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ning of each regular issue of the PCT Gazette.*

(54) Title: A RELEASE COMPOSITION AND METHOD OF PREPARATION

(57) Abstract: A method of producing a thermo-stable biodegradable thermo-stable biomatrix for storage of biological materials in disclosed. The biomatrix contains a bio-polymer selected from the group of xanthan gum, acacia gum, guar gum, gellan, starch or a combination thereof. The biological material can be a wide range of materials including; a bio-inoculant such as Rhizobium, or any microorganism or cellular organism. The biomatrix can be formulated for fast or slow release of the biological materials and maintains activity of the biological materials after being stored for months. In use the biomatrix can be applied in the form of a pellet or mixed with liquid and applied to seeds, plants or soil.



WO 02/15703 A1

TITLE: A RELEASE COMPOSITION AND METHOD OF PREPARATION**TECHNICAL FIELD**

The present invention relates to a release composition and method of preparation. More
5 specifically the present invention relates to a composition preferably of a type that allows the
release of a compound of biological materials contained within the composition once the
composition is placed in water or other solvent.

BACKGROUND ART

10 For the purpose of this specification the term "biological material" is used to encompass, but
is not limited to, any or all of the following: a bio-inoculant, a micro-organism, biological
cells, a part or parts of biological cells, pharmaceuticals, enzymes, hormones, proteins and
other bio-chemicals, unstable compounds and compositions (both biological and non-
biological); and a combination of these.

15

Two known problems are associated with the industrial or agricultural application of
biological materials. Firstly, there is the difficulty of maintaining the biological materials in
a viable state until they are used or required, and especially during the period in which they
may be incorporated in a release mechanism for delivery of the biological material.

20

Secondly, there may be a need for further storage before the release of the biological
material, once it has been applied or distributed to the intended substrate. The delay of
release can be for a number of reasons. For example: the composition may need to be stored
after manufacture so as to be ready when needed (convenience factor); there may need to be
25 a time delay to the point at which release of the biological material starts, so that the material
is released at the intended time for the intended purpose.

In this respect, an example is the release of a hormone for spring growth, intended for
consumption by young steers or calves when they start grazing. The delay can also be
30 intended to engender a weedicide, herbicide or insecticide with long-term action after the
immediate release into the substrate.

For the purposes of this specification the term "substrate" is used to encompass, but is not limited to, agricultural, horticultural, forestry or other commercial substrates, such as grasses and crops, soils (etc); water, waste water, skins of animals and tissues of animals; and solids
5 such as sands and gravels.

There have been numerous solutions disclosed for encapsulating products for fast release of compounds to substrates.

10 An example of such a granule can be found in US patent No. 6087306. This patent discloses a granule, suitable for aqueous spray application, after mixing of the granules with water in a tank, and application through a standard sprayer. The granule incorporates a wettable, dispersible, or water-soluble granule, with an agrochemical or other ingredient suitable for application by aqueous spray. The non-ionic, or predominately non-ionic, water-swella-
15 ble gums and polymers are disclosed as including microbial polysaccharides such as dextran, gellan gum, and xanthan gum, along with other polysaccharides. These gums and the active ingredient are combined along with an inert filler and extruded as a paste. This paste is dried and the result formed into granules in an extruder. The inert material is preferably water-swella-
20 ble. The active ingredient is disclosed as a herbicide, insecticide, nematocide, fungicide or plant growth regulator. A wetter or other adjuvant is optionally added.

US Patent No. 4859377 discloses the use of starch to encapsulate a release granulation. The starch is used substantially as an agent without cross-linking. This also discloses, however, that the starch needs to be treated in order to ensure that it is insect digestible, such that the
25 biological agent being carried within the starch capsule reaches the desired destination.

US Patent 4563344 discloses elements for both quick and slow release pesticides. The biological chemical is enclosed in a capsule incorporating peat, fine and coarse chaff and the woody portions of corncobs. The combinations and proportions are combined in a pre-
30 determined manner which affect the rate of release.

US Patent 4434231 discloses preparation of pellets which incorporate a bio-matrix gel, by the addition of crushed silica which is then homogenised. The resulting pellets are 3 mm in diameter, 33 mm in length and cylindrical in shape. However, there is no disclosure of whether the resultant pellets are fast acting or slow acting or how they breakdown or are
5 activated to break down.

However, all the above disclosed methods of conveying a biological agent to the soil or to a substrate (as herein before defined) disclose disadvantages: the matrix incorporating a biological material needs to be especially treated (as in US 4859377) in order to be adapted
10 to break down at the desired end point and time (for example, inside an insect). Certain types of encapsulating material will only incorporate chemicals, not all types of biological materials (US 4563344). Some of the compositions are disclosed as being most appropriately applied in an aqueous spray for chemical insecticides like *Lambda-cyhalodhrin* and *Pirimicarb* and thus may not be useful for biological materials. (US 6087306). Thus
15 many more steps are required to get the biological material to the intended substrate. This in turn increases the inconvenience and cost of application of the biological material.

It is an object of the present invention to provide a composition and method of preparation, which method is both quick and simple, to provide a thermo-stable bio-degradable medium
20 for transfer of a biological material to a substrate, at which point it breaks down quickly on contact with water.

