aspect of the invention is a composition comprising soil, a plant, a seed, or a seedling treated with HYTa.

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BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a diagram of the test area involving the growth of durum wheat in Sonora, Mexico where HYTa and HYTb were used.

- 5 Figure 2 is the same diagram as Figure 3 and shows zones that were compromised and suffered impairment by external factors during the trial.

Figure 3 graphically depicts the results from treating the soil and foliage of durum wheat with HYTa and HYTb.

- 10 Figure 4 shows the yield of melons as a function of size for soil and foliage that was either not treated or treated with HYTa and HYTb.

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Figure 5 shows the yield of potatoes having diameters greater than 42 mm that were treated with HYTa, HYTb and HYTc as compared to untreated potatoes.

5 DETAILED DESCRIPTION

Microbial compositions comprising HYTa and a second component are disclosed. HYTa is a consortium of microbes derived from further soils and commercial sources. The second component comprises at least one of chitin, chitosan, glucosamine and amino acids. The various microbes in HYTa are capable of fixing nitrogen, digesting proteins and other biopolymers such as chitin and chitosan, providing protection against plant pathogens and supplementing the microbial flora of soil. The microbial compositions or their components are used to treat soil, seeds, seedlings and/or plant foliage.

HYTa

As used herein, the term "HYTa" refers to a consortium of microbes derived from fertile soil samples and commercial sources. HYTa was deposited with the American Tissue Type Culture (ATTC), Rockville, Maryland, on May 19, 2010 with an assigned deposit designation of PTA-10973.

Table 1 identifies some of the microbes in HYTa that are believed to be responsible for the beneficial effects observed when it is used to treat soil and/or foliage.

Table 1

	<u>Bacteria</u>
	I. Azotobacter
	1. <i>Azotobacter vinlandii</i>
5	II. Clostridium
	1. <i>Clostridium pasteurianum</i>
	2. <i>Clostridium beijerinckii</i>
	3. <i>Clostridium sphenoides</i>
	4. <i>Clostridium bifermentans</i>
10	III. Lactobacillus
	1. <i>Lactobacillus paracasei</i> ss. <i>paracasei</i>
	2. <i>Lactobacillus acidophilus</i>
	3. <i>Lactobacillus delbrueckii</i> ss. <i>Bulgaricus</i>
	4. <i>Lactobacillus brevis</i>
15	IV. Bacillus
	1. <i>Bacillus amyloliquefaciens</i> (<i>Bacillus subtilis</i> (SILoSil® BS))
	2. <i>Bacillus thuringiensis</i> var. <i>kurstakii</i> (<i>Bacillus thuringiensis</i> (Strains HD-1))
20	3. <i>Bacillus thuringiensis</i> var. <i>canadensis</i> (<i>Bacillus cereus</i> group)
	4. <i>Bacillus pasteurii</i> (<i>Bacillus cereus</i> group)
	5. <i>Bacillus sphaericus</i> (subgroup I, III, and IV)
	6. <i>Bacillus megaterium</i> (subgroup A)
25	V. Acetobacter or Gluconacetobacter
	1. <i>Acetobacter aceti</i> ss. <i>liquefaciens</i>
	2. <i>Acetobacter aceti</i> ss. <i>xylinum</i>
	VI. Enterococcus
	1. <i>Enterococcus faecium</i> (subgroup A)

VII. Pediococcus

1. *Pediococcus pentosaceus*

VII. Rhizobium

1. *Rhizobium japonicum*

5 **Fungi****I. Saccharomyces**

1. *Saccharomyces cerevisiae*

II. Penicillium

1. *Penicillium roqueforti*

10 **III. Monascus**

1. *Monascus ruber*

IV. Aspergillus

1. *Aspergillus oryzae*

V. Trichoderma

- 15
1. *Trichoderma harzianum* (TRICHOSIL)

Plantae**I. Arthrospiro**

1. *Arthrospira platensis*

II. Ascophyllum

- 20
1. *Ascophyllum nodosum*

Other microorganisms contained in HYTa: *Nitrobacter*,
Nitrosomonads, *Nitrococcus*, *Pseudomonas*, *Micrococcus luteus*,
Actinomycete, *Azotobacter vinelandii*, *Lactobacillus casei*,
25 *Trichoderma harzianum*, *Bacillus licheniformis*, *Pseudomonas*
fluorescens and *Streptomyces*.

Active microbes in HYTa include nitrogen-fixing microorganisms native to soil. These are *Azotobacter vinelandii* and *Clostridium pasteurianum*. *Bacillus subtilis* provides enzymes for breaking down plant residue. *Bacillus cereus* provides additional enzymes to break down plant residue and penicillinase to decrease unwanted bacteria. *Bacillus megaterium* degrades complex sugars after crop residue breakdown. *Lactobacillus* provides food for the microbes in HYTa and controls the pH of the environment. The *Nitrobacter* organisms oxidize ammonia to nitrite (NO₂) while the *Nitrosomonas* microbes oxidize nitrite to nitrate (NO₃).

An important property of HYTa is the fixation of atmospheric nitrogen. The nitrogen fixing capability of the microbes in HYTa is enhanced by the assistance of other organisms in HYTa. Nitrogen fixation requires that phosphorous (P), potassium (K) and carbon (C) be available. HYTa contains microbes that are able to decompose P, K, and C within the soil. In addition, the nitrogen fixing bacteria provide a source of nitrogen for the other microbes in HYTa.

Nitrogen fixation may occur in a non-symbiotic manner by the bacteria *Nitrosomonas*, *Nitrobacter*, *Azotobacter vinelandii*, and *Clostridium pasteurinum* present in HYTa or in a symbiotic manner as occurs in root nodules by way of the *Rhizobium* bacteria.

The carbon required by the nitrogen fixing microbes in HYTa is provided by the C decomposers which convert the complex organic compounds in soil into simple compounds such as sugars, alcohols, and organic acids. The C decomposers include many of the above identified microbes.

Phosphorus is necessary for the nitrogen fixing microbes to proliferate and is obtained from the metabolic activity of the P decomposers which convert immobilized

phosphorus in the soil into a bio-available phosphorus nutrient. P decomposers in HYTa include *Azotobacter*, *Bacillus subtilis*, *Pseudomonas fluorescens* and *Micrococcus luteus*.

5 The potassium required by the nitrogen fixers is provided by the K decomposer microbes present in HYTa which activate the potassium from the soil. K decomposers in HYTa include *Pseudomonas fluorescens*.

 Three important microbes in HYTa are *Bacillus subtilis* (SILoSIL® BS) *Bacillus thuringiensis* strains HD-1
10 and HD-73 (SILoSIL® BT), and *Trichoderma harzianum* (TRICHOSIL). These organisms are present ATTC deposit PTA-10973. They were originally obtained from Biotechnology Agroindustrial S.A. DE C.V., Morelia, Michoacan, Mexico.

15 *Bacillus subtilis* (SILoSIL® BS) is a Gram positive bacterium which is mesophilic and grows at an optimum temperature between 25 and 35°C. It is aerobic and can grow in anaerobic conditions and utilizes a wide variety of carbon sources. It contains two nitrate reductases, one of
20 which is utilized for nitrogen assimilation. It is capable of secreting amylase, proteases, pullulanases, chitinases, xylanases and lipases.

Bacillus thuringiensis (Strains HD-1 and HD-73 (SILoSIL® BT)) are Gram Positive anaerobic facultative
25 bacteria, in the form of a peritrichous flagella. Strains HD-1 and HD-73 synthesize crystals with diverse geometric forms of proteic and insecticide activity during the spore period. Strains HD-1 and HD-73 secrete exochitinases when in a chitin containing medium and can be utilized for the
30 degradation of the crustacean residues during the production of chitooligosaccharides.

Trichoderma harzianum (TRICHOSIL) is a saprophyte fungus. It exhibits antibiotic action and biological

competition and for this reason has biological control properties. It produces enzymes that degrade cell walls or a combination of such activities. It produces glucanases, chitinases, lipases, and extracellular proteases when it
5 interacts with some pathogenic fungi, such as *Fusarium*.

As shown above the metabolism of each group of bacteria are closely interdependent and live in a close symbiotic association for the proper performance of HYTa.

Besides carbon, hydrogen, phosphorus, potassium,
10 sulfur and various trace elements, a mix of special growth factors, such as B complex, free L-amino acids, and ultra soluble trace elements are important for optimal bacterial growth. Fermenting yeasts are incorporated into HYTa to provide these components. The N₂ fixing process requires
15 large amounts of ATP. The amount of ATP naturally present is not enough to fuel biological N₂ fixation. The fermentation of the yeast in HYTa compensates for the large energy deficit. During fermentation, organic acids are formed in the respiratory process and together with the phosphorous
20 released by the P decomposers, form ATP. The ATP is used in the biological nitrogen fixation process.

HYTa contains enzymes and beneficial soil microorganisms that replace those that have been depleted due to the excessive use of chemicals which results in
25 diminishing crop yields. By increasing the microbial activity in the soil with HYTa, the bacteria causes the nutrients and micro-elements to be absorbed (mineralized) more efficiently and effectively by plants.

Humus is transformed by some of the microorganisms in
30 HYTa that impregnate both the soil and the radical apparatus of the plant. This process provides increased nutrition to the plant. This increases the nutrients and the essential

elements available in the soil that can be absorbed by plants.

The use of HYTa alone or in combination with chitin, chitosan, glucosamine and/or amino acids (1) provides
5 nutrients and elements in the soil that increase crop yields by 25-55%, (2) reduces green house gas emissions, (3) increases the efficiency of mineral fertilizers (3) reduces the use of conventional fungicides and other pesticides, (4) increases the production of plant growth regulators, (5)
10 improves soil structure, tilth, and water penetration and retention, (6) cleans up chemical residues and (7) shifts soil pH toward neutral pH.