It is a further object of the present invention to address the foregoing problems or at least to provide the public with a useful choice.
25

Further aspects and advantages of the present invention will become apparent from the ensuing description which is given by way of example only.
30

DISCLOSURE OF INVENTION

According to one aspect of the present invention there is provided a method of preparation of a release composition, which method comprises the steps of:

- (a) preparing a bio-degradable bio-matrix either as a gel or a liquid, incorporating a biological material, wherein said liquid is of high or medium viscosity;
- 5 (b) preparing a dry powder of one or more inert compounds; and
- (c) mixing the preparation of step (a) and the preparation of step (b) to form a homogenous mix;

and wherein the bio-matrix is selected from the group: xanthan gum; acacia gum; guar gum; gellan; starch; and a combination thereof;

- 10 and wherein the biological material is selected from: a bio-inoculant, a micro-organism, biological cells, a part or parts of a biological cell, a vaccine, at least one pharmaceutical compound, at least one enzyme, at least one hormone, at least one protein; at least one bio-chemical, biological unstable composition; at least one non-biological compound; and a combination of these;

- 15 and wherein said composition is thermo-stable and bio-stable in the absence of substantial water.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above, wherein said inert compounds
20 are chemically unreactive with respect to the remaining compounds or elements used in the method.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above, wherein said inert compounds
25 are selected from the group: inorganic compounds; powdered sterilised soils; powdered sterilised substrates; powdered silica; powdered dry organic materials; and a combination thereof.

Advantageously, said organic material may be selected from agricultural materials or waste
30 agricultural materials, for example: corn cobs; chaff; straw. Advantageously, said inert compounds may include peat and sands. Advantageously, the inorganic compound is a clay which is selected from talc, bentonite and diatomaceous earth, and a combination thereof.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above wherein said method includes a further step after step (c) as follows:

- 5 (d) mixing the composition formed in step (c) with at least one clay in powdered form.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above wherein the inert compound
10 used in step (b) and step (d), if present, is clay which is selected from: diatomaceous earth, talc, bentonite, a combination thereof.

According to a further aspect of the present invention there is provided the process for the preparation of a release composition as described above wherein the clay in step (d) (if
15 present) is powdered bentonite.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above, wherein said method includes a further step, after step (c) as follows:

- 20 (e) repeating steps (a) to (c) one or more times with a different bio-matrix and/or a different biological material; and mixing the homogenous mix resultant from each step (c).

According to another further aspect of the present invention there is provided a process for
25 the preparation of a fast release composition as described above, wherein said method includes step (d) after step (e).

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above, wherein said method includes a further step after step (d) as follows:

- 30 (f) mixing the resultant mix of step (d) with water, to form a dough.

According to another further aspect of the present invention there is provided a process for the preparation of a release composition as described above, wherein said method includes further steps after step (f) as follows:

- 5 (g) forcing through or extruding the dough of step (f) in a pelletiser or granulator to form pellets;
- (h) drying the pellets or granules at room temperature.

The pellets or granules may have a size of 0.1 – 20mm diameter. The pellets may be up to 30mm long.

10

Advantageously, the pellets resulting from step (g) may be in the range 0.1 mm to 10 mm in diameter, and up to 10 mm in length.

15 According to a further aspect of the present invention there is provided the method of preparation of a slow release or delayed release composition substantially as described above wherein said bio-degradable bio-matrix has between 15 to 40% water by weight at the end of step (g).

20 It is to be noted that no additional drying of the composition of step (c) is required where a liquid bio-matrix is used.

In the case of a fast release mixture, the granules and/or pellets formed are bio-stable, in that until they come into contact with substantial amounts of water, they are both bio-stable and thermo-stable. However, once water is introduced the structure of the granules/pellets
25 breaks down rapidly, within minutes, to release the biological material. This rapid breakdown optimally occurs in less than one minute.

In the case of fast release compositions, it has been found that the pellets of step (g) break down faster than composition formed in step (c) of the above process.

30

According to a further aspect of the present invention there is provided a method of preparing a slow release or delayed release composition substantially as described above

wherein said composition is a thermo-stable gel and the composition is capable of storage of the biological materials for a period of time.

5 The liquid bio-degradable bio-matrix can be obtained in a number of ways or arrived at through a number of methods of preparation. An example of this is the use of acacia gum mixed with distilled water and agitated; and the further addition of concentrated biological material. This mix can be left to stand for up to three hours before being mixed with the preparation of step (b).

10 Other alternative liquid bio-matrixes which are also bio-degradable and/or thermo-stable can be used. For example the liquid bio-matrix disclosed in specification of US Patent No. 5292507.

15 According to a further object of the present invention there is provided a method of producing a method of preparation of a release composition as described above wherein the biological material is cellular or a micro-organism. The concentration of such biological material at the end of step (g) is hereafter referred to as the "cell concentration". Preferably, the cell concentration is in the range 10^5 cells to 10^{12} cells g^{-1} , more preferably in the range 10^8 to 10^{12} cells g^{-1} , more preferably in the range of 10^9 to 10^{10} cell g^{-1} . Preferably, the
20 biological material may be present in step (a) in a broth, or on a growing medium.

According to another aspect of the present invention there is provided a method of preparing a slow-release or delayed release composition substantially as described above wherein the biological material includes: a pesticide; a viricide; a bacteriacide, a fungicide; and a
25 combination thereof.

According to a further aspect of the present invention there is provided a method for producing a thermo-stable biodegradable medium for storage of biological materials wherein the biological material is a vaccine selected from: a live vaccine; an oral attenuated vaccine;
30 an encapsulated mycobacterium vaccine; and a combination thereof. Examples of the vaccine include Bacille Calmette and Guerin (B.C.G.).

For the purposes of this specification, the term "storage" means a stability of better than LT_{50} with respect to the cell concentration of the biological material. That is, more than 50% of the cells (if cells are the biological material) are viable at the end of the storage period; or more than 50% of the non-living material is viable at the end of the storage period.
5 Advantageously, LT_{50} may be achievable after two months, four to six months, or 12 to 18 months.

For the purposes of this specification, the term "thermo-stable" means a range of temperatures in which the combination of bio-polymer and biological material is stable.
10 This temperature range is 4°C to 40°C, and preferably between 5°C to 30°C.

According to another aspect of the present invention there is provided a method of preparing a slow-release or delayed release composition substantially as described above wherein the biological material is a micro-organism. Said micro-organism may be selected from the
15 group *Serratia*, *Pseudomonas*, *Xanthomonas*, *Rhizobium*. Most preferably the micro-organism is *Serratia entomophila*.

According to another aspect of the present invention there is provided a slow-release or delayed release composition incorporating a biodegradable thermo-stable bio-matrix with a
20 biological material therein, said composition being formed by the above described method.

According to another aspect of the present invention there is provided a fast release composition, suitable for application to a substrate (as defined above), the composition being produced by steps (a) to (c), or steps (a) to (e) of the method as described above.
25

According to a further aspect of the present invention there is provided a method of producing a fast release composition as described above wherein the steps (f) and (g) occur a substantial time after the immediately preceding step. Optimally, the steps (f) and (g) can occur immediately before the composition is applied to a substrate.
30

According to a further aspect of the present invention there is provided a composition, in granular form, produced in accordance with steps (a) to (c) or steps (a) to (e) of the method

of preparation as described above, characterised in that to the composition can be added one or more clays or/and other dry powders or/and sufficient water, and that the composition is capable of being formed into pellets of the type described above.

- 5 According to a further aspect of the present invention there is provided a release composition, produced in accordance steps (a) to (d) of the method of preparation as described above.

10 According to another aspect of the present invention there is provided a slow-release composition in the form of a dough incorporating a biodegradable thermo-stable bio-matrix and biological material, produced by the steps (a) to (e) of the above described method, wherein said dough can be passed through a pelletiser and dried.

15 According to a further aspect of the present invention there is provided a bulk dough release composition produced in accordance with steps (a) to (f) of the method of preparation as described above, and wherein said dough is processed through a pelletiser to form pellets.

20 According to a further aspect of the present invention there is provided pellets produced in accordance with steps (a) to (g) of the method described above.

Preferably, the pellets are of a size that can easily be drilled into soil or other substrate, either alone or in combination with seeds.

25 According to another aspect of the present invention there is provided a slow-release composition in the form of a powder, as prepared by steps (a) to (c) of the above described method, wherein said powder, when mixed with a least one clay and sufficient water, forms a dough capable of passage through a pelletiser.

30 According to another aspect of the present invention there is provided a slow-release composition in the form of a powder, as prepared by steps (a) to (d) of the above described method, wherein said powder, when mixed with sufficient water, forms a dough capable of passage through a pelletiser.

Optionally, the bio-matrix of step (a) will be between 0-50% water by weight. It will be appreciated that the powdered mix after step (c) of the method, or the dough formed after step (e) of the method, is also suitable for storage of the biological material until it is
5 required to be applied to the substrate.

Optionally, the release composition can be used to stabilize biological material for transportation and/or storage and/or delivery.

10 According to a further aspect of the present invention there is provided the use of the release component for delivery of biological material via a spray for application to plants and/or animals, characterised in that the biological material includes an active ingredient to be sprayed over plants and/or animals.

15 According to another aspect of the present invention there is provided a spray solution which includes at least one release composition.

According to a still further aspect of the present invention there is provided a method of inoculating a plant seed with a biological material, said method including the steps of;

- 20 (a) selecting at least one biological material to be used as an inoculant;
(b) preparing the composition by the above described method;
(c) adding the composition to water and mixing to release the biological material into the solution;
(d) soaking the plant seed in said solution to allow the biological material to coat the
25 plant seed.

According to a further aspect of the present invention there is provided a method of inoculating a plant seed with a biological material substantially as describe above, said method including, a further step after step (b), of (bi): adding a powdered compound to the
30 composition, said powdered compound being selected from the group: a second biological material, a dried and powdered composition, a dried and powdered bio-polymer matrix containing a second or a third biological material, a chemical, and a combination of these.

Preferably, the plant seed can be dried at room temperature before drilling or seed broadcast. Preferably, more than one inoculant may be used in step (a) above, each inoculant being for a different purpose. As the bio-matrix is thermo-stable and bio-stable, the seeds need not be
5 drilled or sown immediately after the inoculation process.

According to a further aspect of the present invention there is provided seed inoculated by the method as described above.

- 10 According to a further aspect of the present invention there is provided a composition for application to a substrate (as hereinbefore defined) wherein said composition includes:
- (a) one or more fast release compositions (substantially as hereinbefore described);
 - and
 - (b) one or more slow release compositions.

15

BEST MODES FOR CARRYING OUT THE INVENTION

Example 1

A slow release gel is made with xanthan gum as the bio-polymer and *Serratia entomophila*
20 as the biological material. The cell concentrations are set out in Table 1.