MICROBIAL COMPOSITIONS

15 HYTa can be used, alone or in combination, with one or more components selected from the group of one or more amino acids, chitin, chitosan and/or glucosamine. In some cases, Acetyl-D-glucosamine can be included in the microbial composition. The microbial composition includes any and all
20 combinations of the aforementioned components. Particularly preferred combinations include: (1) HYTa and chitin; (2) HYTa and chitosan; (3) HYTa and glucosamine; (4) HYTa and amino acids; (5) HYTa, chitin and amino acids; (6) HYTa, chitin, chitosan and amino acids; (7) HYTa, chitosan,
25 glucosamine and amino acids; (8) HYTa, chitosan and glucosamine and (9) HYTa, chitin, chitosan, glucosamine and amino acids, the latter being particularly preferred.

When HYTa is grown in the presence of chitin, chitosan and/or amino acids it may contain residual chitin,
30 chitosan and/or amino acids. Under such circumstances, the HYTa culture constitutes the disclosed microbial composition and can be applied directly to soil, seed, seedlings or plant foliage. Alternatively, one or more of the second

components can be added to supplement the second components in the composition or to change its composition.

As used herein, the term "amino acids" refers to a composition containing two or more amino acids. Amino acids
5 include tryptophan, histidine, threonine, tyrosine, valine, methionine, isoleucine, leucine, phenylalanine, lysine, aspartic acid, cysteine, glutamic acid, glutamine, serine, glycine, alanine, proline, asparagine and arginine. In preferred embodiments, amino acids are provided by use of
10 HYTb (See below).

As used herein, the term "chitin" refers to a biopolymer consisting predominantly of repeating units of beta-1-4-linked N-acetyl-D-glucosamine. Chitin is found in the natural environment as a primary structural material of
15 the exoskeleton of animals such as *Arthropoda*, e.g., crustaceans, insects, spiders, etc., *Mollusca*, e.g., snails, squid, etc., *Coelentara*, e.g., organisms such as hydroids and jellyfish, and *Nematoda*, such as unsegmented worms. Chitin is also found in various fungi including members of the
20 genus *Fusarium*. Chitin can be extracted from these natural sources by treatment with alkali, or by a biodegradation process. The molecular weight of chitin varies depending on its source and method of isolation. In preferred
embodiments, the chitin is derived as a solid from the
25 biodegradation of chitin containing *Arthropods* as described in the Bioderpac applications. It is preferred that the chitin have a diameter of about 50 to 75 microns to facilitate its application via drip and spray irrigation systems.

30 As used herein, the term "chitosan" is a polysaccharide consisting predominantly of repeating units of D-glucosamine. Chitosan is obtained by deacetylation of chitin. The degree of deacetylation as compared to chitin

is preferably greater than 50%, 60%, 70%, 80%, 85%, 90% and 95%. It is preferred that the level of deacetylation be sufficient to render the chitosan water soluble at acidic pH. The molecular weight of chitosan varies depending on its source and method of isolation. Chitosan includes chitosan oligomers. In preferred embodiments, chitosan is precipitated at pH 9.0 from the aqueous fraction obtained from the biodegradation of chitin containing *Arthropods* such as described in the Bioderpac applications.

As used herein, the term "chitosan oligomer" refers to chitosan having 2 or more repeating units of D-glucosamine and, in the case of incomplete deacetylation of chitin, one or more units of N-acetyl-D-glucosamine. In preferred embodiments, the chitosan oligomers are derived from the aqueous fraction generated in the biodegradation of chitin containing *Arthropods* such as described in the Bioderpac applications. In some embodiments chitosan oligomers are used as the second component of the microbial composition.

As used herein, the term "glucosamine" refers to an amino monosaccharide. In preferred embodiments it is the sugar residue that forms the backbone of the biopolymers chitin and chitosan. Glucosamine is present in the aqueous fraction generated during the biodegradation of chitin containing *Arthropods* such as described in the Bioderpac applications. Glucosamine induces plants to make chitinase as a defense to chitin containing pathogens.

HYTb and HYTc

As used herein, the term "HYTb" refers to the aqueous fraction and "HYTc" refers to the solid fraction obtained from the biodegradation of *Arthropods* such as shrimp waste. derived from the biodegradation or chitin containing

Arthropods such as described in US Patent Application Serial No. 61/289,706, filed 12/23/09 entitled "Biodegradation of Crustacean By-products", US Patent Application Serial No. 61/299,869, filed 1/29/10 entitled "Biodegradation Process and Microbial Composition" and US Patent Application Serial No. 61/355,365 filed June 16, 2010 entitled "Biodegradation Process and Composition" each of which are incorporated by reference herein in their entirety.

Briefly, in the arthropod biodegradation process a microbial composition is used to degrade the arthropod or waste components of the arthropod. It is a lactic acid fermentation process. The microbial composition contains microbes that produce enzymes that can degrade the chitin containing components of the arthropod to chitin, chitosan, N-acetyl glucosamine and glucosamine. It also contains microbes that produce enzymes that can degrade proteins and fats to produce amino acids and lipids. A preferred microbial composition for arthropod degradation is referred to as HQE. HQE was deposited with the American Type Culture Collection (ATCC) Manassas, VA, USA on April 27, 2010 and given Patent Deposit Designation PTA-10861.

In a preferred embodiment, the marine arthropod is a crustacean and the preferred crustacean is shrimp. Shrimp by-product comprises shrimp cephalothorax and/or exoskeleton.

In the biodegradation process, it is preferred that the fermentation be facultative aerobic fermentation. It is also preferred that the fermentation is carried out at a temperature of about 30°C to 40°C. The pH is preferably less than about 6, more preferably less than about 5.5. However, the pH should be maintained above about 4.3. The fermentation is carried out for about 24-96 hours. In some embodiments, the fermentation is carried out for about 24-48

hours and more preferably 24-36 hours. These fermentation times are far shorter than the typical prior art fermentation times of 10 to 15 days to achieve substantially the same amount of digestion, albeit without detectable
5 formation of chitosan and glucosamine.

The separation of the mixture is preferably by centrifugation. (e.g. about 920 g). Gravity separation can also be used but is not preferred because of the time required to achieve separation.

10 The mixture separates in to three fractions: solid, aqueous and lipid. The solid fraction comprises chitin and is designated HYTc. The aqueous fraction comprises protein hydrolysate, amino acids, chitosan and glucosamine and is designated HYTb. The lipid fraction comprises sterols,
15 vitamin A and E and carotenoid pigments such as astaxanthine.

It is preferred that HQE be used in the biodegradation process. In other embodiments, it is preferred that previously prepared HYTb be added to HQE or
20 the fermentation broth. As described above, HYTb contains amino acids, chitosan, glucosamine and trace elements including calcium, magnesium, zinc, copper, iron and manganese. HYTb also contains enzymes such as lactic enzymes, proteases, lipases, chitinases, lactic acid,
25 polypeptides and other carbohydrates. HYTb can also contain dormant microorganisms from a prior biodegradation process. Such microorganisms can become reactivated and, in combination with HQE, contribute to a more robust biodegradation process as compared to when HQE is used by
30 itself as otherwise described herein

More particularly, the process includes the following steps:

- 5 a. Activation of the microbial cells in a sugar base solution to enhance its growth and the biomass formation.
- b. Milling of the shrimp by-products (cephalothorax and exoskeleton) to make a homogeneous paste.
- c. Homogeneous mixing of the shrimp by-product paste with at least 10% of the activated inoculum.
- 10 d. Adjustment of the pH values to less than 6.0 in the mixture using a citric acid solution to inhibit the growth of micro organisms and to promote the development of microbial cells that constitute the inoculum.
- 15 e. Fermentation of the mixture in a non continuous agitated system at temperatures within a range of 30 to 40°C at least for at least 96 hours maintaining pH at less than 5.0. The pH is monitored periodically. If the pH rises above 5.0,
20 a citric acid buffer is added in an amount to maintain the pH below 5.0.
- f. Centrifugation of the ferment to separate the three principal fractions: chitin, liquid hydrolysate and pigmented paste.
- 25 g. Rinsing of the crude chitin and recollection of the rinse water to recuperate fine solids or minerals.
- h. Drying of the chitin and storage.
- i. Drying and storage of the liquid hydrolysate.
- 30 j. The pigmented paste (lipid fraction) is stored in closed recipients for conservation.

The process and operational fundamentals are better understood with reference to the following detailed description.

Activation of microbial cells

5 A microbial composition as disclosed herein is used as inoculum. The inoculum of HQE has a concentration of microbes of about 2.5 to 3.0% (w/v). HQE is activated by dilution to 5% in sugar cane solution (3.75% final concentration of sugar cane), and incubated at 37 °C for 5
10 days. HYTb (10 ml per liter of culture) is preferably added to provide a source of minerals and naturally derived amino acids. The cellular growth of the microorganisms was estimated by optical density measured at 540 nm. The activation is complete at an optical density of about 1.7.
15 The concentration of microbes after activation is about 1.9 to 3.0% (w/v).

Preparation of samples

The shrimp by-products samples are obtained from shrimp processing plants. Slightly thawed and minced residue
20 (1500 g by batch) is mixed with 99 grams of sugar cane (final concentration 6.6% wt%) and 85.5 ml of activated HQE 5% (v/w) (optical density of cell = 1.7). Then the pH is adjusted to 5.5 using 2 M citric acid.