To 5 grams of dry xanthan gum is added 5 grams monounsaturated oil. The mixture is agitated at room temperature for between 5-10 minutes to form a suspension.

25 90 grams of the micro-organism concentrate is added to the suspension. The mix is agitated for a further 10 minutes at room temperature. The result is a gel matrix.

Equal portions of diatomaceous earth and talc are mixed to form 200 grams of powder. To this powder is added the 100 grams of gel. This can be done, for example, by drying and
30 crumbling the blend of the gel and diatomaceous earth and talc, or by other known means.

To 300 grams of this mixed powder is added 27 grams of bentonite and up to 180 grams of distilled water. This mixture is homogenised and forms a dough. The mixture is passed through a pelletiser (of known type) and/or a die to form pellets of predetermined size and thickness. These pellets are then air dried to between 10-40% moisture content.

5

The results of six tests using the method of example 1 are shown in table 1 below. A comparison is shown of the survival of *Serratia entomophila* in broth at 20°C.

TABLE 1

10

Example 1

Sample #	Initial concentration cfu g ⁻¹	Survival – 1 month cfu g ⁻¹	Survival - 2 months cfu g ⁻¹	Survival - 4 months cfu g ⁻¹	Survival - 6 months cfu g ⁻¹	LT ₅₀ days
60	6.08x10 ⁷	2.72x10 ⁹	1.64x10 ⁹	5.02x10 ⁸	1.52x10 ⁸	60-120
61	1.32x10 ⁹	6.98x10 ⁹	3.05x10 ⁹	1.23x10 ⁹	6.81x10 ⁸	120-180
62	2.77x10 ⁹	6.94x10 ⁹	7.53x10 ⁹	4.42x10 ⁶	6.68x10 ⁶	60-120
63	3.53x10 ⁹	4.59x10 ⁹	1.24x10 ⁹	4.98x10 ⁹	3.87x10 ⁹	>180
64	8.25x10 ⁸	3.02x10 ⁹	1.75x10 ⁹	3.78x10 ⁹	1.78x10 ⁹	>180
213	4.65 x 10 ⁹	-	-	3.72x10 ⁹	2.04x10 ⁹ *	~165

Comparison

	Initial cfu g ⁻¹	1 week cfu g ⁻¹	2 week cfu g ⁻¹	3 week cfu g ⁻¹	LT ₅₀ Days
Broth at 20°C	6.67x10 ¹⁰	4.52x10 ¹⁰	1.22x10 ¹⁰	3.83x10 ⁸	<14

15 **Example 2**

A slow release gel is made with xanthan gum as the bio-polymer and *Serratia entomophila* as the biological material. The cell concentrations are set out in Table 2.

To 7.5 grams of dry xanthan gum is added 42.5 grams distilled water. The mixture is agitated at room temperature for between 5-10 minutes to form a suspension. Alternatively, a 50% solution of xanthan gum medium may be used for the gel.

- 5 50 grams of the micro-organism concentrate is added to the suspension. The mix is agitated for a further 10 minutes at room temperature. The result is a gel matrix.

Equal portions of diatomaceous earth and talc mixed to form 100 grams of powder. To this powder is added 100 grams of gel. This can be done, for example, by drying and crumbling
10 the blend of the gel and diatomaceous earth and talc, or by other known means.

To 200 grams of this mixed powder is added 20 grams of bentonite and between 75-95 grams of distilled water. This mixture is homogenised and forms a dough. The mixture is passed through a pelletiser (of known type) and/or a die to form pellets of predetermined size
15 and thickness. These pellets are then air dried to between 10-40% moisture content.

The pellets, depending on the die, may vary in size from 0.1-20 mm in diameter and from 0.1-20 mm in length. Preferably, if the pellets are to be drilled, a diameter of less than 3 mm is used. The pellets are stored in the absence of moisture at room temperature, until
20 required.

Data for the survival of micro-organisms in a slow release pellet form over a period of months from 1-6 is shown in Table 2.

25 Alternatively, the powdered mix before the addition of the bentonite, or the dough after the addition of water, may be stored in the absence of moisture. The additional steps described above can be performed in a one or two stage process, depending on when the pellets or the composition is needed. Thus the next step can be separated in time from the immediately preceding mixing step and also from the step of addition of water. Alternatively, these steps
30 can be performed immediately one after the other, and immediately before the pellets are to be used.

The pellets, if small, can be mixed with seed and passed through a drill in known manner and directly drilled into the soil. Alternatively, granules or pellets from the composition may be broadcast in known manner to a substrate of plants or soils or a combination thereof.

5

TABLE 2

Example 2

Sample #	Initial Concentration cfu g ⁻¹	Survival – 1 month cfu g ⁻¹	Survival - 2 months cfu g ⁻¹	Survival - 4 months cfu g ⁻¹	Survival - 6 months cfu g ⁻¹	LT ₅₀ days
14	1.46x10 ¹⁰	2.63x10 ⁹	4.76x10 ⁹	2.76x10 ⁸	7.90x10 ⁵	60-120

Example 3

A slow release gel is made with guar gum as the bio-polymer and *Serratia entomophila* as the biological material. The cell concentrations are set out in Table 3.

To 5.6 grams of dry guar gum is added 0.5 grams of 0.5M sodium hydroxide. The mixture is agitated at room temperature for between 5-10 minutes to form a suspension.

100 grams of the micro-organism concentrate is added to the suspension. The mix is agitated for a further 10 minutes at room temperature. The result is a gel matrix.

Equal portions of diatomaceous earth and talc are mixed to form 150 grams of powder. To this powder is added 106.1 grams of gel. This can be done, for example, by drying and crumbling the blend of the gel and diatomaceous earth and talc, or by other known means.

To 256.1 grams of this mixed powder is added 18g of bentonite and up to 70 grams of distilled water. This mixture is homogenised and forms a dough. The mixture is passed through a pelletiser (of known type) and/or a die to form pellets of predetermined size and thickness. These pellets are then air dried to between 10-40% moisture content.

A second sample is made by comparison without sodium hydroxide. The same method as above is used, however the quantities used are: 2.1 grams of dry guar gum added to 60 grams

of micro-organism concentrate. The mix is agitated for 10 minutes at room temperature. The result is a gel matrix. A further 110-grams of diatomaceous earth-talc powder (mixed in equal proportions) is added to the gel matrix. After this, a further 10 grams of bentonite and 135 grams of distilled water is added. This mixture is formed into a dough and pelletised as above.

The results for both treatments are shown in Table 3 below.

TABLE 3

Example 3

Sample #	Initial Concentration cfu g ⁻¹	Survival – 1 month cfu g ⁻¹	Survival – 2 months cfu g ⁻¹	Survival – 4 months cfu g ⁻¹	LT ₅₀ days
148	6.25 x 10 ⁹	8.83 x 10 ⁹	5.56 x 10 ⁹	1.45 x 10 ⁹	60-120
149	5.31 x 10 ⁹	8.58 x 10 ⁹	5.51 x 10 ⁹	2.64 x 10 ⁷	60-120

Example 4

The same composition and method was used as for Example 1, however the microbe *Pseudomonas fluorescence* was used in the microbial concentrate.

Results were taken of the cell concentration initially and after two months. The results are shown below in Table 4.

TABLE 4

Example 4

Sample #	Initial Concentration cfu g ⁻¹	Survival – 2 month cfu g ⁻¹	LT ₅₀ days
288	4.18 x 10 ⁹	1.26 x 10 ¹⁰	>60

Example 5

A fast release composition is made by combining 20 grams of acacia gum with 30 grams of distilled water. The mix is agitated at room temperature. To this mix is added 50 grams of concentrated biological material containing *Bacillus mycoides*. This mixture is agitated at room temperature and left to stand for between 1½ to 3 hours at room temperature.

A powder is formed of diatomaceous earth and talc in the ratio 50:50 (by weight). The solution of acacia gum and biological material is added to 200 grams of the powdered mix and mixed thoroughly.

A second mixture was prepared using the above method. However, the quantities used were 16 grams of acacia gum, 8 grams of 0.5% yeast, 12 grams of distilled water, 36 grams of *Trichoderma* spores and 130 grams of a 50:50 mix of diatomaceous earth and talc (by weight).

Either mixture can either be stored at room temperature until required.

The survival rates for the biological materials of Example 5 were tested and the results are set out in Table 5 below.

TABLE 5**Example 5**

Sample #	Organism	Initial cfu g ⁻¹	3 months cfu g ⁻¹	6 months cfu g ⁻¹	8 months cfu g ⁻¹	LT ₅₀ days
186	<i>Bacillus Mycoides</i>	8.47x10 ⁵	8.25x10 ⁵	8.00x10 ⁵	4.3x10 ⁵	>240
171	<i>Trichoderma</i>	2.96x10 ⁹	2.02x10 ⁹	1.81x10 ⁹	7.81x10 ⁸	120-180

Example 6

A fast release composition was made by combining 2.5 grams of gellan gum with 50 grams of concentrated biological material containing *Bacillus mycoides*. This mixture is agitated at room temperature and left to stand for between 1½ to 3 hours at room temperature.

A powder is formed of diatomaceous earth and talc in the ratio 60:40 (by weight). The solution of gellan gum and biological material is added to 100 grams of the powdered mix and mixed thoroughly. To this resulting mixture a further 15 grams of bentonite and 100
5 grams of distilled water is added.

A second mixture was prepared using the same composition and method as used above, however the microbe *Pseudomonas fluorescense* was used in the microbial concentrate.

10 A third mixture was prepared using the above method, however the quantities used were 3 grams of gellan gum, 9 grams of 0.5% yeast, 18-grams of distilled water, 28 grams of *Trichoderma* spores and 120 grams of a 50:50 mix of diatomaceous earth and talc (by weight), 15 grams of bentonite and 110 grams of distilled water.

15 Results were taken of the cell concentration initially and after various monthly intervals thereafter. The results are shown below in Table 6.

TABLE 6

Example 6

Sample #	Organism	Initial cfu g ⁻¹	2 months cfu g ⁻¹	3 months cfu g ⁻¹	6 months cfu g ⁻¹	8 months cfu g ⁻¹	LT ₅₀ days
185	<i>Bacillus mycoides</i>	1.61x10 ⁶	-	1.40x10 ⁶	1.02x10 ⁶	8.03x10 ⁵	>240
289	<i>Pseudomonas fluorescense</i>	1.73x10 ⁹	1.86x10 ⁹	-	-	-	>60
172	<i>Trichoderma</i>	1.2x10 ⁷	-	1.3x10 ⁷	4.6x10 ⁶	6.7x10 ⁵	90-180

20

Example 7

A fast release composition is made by combining 3.5 grams of starch with 50 grams of hot water (70°C to 80°C). The mix is held in this state for 10 minutes and then cooled to room
25 temperature. To this mix is added 50 grams of concentrated biological material (in this

example *Serratia entomophila*). This mixture is agitated at room temperature and left to stand for between 1 ½ to 3 hours at room temperature.