Fermentation control

25 The mixture is incubated at 36 °C with a non continuous agitation for 96 h. During the fermentation process, the pH is monitored by using a potentiometer, and the total titratable acidity (TTA, %) was determined by titration with 0.1 N NaOH until a pH of 8.5 is obtained. The
30 TTA is expressed as a percentage of lactic acid.

Conditions of separation

The fermentation product is a viscous silage which has an intense orange color, due to the astaxanthine

presence. The ensilage is centrifuged (5 °C) at 1250 rpm (930g) for 15 min to obtain the chitin, the liquid hydrolysates, and the pigment paste. The upper phase (pigment paste) is separated manually. The liquid hydrolysates are separated by decantation, and the sediment that constitutes the raw chitin is washed with distilled water to separate fine solids. The resulting liquid is collected and dried. The raw chitin, liquid hydrolysates and fine solids are dried at 60 °C. All the fractions are stored to protect them from light.

Other microbial compositions for the production of HYTb and HYTc are set forth in the following Table 2.

Table 2
Culture Composition

Microorganism	1	2	3	4	5	6	7	8	9	10
<i>Bacillus subtilis</i>	X	X	X	X		X	X	X		X
<i>Bacillus cereus</i>	X	X		X		X		X		X
<i>Bacillus megaterium</i>	X	X								
<i>Azotobacter vinelandii</i>	X	X		X		X		X		X
<i>Lactobacillus acidophilus</i>	X	X	X	X	X		X	X	X	
<i>Lactobacillus casei</i>	X	X		X	X			X	X	
<i>Trichoderma harzianum</i>	X	X	X	X		X	X	X		X
<i>Rhizobium</i>	X	X		X		X		X		X

<i>japonicum</i>										
<i>Clostridium pasteurianum</i>	X	X			X	X			X	X
<i>Bacillus licheniformis</i>	X	X	X		X	X	X		X	X
<i>Pseudomonas fluorescens</i>	X	X	X		X	X				
<i>Bacillus thuringiensis</i>	X					X	X	X	X	X
<i>Streptomyces</i>	X				X	X	X	X	X	X
<i>Nitrobacter</i>	X						X	X	X	X
<i>Micrococcus</i>	X						X	X	X	X
<i>Proteus vulgaris</i>	X						X	X	X	X

These microorganisms are preferably derived from HQE and are referred to as *Bacillus subtilis* ((SILoSil® BS), *Bacillus cereus* (Bioderpac, 2008), *Bacillus megaterium* (Bioderpac, 2008), *Azotobacter vinelandii* (Bioderpac, 2008), *Lactobacillus acidophilus* (Bioderpac, 2008), *Lactobacillus casei* (Bioderpac, 2008), *Trichoderma harzianum* (TRICHOSIL), *Rhizobium japonicum* (Bioderpac, 2008), *Clostridium pasteurianum* (Bioderpac, 2008), *Bacillus licheniformis* (Bioderpac, 2008), *Pseudomonas fluorescens* (Bioderpac, 2008), *Bacillus thuringiensis* strains HD-1 and HD-73 (SILoSil® BT), *Streptomyces* (Bioderpac, 2008), *Micrococcus* (Bioderpac, 2008), *Nitrobacter* (Bioderpac, 2008) and *Proteus* (Bioderpac, 2008). Each of these organisms can be readily isolated from HQE and recombined to form the disclosed

microbial composition to degrade arthropods to make HYTb and HYTc.

HYTb contains amino acids (about 12 wt%), chitosan (about 1.2 wt%), glucosamine (about 1 wt%) and trace elements (about 6 wt%) including calcium, magnesium, zinc, copper, iron and manganese. It also contains enzymes such as lactic enzymes, proteases, lipases, chitinases among others, lactic acid, polypeptides and other carbohydrates. The specific gravity of HYTb is typically about 1.050-1.054. The average amino acid content in HYTb for certain amino acids is set forth in Table 2.

Table 3

Amino acid profile dry powder hydrolysates
(mg per g dry weight)

Amino acid	Dry powder hydrolysates
Aspartic acid	38
Glutamic acid	39
Serine	16
Histidine	9
Glycine	28
Threonine	14
Alanine	36.1
Proline	25.8
Tyrosine	70
Arginine	22.2
Valine	20
Methionine	16.4
Isoleucine	18.3
Tryptophan	3.1
Leucine	23
Phenylalanine	39
Lysine	13

Amino acid	Dry powder hydrolysates
Total	431

In some embodiments, HYTb can constitute a second component that is either combined with HYTa or used separately as a soil amendment and/or as a foliage spray.

5 The primary component of HYTc is chitin. It has an average molecular weight of about 2300 daltons and constitutes about 64 wt% of the composition. About 6 % of HYTc contains minerals including calcium, magnesium, zinc, copper, iron and manganese, about 24 wt% protein and 6%
10 water. It has a specific gravity of about 272 Kg/m³. In some embodiments, HYTc can constitute a second component that is either combined with HYTa or used separately as a soil amendment and/or as a foliage spray.

 HYTa is preferably used with HYTb and HYTc either in
15 combination or separately as a soil amendment or foliage spray.

 The microbes in HYTa require the trace elements calcium, magnesium, sulfur, boron, manganese, zinc, molybdenum, iron, copper, sodium, and silicon. These
20 important trace elements can be often obtained from toxic chemical reactions which are not suitable for organic certified products. Accordingly, it is preferred that these trace elements be obtained from an organic source such as HYTb and/or HYTc.

25

Activation of HYTa

 The aforementioned microbial compositions can be used to treat soil, seeds, seedlings and/or plant foliage. However, HYTa is first activated before use.

In preferred embodiments, HYTa is activated by incubating an inoculum of HYTa in an aqueous solution for 24-168 hours to allow the microbes to grow and reproduce before being used in the process of treating soil, seeds, seedlings and/or plant foliage. The conditions of the incubation influence the overall initial properties of HYTa.

In one embodiment, an inoculum of HYTa is diluted with water in a ratio of 1/100 and allowed to incubate at a temperature of approximately 36°C at a pH of 6.8-7.1 for about 24 to about 168 hours (7 days). HYTb can optionally be used during this activation. The nitrogen fixing microbes *Azotobacter vinelandii* and *Clostridium pasteurianum* proliferate under reduced nitrogen growth conditions. In addition, as the oxygen concentration decreases, *Lactobacilli*, including *Lactobacillus acidophilus* and *Lactobacillus casei*, proliferate. The colony forming units (CFUs) for some of the bacteria in activated HYTa are set forth in Table 3:

Table 4

<i>Azotobacter vinelandii</i>	101,050,000 Cfu/mL
<i>Clostridium pasteurianum</i>	104,275,000 Cfu/mL
<i>Bacillus subtilis</i>	1,100,000 Cfu/mL
<i>Bacillus cereus</i>	25,000 Cfu/mL
<i>Bacillus megaterium</i>	10,000 Cfu/mL
<i>Lactobacillus</i>	500,000 Cfu/mL
<i>Nitrobacter</i>	5,000 Cfu/mL
<i>Nitrosomonas</i>	2,500 Cfu/mL
Total	206,967,000Cfu/mL

The HYTa obtained after this incubation retains the beneficial properties of HYTa but is particularly suited as a soil amendment for treatment of nitrogen-depleted soils given the nitrogen-fixation capabilities of *Azotobacter vinelandii* and *Clostridium pasteurianum*.

If soil pathogens such as filamentous fungi from the genus *Fusarium* or nematodes are present, or believed to be present, HYTa can be activated under substantially the same conditions but in the presence of chitin. The chitin stimulates the expansion of the chitin responsive microbes such as *Pseudomonas fluorescens*, *Trichoderma harzianum*, *Bacillus thuringiensis*, *Streptomyces* sp., *Nitrobacter* sp., *Micrococcus* sp., and *Bacillus subtilis*. HYTa obtained under these conditions has an antifungal, fungicidal, antinematode, nematodicidal and insecticidal properties to the extent such pathogens contain chitin. Such microbial compositions can be applied directly to the soil or to seed, seedlings and/or plant foliage. Such microbial compositions also have the ability to fix nitrogen as in the aforementioned incubation in the absence of chitin.

In addition to incubating with chitin, HYTa can be activated with chitin and amino acids. A preferred source of chitin is HYTc. When HYTc is used the protein and minerals in HYTc are also present during the activation.

Further, HYTa can be activated in the presence of amino acids and chitosan. A preferred source of amino acids and chitosan is HYTb. When HYTb is used glucosamine and the other components of HYTb are also present during the activation.

Optionally, HYTa can be incubated with chitin, amino acids and chitosan. A preferred source of chitin is HYTc. A preferred source for amino acids and chitosan is HYTb. When

HYTb and HYTc are used the other components in these formulations are also present during activation.

Use of Activated HYTa

5 Activated HYTa can be used alone or in combination with other components such as chitin, chitosan, glucosamine and amino acids to treat soil, seed, seedlings or foliage. In some embodiments, combinations of these components can be applied as a mixture. In other embodiments, they can be
10 applied separately. In still other embodiments, the components can be applied at different times.