A powder is formed of diatomaceous earth and talc in the ratio 60:40 (by weight). The solution of starch and biological material is added to 175 grams of the powdered mix, 26 grams of bentonite and 150 grams of distilled water, and the resulting solution is mixed thoroughly.

A similar fast release composition is made by combining 6 grams of starch with 100 grams of concentrated biological material (*Serratia entomophila*). A powder is formed of diatomaceous earth and talc in the ratio 60:40 (by weight). The solution of starch and biological material is added to 175 grams of the powdered mix, 12 grams of bentonite and 180 grams of distilled water and the resulting solution is mixed thoroughly.

The survival rates for both of the above options were tested at 2 ½ months, 3 ½ months and 5 ½ months. The results are set out in Table 7 below.

TABLE 7

Example 7

Sample #	Initial Concentration cfu g ⁻¹	Survival – 2½ months cfu g ⁻¹	Survival – 3½ months cfu g ⁻¹	Survival – 5½ months cfu g ⁻¹	LT ₅₀ days
208	1.79x10 ⁹	1.14x10 ⁹	2.8x10 ⁹	8.59x10 ⁸	>165
225	4.77x10 ⁹	5.26x10 ⁹	2.49x10 ⁹	1.25x10 ⁹	~165

Example 8

The success or otherwise of the biomatrix (biofungicide) in aiding root rot disease control was compared against a control sample where no treatment occurred and a seed treated with a known chemical fungicide.

The seeds chosen were pea seeds which are prone to *Aphanomyces* root rot. The microbe *Bacillus mycoides* is known to control this disease, as is the chemical fungicide used in the trial as a positive control. It was expected that the positive control fungicide would have a heightened effect as the fungicide has a broad range of fungicidal properties.

5

A matrix treated according to Example 6 was placed at the bottom of each drilled hole, upon which the pea seed was placed.

The results of the test are shown below in Table 8 where the better yield of the plant relates to the lower presence of *Aphanomyces* root rot.

10

TABLE 8

Example 8

Treatment	Initial Spore g ⁻¹	% Plot Stand	Pods per plant	Pod Weight per plant (g)	Yield of Pods per Plot (g)
Chemical fungicide	870	92.78	2.78	15.07	503.15
Control	4.0 x 10 ¹	77.22	2.59	14.37	399.62
Biofungicide	1.5 x 10 ⁶	79.44	3.26	16.86	482.20

It will be appreciated by those skilled in the art that a number of compounds, biological material or other chemicals can be added to the suspension prior to the inoculation of the plant seed. Such compounds can be for the promotion of plant growth, the promotion of growth of the animal feeding off the plant, etc.

It will also be appreciated that the suspension generated in the Example 8 above may be sprayed in known manner, to broadcast the biological material onto a substrate. Such spraying can be done either before or after planting.

It will also be further appreciated that the seeds in Example 8 above may be inoculated with the bio-matrix by soaking the seeds in the suspension prior to drilling.

25

Example 9

A further experiment to assess the success or otherwise of the bio-matrix in control of grass grubs at filed level was compared against a control sample where no treatment occurred and a field drilled with two separate release formulations containing *Serratia entomyphila*.

- 5 *Serratia entomyphila* is a microbe that is efficient at controlling grass grub, a common disease in pasture grass.

The above samples were compared to positive controls where a liquid, product was applied through a modified see drill onto the paddock.

10

The first release solution is that produced by example 2 and the second, by example 7.

- The paddock kept isolated via a fence. Presence of the disease in grass grubs was tested after 6 weeks in the case of the fast release formulation and after 10 weeks in the case of the slow release formulation. The results are shown below in Table 9. Ideally the higher the percentage of disease prevalence in the grass grub, the better the rate of success of the formulation i.e. the higher the occurrence of the disease in the grass grub relates to the presence of *Serratia*.

TABLE 9

20

Example 9

Treatment	Sample Set 1 Average	Sample Set 2 Average	Trial Average
Control	9.2%	2.7%	6.0%
Drilled Fast Release	25.2%	5.0%	15.1%
Drilled Slow Release	24.7%	18.4%	21.6%
Liquid (into SR)	23.5%	32.3%	27.9%
Liquid (into FR)	25.7%	9.1%	17.4%

Aspects of the present invention have been described by way of example only and it should be appreciated that modifications and additions may be made thereto without departing from the scope thereof.

THE CLAIMS DEFINING THE INVENTION ARE:

1. According to one aspect of the present invention there is provided a method of preparation of a release composition, which method comprises the steps of:
 - (a) preparing a bio-degradable bio-matrix either as a gel or a liquid, incorporating a biological material, wherein said liquid is of high or medium viscosity;
 - (b) preparing a dry powder of one or more inert compounds; and
 - (c) mixing the preparation of step (a) and the preparation of step (b) to form a homogenous mix;and wherein the bio-matrix is selected from the group: xanthan gum; acacia gum; guar gum; gellan; starch; and a combination thereof;
and wherein the biological material is selected from: a bio-inoculant, a micro-organism, biological cells, a part or parts of a biological cell, a vaccine, at least one pharmaceutical compound, at least one enzyme, at least one hormone, at least one protein; at least one bio-chemical, biological unstable composition; at least one non-biological compound; and a combination of these;
and wherein said composition is thermo-stable and bio-stable in the absence of substantial water.
2. A method of preparation of a release composition as claimed in claim 1 wherein said inert compounds are chemically unreactive with respect to the remaining compounds or elements used in the method.
3. A method of preparation of a release composition as claimed in either claim 1 or claim 2 wherein said inert compounds are selected from the group: inorganic compounds; powdered sterilised soils; powdered sterilised substrates (as hereinbefore defined); powdered silica; powdered dry organic materials; and a combination thereof.
4. A method as claimed in claim 3 wherein said organic material is selected from: corn cobs; chaff; straw; or a combination thereof.

5. A method as claimed in any one of claims 3 and 4 wherein said inert compounds is selected from: peat; sands; or a combination thereof.
6. A method as claimed in any one of claims 3 to 5 wherein the inorganic compound is a clay.
7. A method as claimed in any one of claims 3 to 5 wherein the inorganic compound is selected from: talc; bentonite; diatomaceous earth; and a combination thereof.
8. A method of preparation of a release composition as claimed in any one of the previous claims wherein said method includes a further step, after step (c), as follows:
 - (d) mixing the composition formed in step (c) with at least one further inert compound in powdered form.
9. A method of preparation of a release composition as claimed in any one of the previous claims wherein the inert compound used in step (b) and in step (d), if present, is selected from: diatomaceous earth; talc; bentonite; or a combination thereof.
10. A method of preparation of a release composition as claimed in claim 9 wherein the inert compound in step (d) (if present) is powdered bentonite.
11. A method of preparation of a release composition as claimed in any one of the previous claims wherein said method includes a further step, after step (c), and after step (d) if present, as follows:
 - (e) repeating steps (a) to (c) at least once more with a different bio-matrix and/or a different biological material; and mixing together the homogenous mix resultant from all steps (c).
12. A method of preparation of a release composition as claimed in any one of the previous claims wherein said method includes a second step (d) after step (e), and wherein the composition is a fast release composition.

13. A method of preparation of a release composition as claimed in claims 8 to 12 wherein said method includes a further step after step (c) and after step (d) if present, as follows:
(f) mixing the resultant mix of any step (d) with water, to form a dough.
14. A method of preparation of a release composition as claimed in claim 13, wherein said method includes further steps after step (f) as follows:
(g) forcing through or extruding the dough of step (f) to form pellets.
15. A method as claimed in claim 14 wherein the pellets resulting from step (h) have a size of 0.1 – 20 mm diameter and up to 30 mm long and are of a shape selected from round, ovoid, cylindrical, spherical and rectangular.
16. A method as claimed in claim 15 wherein the pellets resulting from step (g) are in the range 0.1 mm to 10 mm in diameter and up to 10 mm long.
17. A method of producing a release composition as claimed in any one of the previous claims wherein the biological material is cellular or is micro-organism.
18. A method as claimed in claims 14 to 17 wherein the concentration of biological material, at the end of step (g), is in the range 10^5 cells to 10^{12} cells g^{-1} .
19. A method as claimed in claim 18 wherein the cell concentration is the range 10^8 to 10^{12} cells g^{-1} .
20. A method as claimed in either claim 19 or claim 20 wherein the cell concentration is the range 10^9 to 10^{10} cells g^{-1} .
21. A method as claimed in any one of the previous claims wherein the biological material in step (a) is in a state selected from: a broth and on a growing medium.

22. A method of preparing a release composition as claimed in any one of the previous claims wherein the biological material is selected from: a pesticide; a viricide; a bacteriacide, a fungicide; and a combination thereof.
23. A method of preparation of a release composition as claimed in any one of the previous claims wherein the biological material is a vaccine selected from: a live vaccine; an oral attenuated vaccine; an encapsulated mycobacterium vaccine; and a combination thereof.
24. A method as claimed in claim 23 wherein the vaccine Bacille Calmette and Guerin (B.C.G.) is used.
25. A method as claimed in any one of the previous claims wherein the biological material is cellular or a micro-organism and storage of the composition is at a stability of better than LT_{50} with respect to the cell concentration for the length of time of storage.
26. A method as claimed in claim 25 wherein the temperature range for storage is 4°C to 40°C.
27. A method as claimed in either claim 25 or claim 26 wherein the temperature range for storage is 5°C to 30°C.
28. A method of preparation of a slow release composition as claimed in any one of claims 14 to 27, wherein said method includes further steps after step (g) as follows:
 - (h) drying the pellets or granules at room temperature.
29. A method of preparation of a slow release composition as claimed in claim 28 wherein said bio-degradable bio-matrix has between 15 to 40% water by weight at the end of step (g).
30. A method of preparing a slow-release composition as claimed in any one of the previous claims wherein the biological material is a micro-organism.