 In one embodiment, activated HYTa can be applied to soil, seeds or seedlings, or used in foliar applications by direct application to foliage. However, when plant
15 pathogens are present, it is preferred that microbial composition comprises activated HYTa, chitin and/or chitosan. Alternatively, the HYTa can be activated in the presence of chitin. Chitosan is known to have bactericidal, fungicidal, and antiviral properties, as well as its ability
20 to stimulate plant growth and to induce plant resistance to pathogens. In other embodiments, glucosamine is a part of the microbial composition

 In a preferred embodiment, the activated HYTa alone or in combination with chitin (preferably HYTc) and/ or
25 chitin, chitosan, and amino acids (preferably HYTb and HYTc), is applied to soil, seeds, seedlings and/or foliage. It is preferred that HYTa be used in combination with chitin, chitosan, glucosamine and amino acids. HYTc is the preferred source of chitin while HYTb is the preferred
30 source of chitosan, glucosamine and amino acids. However, the components of the microbial composition namely HYTa, chitin, chitosan, glucosamine and amino acids can be applied separately or in any combination or sub-combination. They

can be applied at the same time or sequentially, in any given order. However, the preferred mode of application is to initially apply all at the same time. The application of the foregoing components provide for the direct treatment of plant pathogens, the induction of plant pathogen resistance pathways, and the nourishment of the HYTa microbes, the indigenous nonpathogenic soil flora, and the plant.

When soil is initially treated with a microbial composition comprising activated HYTa alone, the microbes present in the composition have an opportunity to populate the soil and to alter its taxonomic composition. In some situations, the initial colonization by HYTa provides little or no nutrients to the plant. In such instances, it is important to maintain a nutrient reserve to sustain both the growth of the microbes while colonizing the rizosphere and the growth of the plants in the soil. It may be necessary to repeat the application of HYTa, depending on the plant's growth cycle and nutritional regime. In other cases, it may be sufficient to provide additional applications of amino acids, chitin and/or chitosan, eg. HYTb and HYTc, to the previously treated soil.

When HYTa is used in combination with, for example, HYTb and HYTc, addition nutrients are available to the HYTa microbes and the plants present in the treated soil.

Table 5 sets forth a typical fourteen week program for the application of HYTa, HYTb and HYTc to drip irrigated crops cultivated in soil. The values are per hectare. For HYTa and HYTb, the values represent liters per week. For HYTc, the values represent kilograms per week.

Table 5

Lts/kg/ Week	W1	W2	W3	W4	W5	W6	W7	W8	W9	W10	W11	W12	W13	W14
HYT-A	3	0	0	1	0	1	0	1	0	1	0	1	0	1

HYT-B	10	5	0	3	2	3	2	3	2	3	2	3	2	3
HYT-C	1			1				1				1		

The pulse in which the microbial composition is injected to the irrigation system should be one in which the microbial composition is able to reach the root system and stay there over night while the system is off. For maximum performance of HYTc, it should be applied at the same time as a mixture with HYTa. The protocol should be continued as long as the plant continues in production. This protocol covers all plant stages including germination, root formation, plant growth, flowering, fruit setting, fruit formation harvesting and re-harvest. This protocol is designed for maximum yield potential covering nutritional aspects, biostimulation aspects and protection against diseases such as nematodes and fungi.

The process can be carried out by contacting soil to form a treated soil. In some cases the process is repeated. In some cases, plants, seedlings or seeds are already present in the soil prior to treatment with the microbial composition. In other cases, plants, seedlings or seeds are transplanted to the soil after treatment with the microbial composition.

In general, before application the number of hectares or acres to be treated is determined. Then the recommended amount of activated HYTa per hectare or acre is multiplied by the area to be treated and diluted in sufficient water to irrigate or spray the soil or crop on the area to be treated. The same procedure can be followed for liquid HYTb. HYTc, being a solid, can be applied directly as a solid or as a suspension in water. HYTc is preferably ground to micron size particles prior to use.

The process can be carried out with infertile soil. Such soils generally are those where at least one of low cation exchange capacity, low water holding capacity, low organic matter content and low levels of available nutrients is present. In general, infertile soil does not support vigorous plant growth and/or produces low crop yields.

For non-soil systems such as hydroponics, the same protocol applies but with a daily distribution following the ferti-irrigation program.

10 The microbial compositions can be used in connection with any plant including but not limited to alfalfa, banana, barley, broccoli, carrots, corn, cucumber, garlic, grapes, leek, melon, onion, potato, raspberry, rice, soybean, squash, strawberry, sugarcane, tomato and watermelon.

15 When applied as a soil amendment, the microbial composition containing HYTa, chitin, amino acids and chitosan enhances crop production on average about 25%-55% as compared to the 15-25% increase in crop production observed for E2001. From Karl Co. SA de CV, Navojoa, 20 Sonora, Mexico.

The microbial composites can also result in a decrease in the amount of chitin used. For example, chitin has been used as a soil amendment in the prior art. Typically, about 600 kg of chitin were used per hectare. 25 However, beneficial effects of such use were not observed for up to six months. When HYTa was activated in the presence of chitin and then combined with chitin and applied as a soil amendment, beneficial effects were observed after seven days with the use of only 4-6 kg of chitin per 30 hectare.

Although the disclosure is directed primarily to the use of the disclosed microbial compositions for agricultural applications, such compositions or their components and

processes can also be used in horticultural applications to improve the production of foliage and flowers and decrease the use of conventional insecticides and fungicides.

When activated HYTa is applied to soil, seed seedling or foliage it forms treated soil, treated seed, treated seedling, treated foliage and treated plants. HYTa is a novel microbial composition. Therefore the soil, seed, seedling, foliage and plants treated with HYTa are also novel.

Treated soil is defined as soil that contains one or more microbes that are unique to HYTa dispersed within the treated soil. Such microbes can be detected in the treated soil genetically by using a BioChip that detects microbial populations based on DNA. See e.g. US Patent Publication 2007/0015175, incorporated herein by reference. Other methods, such as PCR, which know to those skilled in the art can also be used. Microbes in HYTa that are particularly preferred are *Bacillus subtilis* (SILOsil® BS), *Bacillus thuringiensis* strain HD-1, *Bacillus thuringiensis* strain HD-73 (SILOsil® BT) and *Trichoderma harzianum* (TRICHOSIL) each of which can be isolated from the HYTa deposit or obtained from Biotechnologia Agroindustrial S.A. DE C.V., Morelia, Michoacan, Mexico. *Trichoderma harzianum* (TRICHOSIL) is most preferred as it is important during the activation of HYTa in that it causes inter-component synergies among the other microbes in HYTa. Identification of one or more of these microorganisms can be further combined with the identification of other microbes in HYTa, if necessary, to confirm the presence of HYTa or that HYTa was present. *Bacillus subtilis* (SILOsil® BS): ATCC designation PTA-12153, deposited on October 7, 2011. *Bacillus thuringiensis* strains HD-1 and HD-73 (SILOsil® BT): ATCC designation PTA-12967, deposited on May 31, 2012. *Trichoderma harzianum* (TRICHOSIL): ATCC designation PTA-12152, deposited on

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October 6, 2011.

Treated seed, seedlings, foliage and plants are similarly defined. In these cases, the microbes of HYTa are found on the surfaces of the treated seed, seedlings, foliage and plants.

5 As used herein, the term "consisting essentially of" in connection with HYTa, HYTb and HYTc means any of HYTa, HYTb and/or HYTc alone or in combination without additional microbes.

Example 1

10 The following example compares the growth of Persian cucumber plants using HYTa, HYTb and chitosan as compared to a control which was not treated with HYTa, HYTb and chitosan.

15 During the development of seedlings of Persian cucumber, seeds were incubated for three hours in a mixture of 1 liter of water and 250 grams of HYTc. A bag of peat moss and 250 grams of micronized 200 mesh (approximately 75 micron diameter) chitin (HYTc) per bag of peat moss were blended. The seeds were planted in the peat moss/chitin mixture at 18-24[deg.]C. The
20 plant development after five days following treatment with HYTc was comparable to 9 days of development without the treatment.

The treated and control seedlings were transplanted into 1 hectare of soil in a green house. The HYTa and control soil were treated as set forth in Table 5

25

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Table 6

	HYT-A	Control
Nitrogen fertilizer	150 Kg	280 Kg
Potash	160 Kg	250 Kg
Calcium	80 Kg	130 Kg
Phosphorous	200 Kg	320 Kg
Magnesium	20 Kg	45 Kg
Trace elements	10 liters	22 liters
Fungicides	0	20 liters
Insecticides	0	16 liters
agricultural soap made from palm and olive oil	10 iters	0

The soil containing the HYTa treated seedlings was treated with 2 liters HYTa and 7 liters of HYTb over time.

5 HYTa was diluted in 200 liters of water and activated without the presence of HYTb or HYTc.

At week two, one liter of HYTa and three liters of HYTb were applied to the soil and two liters of HYTb were applied to the foliage of the HYT treated plants.

10 There was a significant increase in the yield of cucumbers over the control. The control plants produced 3,000 twenty five pound boxes while the HYT treated plants produced 4,300 boxes. Accordingly, this example demonstrates a significant increase in yield using HYT and a decrease in

the amount of fertilizer, insecticides, fungicides and other components otherwise needed.

Example 2

Septoria leaf and early blight as well as infection of Roma and beefsteak tomatoes with *Phytophthora infestans* can be treated by the protocol set forth in Table 7. All values are per hectare.

10 **Table 7**

	Start	Per day	Duration	Application
HYTa	3 litres	0	10 days	Spraying
HYTa	2 litres	0	10 days	Drip System
HYTc	20 Kg	2 Kg	10 days	Spraying
HYTc	500 grms	0	0	Drip System
HYTb	1 liter	1 liter	10 days	Spraying

HYTa was diluted in 200 liters of water and activated with HYTc.

This treatment resulted in control of these
15 infections.

Example 3

10 acres of Roma tomatoes were treated with 4 liters of HYTa, 10 liters of HYTb and 30 pounds of chitin.

20 The application protocol was as follows for 10 acres:

Table 8

Lts/Lb/Week	W1	W2	W3	W4	W5	W6	W7
*HYT-A	3	0	0	0	1	0	0
**HYT-A	2	0	0	0	1	0	0
*HYT-B	6		5		5	0	
**HYT-B	4		2		2		
*HYT-C	5				5		
**HYT-C	10		5		5		

* Irrigation system: Spraying (foliage)

** Irrigation system: Drip Tape

5

The values are in liters for HYTa and HYTb and pounds for HYTc. The crop yield was 46 tons of tomatoes per acre as compared to 31 tons per acre for the control. This is a 36% increase in yield.

10

Example 4

Root-Knot nematode *Meloidogyne* spp. and white mold disease caused by *Sclerotinia sclerotiorum* were identified as problematic for the growth of carrots. Figure 2A shows the foliage and carrots obtained from such soil.

15

The following protocol was used to treat a hectare. One Kg of HYTc was applied to the soil at the time of transplantation. Two weeks later 1 Kg of HYTc and 1.5 liters of HYTa was applied. Two weeks later 2 Kg of HYTc and 1 liter of HYTb was applied. Thirty days later 1.5 Kg of HYTc, 1 liter of HYTb and 1 liter of HYTa were applied.

20

The root galls caused by the nematode infection was no longer present on the carrots after the treatment. The cottony soft rot caused by white mold was also absent from the carrots after treatment.

25

Example 5

HYTa, HYTb and HYTc can be used to eradicate and control ROYA (*Puccinia dracunculina*) on Tarragon (*Artemisia dracunculus* L.). A total of 6 liters of HYTa, 15 liters of
5 HYTb and 900 grams of HYTc were applied to each hectare.

The following protocol was used:

Table 9

Product	Dosage	Round time	Application
HYTa	2 liters	5 days	spraying
HYTb	5 liters	5 days	spraying
HYTc	300 grms	5 days	spraying

10

The protocol was repeated twice. This treatment reduced damage from ROYA on treated foliage.

Example 6

15 This example discloses a summary of tests carried in cooperation with and under the supervision of the Centre International of Maize and Wheat Improvement Center (known and referred to herein as the "CIMMYT")
<http://www.cimmyt.org/>.

20 This example presents the final data from the harvest of the different treatments. CIMMYT staff performed the collection of samples in accordance with its scientific methodologies and information.

25 These tests were designed to demonstrate the following key benefits of using HYTa alone or in combination with HYTb: (1) the ability to maintain high-performance growth with different regimes of fertilizer and minerals,

(2) improving the performance of the system through the use of HYTa or HYTa in combination with HYTb, and (3) the ability to restore soil health soil and increased the levels of fertility through the repeated use of HYT programs.

5 The objective of the test was to determine the effect of the levels of tillage and the handling of straw in two different environments of soils (neighborhood and alluvium), investigate the efficiency of the different forms, types and doses of mineral fertilizers in combination with HYT of
10 Agrinos to make more efficient use of these inputs in order to increase the profitability of the cultivation of wheat to the producer.

Areas of test

 These trials were performed in an agricultural field
15 associated with the assignee which has been used widely for the development of soil remediation products, as well as for the production of cash crops. This agricultural field was treated over the last eleven years with E2001 and related products from Karl Co. SA de CV, Navojoa, Sonora, Mexico and
20 more recently with HYTa and HYTb

 This area of trials is identified by CIMMYT under the coupon code Z 702 Module Agirnos-CIMMYT and is in the District of irrigation Nr. 38, module 4, section 15, rolls of irrigation 1049-0 and 1115-0.

25 One of the main attributes of HYT™ products is its ability to improve (instead of degrade) agricultural soils with continuous use. In order to demonstrate this attribute of the HYT™ product, the trials have included a test area which was not treated with mineral inputs, E 2001 or any HYT
30 product. The performance of plants of the crop in this area depends entirely on the state of the soil prior to planting.

Other information

-Types of crops: wheat

-Variety: ATIL (durum wheat)

-Sowing date: 23 December in dry soil and wet soil in areas 2 and 1 of Figure 3. Planting was delayed until January 14, of the following year due to a flood caused by an irrigation problem in the adjoining plot.

-Date of harvest: May 20-23, (approximately 4 months after planting)

-Size of the test area: 15 hectares

-Mineral fertilization: was used as the basis for fertilization which is considered as the best practice of deossification of mineral nutrients NPK generally accepted in the region (BNFP = Best nitrogen fertilizer practice).

Table 10

Mineral fertilizers and HYTTM protocols

Treatment	Description	Initial Application	Second Application	Third Application
Treatment 1	Control area; no fertilizer	0 units NPK, 0 liters of HYTa or HYTb	0 units of NPK, 0 liters of HYTa orHYTb	0 units of NPK, 0 liters of HYTa or HYTb
Treatment 2	50% BNFP + HYTa and HYTb	103 units of N, 52 units of P, 1 liter of HYTa	1 liter of HYTa; 1 liter of HYTb	1 liter of HYTa; 1 liter of HYTb
Treatment 3	HYTa + HYTb	1 liter of HYTa	1 liter of HYTa; 1 liter of HYTb	1 liter of HYTa; 1 liter of HYTb
Treatment 4	100% BNFP	149 units of N,	61 units of N	

52 units of
P

In addition to the main protocols described above, some areas were tested and harvested separately with some additional component, with the aim of obtaining an extra point of reference and expand the possibilities for analysis. The designated test was as follows:

TRT 5: Biological treatment HYT more 100% traditional mineral fertilization programme: initial application: 1 litre of HYT + 103-52-0 (NPK), second application 1 litre of HYT + 1 litre of HYT B + 61-0-0 (NPK), third implementation 1 liter of HYT B. This further treatment was recommended by CIMMYT in order to observe the behavior of the traditional program of more complete mineral nitrogen program HYT a + b in the performance of the grain of wheat. Only an area of 4 rows was dedicated to this treatment and the information was collected only by CIMMYT staff.

A diagram of the test area is shown in Figure 1 where "TRT" refers to the above identified treatment.

External factors in the test area

Some areas of the test zone were compromised and suffered impairment by external factors. The results of these areas have been excluded from the final results of the harvest to allow a reliable comparison. Reference is made to Figure 2. These external factors were as follows:

Area highlighted 1 (Zone 1): The wheat variety used is very susceptible to "Chahuistle". Due to the proximity of areas 1 and 5 to high voltage electric lines, the plane could not apply the product on these areas and consequently these areas suffered a higher incidence of the pathogen, causing a significant loss of performance potential.

Highlighted areas 1 and 2 (Zone 2) suffered flooding due to problems of irrigation in the surrounding plots, delaying the planting in 20 days and being affected by the "chahuistle".

5 Highlighted areas 9 and 10 (Zone 3) suffered from irregular irrigation due to the topography of the ground which makes erratic performance of the crop having low and high areas causing non-uniform irrigation that affects the average of the performance.

10

Table 11
Final data of total harvest reported for different treatments

Lot	metric	Lot	metric
	tonnes per hectarea		tonnes per hectarea
Area 1	4.1*	Area 9	5.4 / 6.6*
Area 2	4.1*	Area 10	6.2*
Area 3	7.8	Area 11	6.8
Area 4	8.3	Area 12	8.3
Area 5	4*	Area 13	8.3
Area 6	7.1	Area 14	7.7
Area 7	7.4	Area 15	8.2
Area 8	7.9	Area 16	9.0

15

*external factors affected this result

Table 12
Results of Trial

Performance			
Tonnes/Ha*	Clay*	Alluvial*	Average*
100% BNFP (Treatment 4)	7.45	7.40	7.4
50% BNFP plus HYTa and HYTb (Treatment 2)	7.40	7.20	7.8
Control (Treatment 1)	7.90	8.30	8.2
HYTa and HYTb only (Treatment 3)	8.30	6.50	8.7

*results from areas where external factors affected

5 results are not included

In comparison with the average wheat yield expected in the region the repeated historical use E 2001 and HYT[™] has contributed to the significant increase in performance of 36% as compared to standard fertilization only. See
 10 Figure 3. This happened without adding any additional element of NPK or HYT during this agricultural cycle given that previous E 2001 and HYT applications had already restored the activity and biodiversity of colonies of benign soil microbes creating high levels of organic matter and
 15 nutrients available in the soil.

Adding the HYTa and HYTb to the soil-plant cycle system continues the improvement of the capacity of the soil to provide nutrients to plants, increasing the capacity of biological nitrogen fixation. See Figure 3.

20 Various combinations of standard fertilization regimes, alone or in combination with HYTa and HYTb do not seem to improve the results as compared to the use of HYTa

and HYTb. This may be due to the existence of sufficient stored nutrients as biomass in the soil from previous years.

When the soil and ecosystem have sufficient available nutrients, either through FBN and/or high levels of biomass in the soil, the addition of more (NPK) fertilizer destabilized the biological balance and interfered with the patterns of absorption of nutrients from the plants, possibly changing the capacity of the cultivation of guiding their own nutritional program given the resources available in the soil and active biomass in its roots.

Example 7

Field experiments were conducted at Pantnagar India under the project entitled 'Agronomic evaluation of HYT (HYTa, HYTb and HYTc and foliar spray of Suryamin). The details are given below.