31. A method of preparing a slow-release composition as claimed in claim 30 wherein said micro-organism may be selected from the group *Serratia*, *Pseudomonas*, *Xanthomonas*, *Rhizobium*, and a combination thereof.
32. A method of preparing a slow-release composition as claimed in claim 30 wherein the micro-organism is *Serratia entomophila*.
33. A method of preparing a fast release composition as claimed in any one of the previous claims wherein the pellets formed are both bio-stable and thermo-stable and wherein when introduced to water, in a time of minutes, the physical structure of the granules or pellets disintegrates to release the biological material.
34. A method of preparing a fast release composition as claimed in claim 17 wherein the time is less than one minute.
35. A method of producing a fast release composition as claimed in either claims 13 or 14 wherein the steps (f) and (g) occur a substantial time after the immediately preceding step.
36. A method of producing a fast release composition as claimed in claim 35 wherein, the steps (f) and (g) occur immediately before the composition is applied to a substrate.
37. A release composition formed by the method of any one of the previous claims.
38. A release composition produced by the method of any one of claims 1 to 36 where the composition is used to stabilize biological material for any one of: transportation; storage; delivery and a combination thereof.
39. A slow release composition formed by any one of claims 1 to 32.
40. A fast release composition formed by any one of claims 1 to 27 and 33 to 36.

41. Granules produced by the method as claimed in any one of claims 1 to 12.
42. A dough produced by the method as claimed in claim 13.
43. Pellets produced by the method as claimed in any one of claims 14 to 16.
44. The use of the release composition for delivery of biological material via a spray for application to plants and animals, characterised in that the biological material includes an active ingredient to be sprayed over plants and animals.
45. A spray solution as claimed in claim 44 that includes at least one release composition as claimed in claims 37 to 40.
46. According to another aspect of the present invention there is provided a method of inoculating a plant seed with a biological material, said method including the steps of;
- (a) selecting at least one biological material to be used as an inoculant;
 - (b) preparing the composition by the method as claimed in any one of claims 1 to 36;
 - (c) adding the composition to water and mixing to release the biological material into the solution; and
 - (d) soaking the plant seed in said solution to allow the biological material to coat the plant seed.
47. A method of inoculating a plant seed with a biological material as claimed in claim 46 wherein said method includes a further step after step (b), of:
- (bi) adding a further powdered compound to said composition, said powdered compound being selected from the group: a second biological material, a dried and powdered composition, a dried and powdered bio-polymer matrix containing at least a second biological material, a chemical, and a combination of these.
48. A method as claimed in either claim 46 or claim 47 wherein more than one inoculant is used in step (a) each inoculant being for a different purpose.

49. A method as claimed in either claim 47 or claim 48 wherein the plant seed is dried at room temperature before drilling or seed broadcast.

50. Seed inoculated by the method as claimed in any one of claims 46 to 49.

51. A composition for application to a substrate (as hereinbefore defined) wherein said composition includes:

- (a) one or more fast release compositions as produced by the method of any one of claims 1 to 27 and 33 to 36; and
- (b) one or more slow release compositions as produced by the method of any one of claims 1 to 32.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/NZ01/00168

A. CLASSIFICATION OF SUBJECT MATTER																						
Int. Cl. ⁷ : A01N 63/00, A01N 25/22, A61K 047/36, C12N 1/00																						
According to International Patent Classification (IPC) or to both national classification and IPC																						
B. FIELDS SEARCHED																						
Minimum documentation searched (classification system followed by classification symbols)																						
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched																						
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) DWPIDS, CA, AGRICOLA, MEDLINE; KEYWORDS: GUM, STARCH, GELLAN, GEL, STORAGE, STABLE, THERMOSTABLE, RHIZOBIUM, BCG, SERRATIA, PSEUDOMONAS,																						
C. DOCUMENTS CONSIDERED TO BE RELEVANT																						
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.																				
X	US 5292507 A (CHARLEY) 8 March 1994 Column 3 line 32 to column 4 line 64, claims	1-22, 25-51																				
X	US 5916029 A (SMITH et al) 29 June 1999 Whole document	1-22, 25-27, 37-38, 41-51																				
X	WO 92/20229 A (KOREA RES. INST. OF CHEMICAL TECHNOLOGY) 26 November 1992 Page 3 line 13 to page 7 line 19, Table 1, Claims	1-22, 25-51																				
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Date of the actual completion of the international search 26 October 2001		Date of mailing of the international search report - 1 NOV 2001																				
Name and mailing address of the ISA/AU AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustalia.gov.au Facsimile No. (02) 6285 3929		Authorized officer ROSS OSBORNE Telephone No : (02) 6283 2404																				

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ01/00168

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5021350 A (JUNG) 4 June 1991 Whole document	1, 2, 21, 25-30, 37-38, 46-50
X	US 5401506 A (CHANG) 28 March 1995 Whole document	1, 2, 21-22, 25-30, 37-39, 41-43
X	US 5686385 A (AKASHI) 11 November 1997 Column 3 line 59 to column 6 line 53, claim 1, 14	1-16, 22, 28-29, 33-51
X	EP 234670 A (BOOTS COMPANY PLC) 2 September 1987 Whole document	1, 2, 39, 51

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.
PCT/NZ01/00168

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Patent Document Cited in Search Report				Patent Family Member			
US	5292507	CA	1300538				
US	5916029	AU	25600/97	CA	2208896	EP	818135
WO	92/20229	AU	17587/92	BR	9205317	CN	1066959
		US	5273749	EP	540713		
EP	234670	AU	67625/87	AU	79462/91	CA	1313133
		FI	870155	IL	81281	NO	870181
		NZ	218907	US	5415871	ZA	8700173
US	5021350	AU	81153/82	BR	8201151	CA	1179618
		ES	510174	FI	820572	FR	2501229
		SE	8201348	US	5021350		
US	5401506	AU	91265/91	EP	561990	WO	9210170
		GB	2255189				
US	5686385	CA	2086077	EP	548901	US	5686385
		JP	5238904				
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