	Crop	:	WHEAT
	Design used	:	RBD
	Replication	:	3
20	Date of sowing	:	19.11.2010
	Variety	:	PBW-550
	Gross plot size	:	6.0 m x 4.0 m = 24 m ²
	Treatments	:	12
	Treatment details		
25	T₋₁ : Recommended NPK dose		
	T₋₂ : T-1+soil application of HYTa (activated for 72 hrs) @ 1L at the time of sowing		
	T₋₃ : T-1+ foliar application of HYTb @ 2L at the time of flower initiation/ panicle initiation)		
30	T₋₄ : T-1+soil application of HYTc @ 2 kg at the time of sowing		

T-5: T-1+ HYTa (activated for 72 hrs) @ 1Liter + HYTc@ 2kg/ha as soil application at sowing

T-6: T-1+ foliar application of HYTb @ at the time of flower initiation/ panicle initiation + 2L+HYTc @ 2kg/ha as soil application at sowing

T-7: T-1+HYTa (activated for 72 hrs) @1L +HYTb @ 2L at the time of flower initiation/ panicle initiation+ HYTc @2 kg/ha as soil application at sowing

T-8: ½ NPK dose + HYTa (activated for 72 hrs) @ 1L+ foliar application of HYTb @ 2L+HYTc @ 2kg/ha as soil application at sowing

T-9: T-1+ soil application of HYTa (activated for 72 hrs) @ 2L/ha+ foliar application of HYTb @ 5L+HYTc @ 5kg/ha at sowing

T-10: ½ NPK dose +HYTa (activated for 72 hrs) @ 2L+foliar application of HYTb @ 5L+HYT-C @ 5kg/ha as soil application at sowing

T-11: T-1+ 1L/ha of Shriram Suryamin as foliar application at flower initiation

T-12: T-1+1L/ha of Shriram Suryamin as foliar application each at tillering and at flower initiation

The biological, grain and straw yields are set forth in Table 13.

Table 13

Effect of different HYT organic product on biological, grain and straw yield of wheat crop.

Treatments	Biological Yield (q/ha)	Grain Yield (q/ha)	Straw Yield (q/ha)
T ₁ : Rec. NPK	82.63	32.00	50.63
T ₂ : T ₁ + HYT-A @ 1.0 l/ha	85.43	33.99	51.50

Treatments	Biological Yield (q/ha)	Grain Yield (q/ha)	Straw Yield (q/ha)
T ₃ : T ₁ + HYT-B @ 2.0 l/ha	87.50	35.30	52.20
T ₄ : T ₁ + HYT-C @ 2.0 kg/ha	76.40	31.33	45.07
T ₅ : T ₁ + HYT-A +C	87.63	37.90	49.73
T ₆ : T ₁ + HYT-B+C	84.53	34.80	49.73
T ₇ : T ₁ + HYT-A +B+C	86.93	35.80	51.93
T ₈ : ¹ / ₂ NPK+HYT-A+B+C	56.90	27.13	29.77
T ₉ : T ₁ + HYT-A +B+C (higher dose)	88.60	40.27	48.33
T ₁₀ : ¹ / ₂ NPK+HYT-A+B+C (higher dose)	58.37	28.30	30.07
T ₁₁ :T ₁ + Suryamin (one spray)	82.87	32.93	49.93
T ₁₂ :T ₁ + Suryamin (two spray)	83.57	33.17	50.40
S.Em (5%)	4.33	0.92	4.27
CD (5%)	12.68	2.71	12.54

Table 14 compares the results for grain yield for the various treatments with the different HYT components and combinations.

Table 14

	Grain Yield above 32 kilo/hectare	CD 2.71	Std. Error 0.92
Control	0		
HYTa	1.99	NS	*
HYTb	3.3	*	*
HYTc	-0.67	NS	NS
HYTa + c	5.9	*	*
HYTb + c	2.8	*	*
HYTa + b + c (1L/2L/2Kg)	3.8	*	*
HYTa + b +c (high dose of a, b and c: 2L/5L/5Kg)	8.27	*	*

NS = Not statistically significant as compared to control

5 * = Statistically significant as compared to control

As can be seen, the separate use of HYTa and HYTb improved grain yield by 1.99 and 3.3 kilo per hectare respectively while the use of HYTc alone caused a decrease in yield. When HYTa was combined with HYTc the yield increase was 5.9 kilo per hectare which is greater than the sum of the results when used separately. The use of HYTb and

HYTc resulted in an increase of 2.8 kilo per hectare while the use of HYTa, HYTb and HYTc caused an increase of 3.8 kilo per hectare. The greatest increase in grain yield was observed for HYTa, HYTb and HYTc used at the higher doses indicated. This resulted in an increase of over 25% in grain yield over the control, i.e. an 8.3 kilo per hectare increase.

Example 8

This example sets forth results of the growth of squash in infertile soil.

In these experiments squash seedlings were planted in 5 gallon pots containing infertile "Superstition" sand. The combinations of HYT A, B, and C were applied as set forth in Table 16. The seedlings were planted December 21 and harvesting began January 20 and continued through February 27 of the following year.

Table 15

Date	Treatment rates applied
December 16	200 mL activated HYT A, 10 mL HYT B, and 200 g HYT C per pot.
December 23	10 mL HYT B per pot.
December 30	5 mL HYT B per pot.
January 5	5 mL HYT A and 5 mL HYT B per pot.
January 19	10 mL HYT B per pot.
January 27	10 mL HYT A and HYT B per pot.
February 3	10 mL HYT B per pot.
February 10	10 mL HYT A and 10 mL HYT B
Feb. 17	Applied 5 mL HYT A and 5 mL HYT B to each pot.
Feb. 25	Applied 5 mL HYT B to each pot.

Date	Treatment rates applied
March 1	Applied 5 mL HYT A and 5 mL HYT B to each pot.

The results are set forth in Table 16.

Table 16

Treatment	Yield (g/pot)
Control	0
HYTa	251.6
HYTb	137.9
HYTc	62.6
HYTa + HYTb	472.1
HYTa + HYTc	0
HYTb + HYTc	0
HYTa + HYTb + HYTc	62.3
Stat. A	**
B	**
C	NS
A*B	NS
A*C	**
B*C	NS

5

NS is not significant at $P < 0.05$.

*, ** are statistically significant at
 $P < 0.05$ and $P < 0.01$, respectively.

As can be seen, there was a substantial increase in
 10 yield of zucchini squash when HYTa and HYTb were separately
 used and when HYTa and HYTb were used in combination.

Example 9

The following protocol was used to treat melons.

Table 16

PRODUC T	STAGE ▶	BEGINNING			FLOWERING			MATURATI ON			15 DAYS AFTER			Read y to Harv est			
	APPLI CA- TION	Dose (Kg or Lt / Ha)															
HYT-a	▼																
	Groun d	3	0. 3	0.2 5				1			0.5		0. 25	0. 25			
HYT-b	Folia r			1		1		1		1		1		1			
	Groun d	3	1	1				3			2		2	2			

The results are set forth in Table 17 and Figure 4.

5

Table 17

WITH HYTa and HYTb TREATMENT							
ACCUMULATED				INFORMATION PER HA			
SIZE	PIEC ES	LBS	BOXE S		PIEC ES	LBS	BOXE S
9	299	1,289 .860	33		7,40 7	31,958. 395	823
12	838	2,859 .835	70		20,7 90	71,007. 580	1,73 3
15	543	1,719 .652	36		13,5 93	43,059. 160	906
18	241	604.1 31	13		6,07 4	15,266. 914	337

TOTAL	1,92	6,473	153	47,8	161,292	3,79
	1			64		9

WITHOUT TREATMENT							
ACCUMULATED				INFORMATION PER HA			
SIZE	PIEC ES	LBS	BOXE S		PIEC ES	LBS	BOXE S
9	181	727.9 36	20		4,48 1	18,026. 691	498
12	493	1,650 .330	41		12,2 47	41,030. 815	1,02 1
15	463	1,489 .587	31		11,5 68	37,168. 321	771
18	176	465.5 97	10		4,43 2	11,725. 728	246
TOTAL	1,31	4,333	102		32,7	107,952	2,53
	3				28		6

PERCENTAGE	46.2	49.8
INCREASE PER HA	5%	1%

Example 10

Squash were grown according to the following protocol.

5

Table 18

PRODUCT	APPLICATION	DOSE PER HECTARE					
		03/07	03/10	03/18	04/04	04/11	04/18
Hyt-a	Soil	2			1		

Hyt-b	Foliar		1	1	3	1	1
	Soil	1	1	1		1	1

The results are set forth in Table 19

Table 19

SIZE	BOXES/HA		DIFFERENCE IN BOXES/HA	DIFFERENCE IN %
	CONTROL	TREATED.		
X	188	228	40	17.54%
XX	63	145	82	56.55%
XXX	47	95	48	50.52%
BRUCE	29	30	1	3.33%

5

Example 11

A trial with HYTa and HYTb was conducted in Norway on a potato crop. Tests with and without HYTa and HYTb were treated with 50 or 100 kg Nitrogen fertilizer/ha. Pesticides were used in normal amounts.

10 At the time of first emergence (June 14), 0.2 liers of HYTa and 0.6 liters of HYTb was applied per decare. After the last application of dirt (July 20), 0.2 liter of HYTa, 0.2 liter of HYTb and 50 grams of HYTc were applied per decare. The results are shown in Figure 5

15 Use of HYTa and HYTb gave a yield increase of up to 17% as compared to the control. In addition, there was less potato blight on the HYTa and HYTb treated crop as compared to control.

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The reference in this specification to any prior publication (or information derived from it), or to any matter which is known, is not, and should not be taken as an acknowledgment or admission or any form of suggestion that that
5 prior publication (or information derived from it) or known matter forms part of the common general knowledge in the field of endeavour to which this specification relates.

Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and
10 variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integers or steps.

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THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A composition comprising HYTa and at least one of chitin, chitosan, glucosamine and amino acids.

5

2. The composition of claim 1 comprising:

HYTa and chitin;

HYTa and chitosan;

HYTa and glucosamine;

10

HYTa and amino acids;

HYTa, chitin, and amino acids;

HYTa, chitosan, and amino acids;

HYTa, chitin, chitosan, and amino acids;

HYTa, chitosan, glucosamine and amino acids; or

15

HYTa, chitin, chitosan, glucosamine, and amino acids.

3. The composition of claim 1, comprising HYTa and at least two of chitin, chitosan, glucosamine and amino acids.

20 4. The composition of claim 1, comprising HYTa and at least three of chitin, chitosan, glucosamine and amino acids.

5. The composition of any of claims 1 through 4, where said chitin is from HYTc.

25

6. The composition of any of claims 1 through 5, where said chitosan, glucosamine and amino acids are from HYTb.

7. A composition comprising HYTa and at least one of HYTb and
30 HYTc.

8. A composition comprising HYTa, HYTb and HYTc.

9. A process comprising contacting soil, seed, seedling or plant foliage with HYTa to form treated soil, seed, seedling, or plant foliage.

5

10. The process of claim 9, further comprising contacting soil, seed, seedling or plant foliage with at least one of chitin, chitosan, glucosamine and amino acids.

10 11. The process of claim 10, further comprising contacting soil, seed, seedling or plant foliage with two or more of chitin, chitosan, glucosamine and amino acids.

12. The process of claim 10, further comprising contacting
15 soil, seed, seedling or plant foliage with three or more of chitin, chitosan, glucosamine and amino acids.

13. The process of claim 10, further comprising contacting soil, seed, seedling or plant foliage with chitin, chitosan,
20 glucosamine and amino acids.

14. The process of any of claims 9 through 13, where said chitin is from HYTc.

25 15. The process of any of claims 9 through 14, where said chitosan, glucosamine and amino acids are from HYTb.

16. The process of claims 9 through 15, wherein said HYTa is activated in an aqueous solution for 24-168 hours before said
30 contacting.

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17. The process of any one of claims 9 through 16, wherein said method further comprises contacting said treated soil or foliage in said treated soil with one or more of HYTa, chitin, chitosan, glucosamine and amino acids.

5

18. The process of any one of claims 9 through 16 wherein said method further comprises contacting said treated foliage or soil containing a plant with said treated foliage with one or more of HYTa, chitin, chitosan, glucosamine and amino acids.

10

19. The process of any one of claims 9 through 18, wherein said plants, seedlings or seeds are present in said soil prior to said contacting step.

15

20. The process of any one of claims 9 through 18, wherein said treated plants, seedlings or seeds are transplanted to said treated soil.

20

21. Activated HYTa made by incubating HYTa in the presence of HYTc for 24-168 hours.

22. A process comprising combining activated HYTa and at least one of HYTb and HYTc to form a mixture.

25

23. The process of claim 22, further comprising applying said mixture to soil, foliage, seed, or seedlings.

24. A composition consisting essentially of HYTa and at least one of HYTb or HYTc.

30

25. A composition consisting essentially of HYTa, HYTb and HYTc.

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26. A composition comprising soil, a plant, a seed, or a seedling treated with HYTa.

5

Alluvial	Area 9	TRT 3	Area 10	TRT 2	Area 11	TRT 4	Area 12	TRT 1	Area 13	TRT 1	Area 14	TRT 4	Area 15	TRT 2	Area 16	TRT 3
	Area 4	TRT 3			TRT 3			TRT 1			TRT 1			TRT 1		
	Area 3	TRT 4			TRT 4			TRT 2			TRT 2			TRT 2		
	Area 2	TRT 2			TRT 2			TRT 3			TRT 4			TRT 4		
Clay	Area 1	TRT 1			TRT 1			TRT 3			TRT 3			TRT 3		
	Area 3	TRT 4			TRT 4			TRT 2			TRT 2			TRT 2		
	Area 2	TRT 2			TRT 2			TRT 3			TRT 4			TRT 4		
	Area 1	TRT 1			TRT 1			TRT 3			TRT 3			TRT 3		

FIG. 1

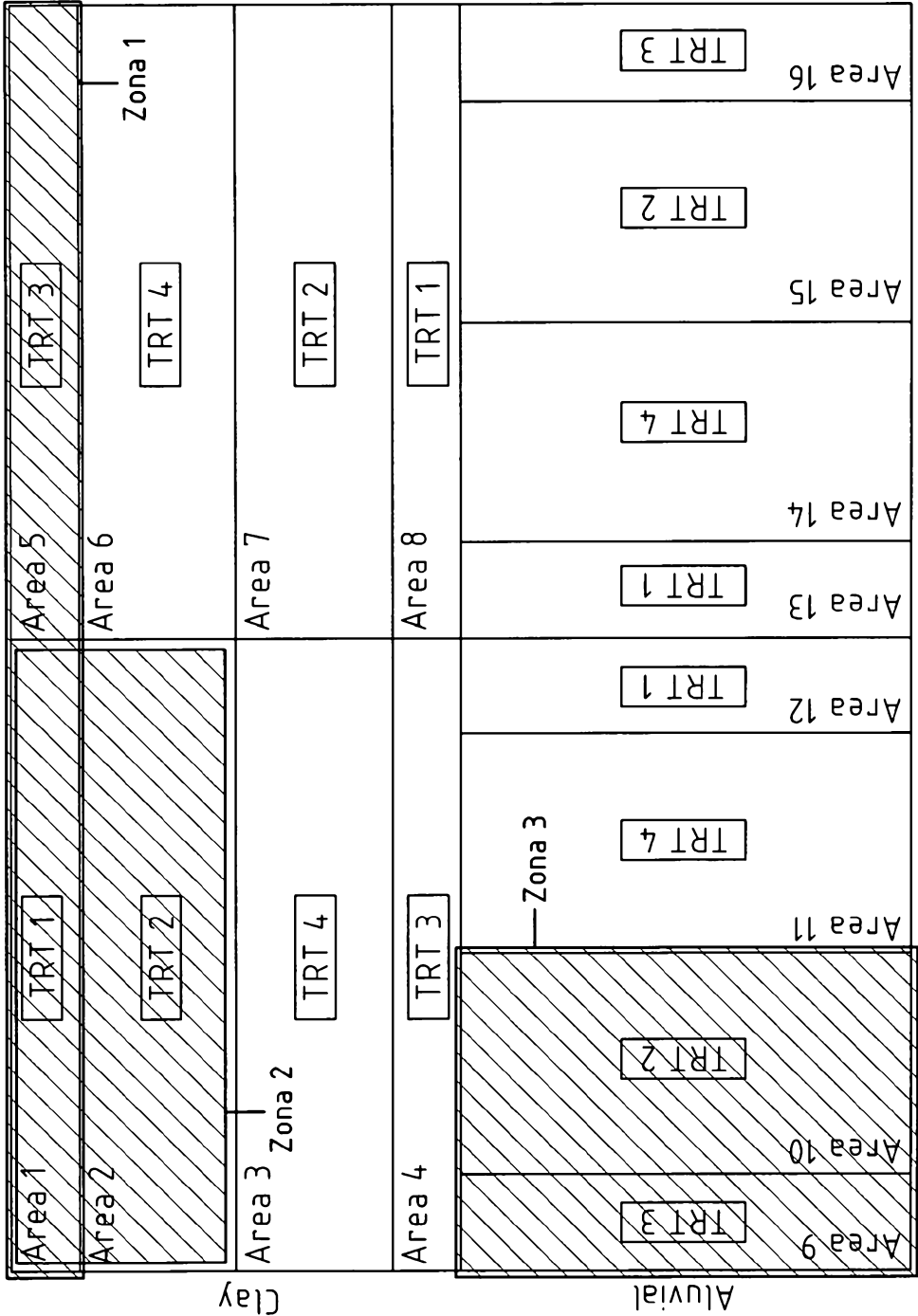


FIG. 2

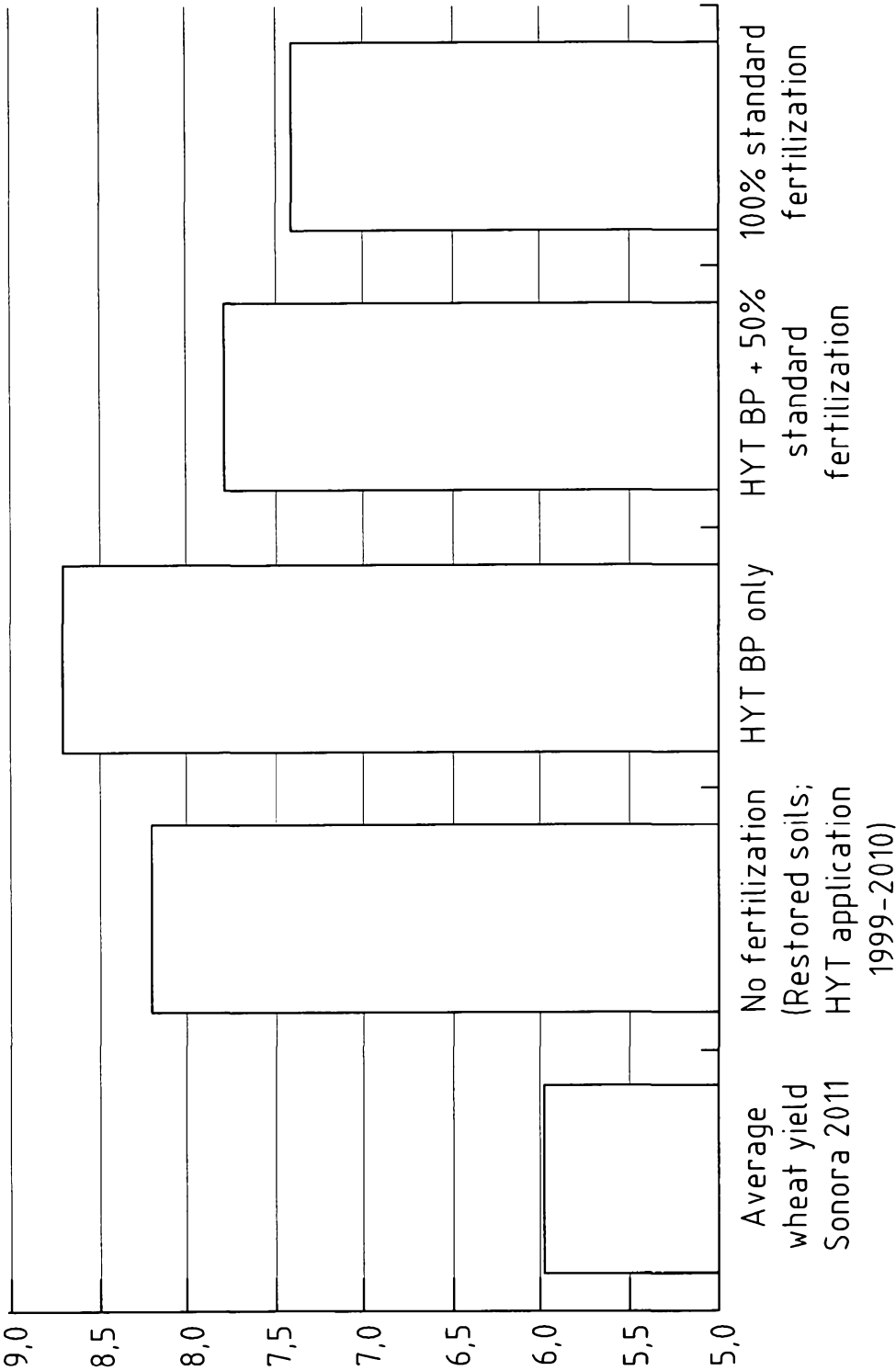


FIG. 3

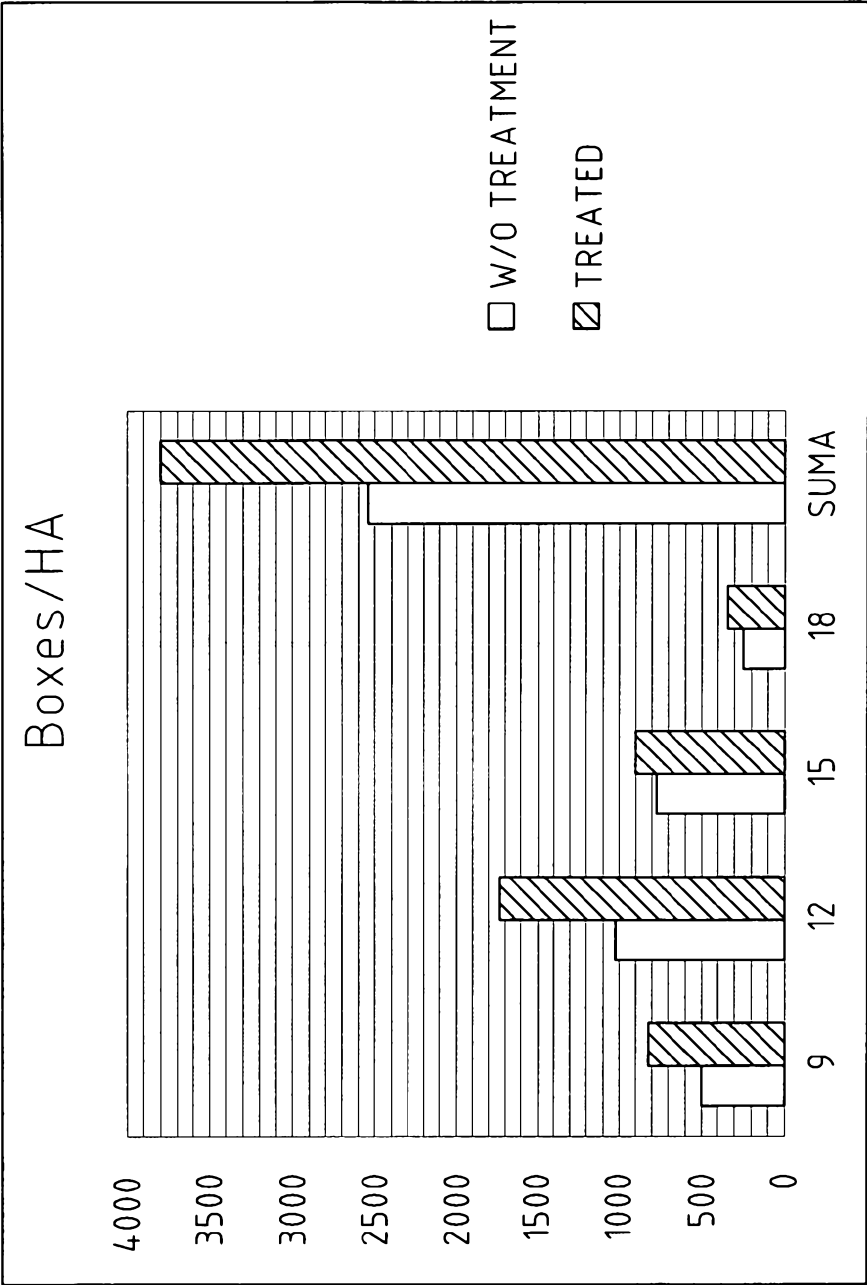


FIG. 4

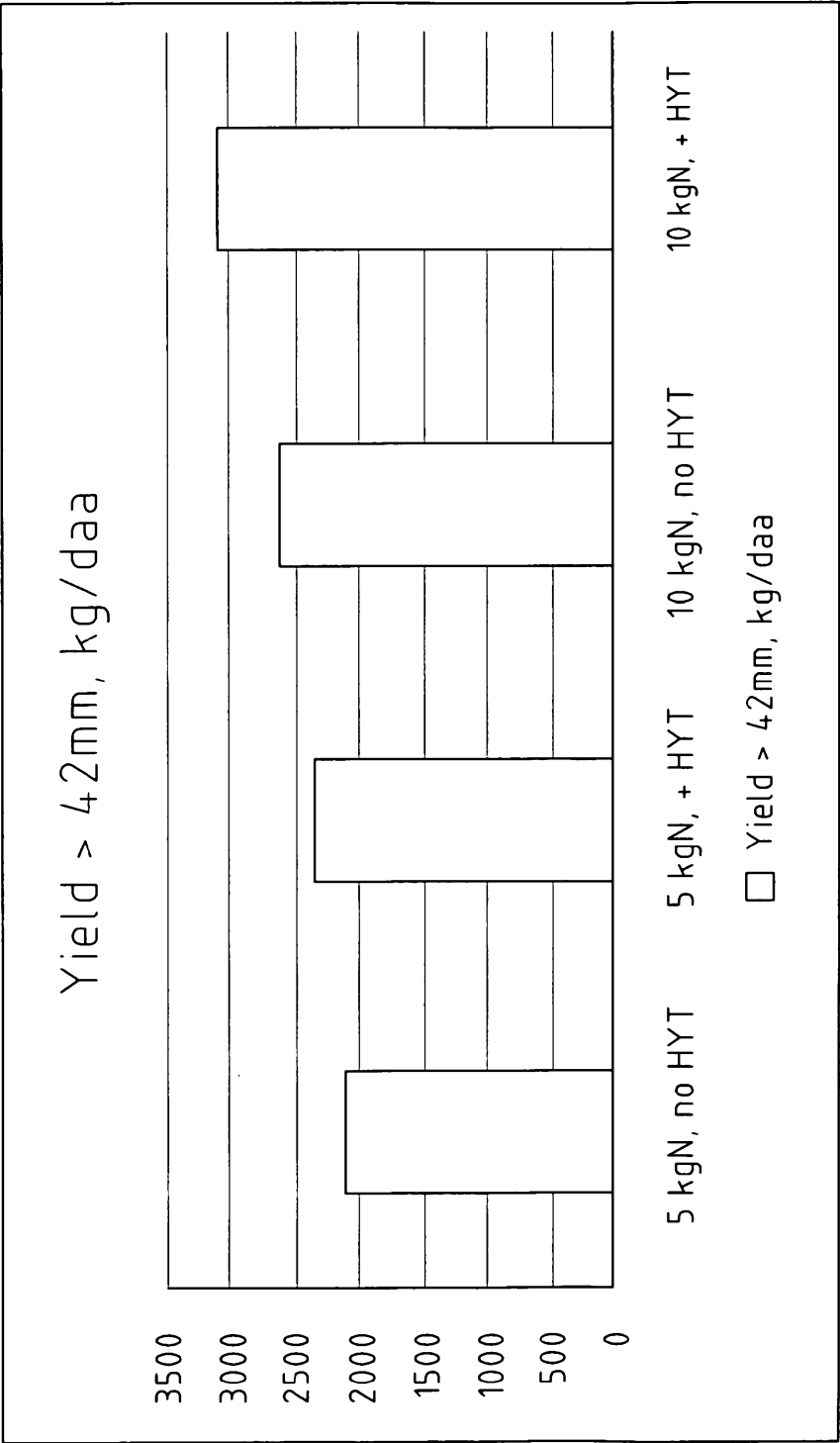


FIG. 5