



## (51) International Patent Classification:

C05G 3/00 (2006.01) C05C 1/02 (2006.01)  
C05C 9/00 (2006.01) C05C 11/00 (2006.01)

## (21) International Application Number:

PCT/US2015/027725

## (22) International Filing Date:

27 April 2015 (27.04.2015)

## (25) Filing Language:

English

## (26) Publication Language:

English

## (30) Priority Data:

61/985,177 28 April 2014 (28.04.2014) US

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## (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM,

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(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

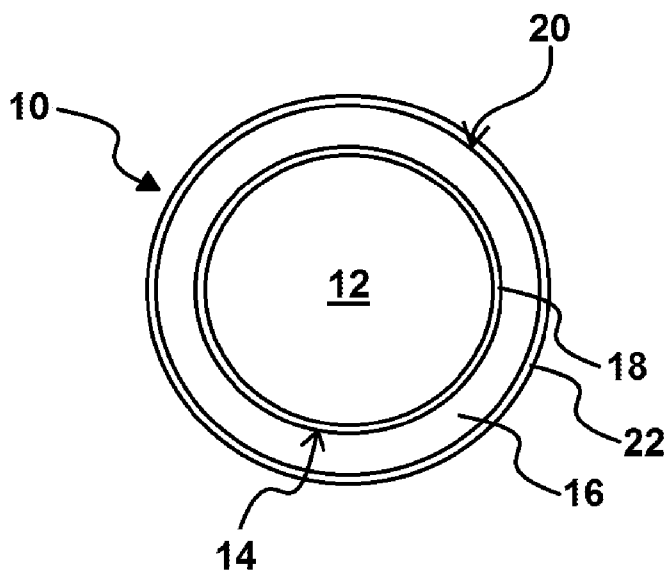
## Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))
- of inventorship (Rule 4.17(iv))

## Published:

- with international search report (Art. 21(3))

## (54) Title: METHOD OF FORMING AN ENCAPSULATED FERTILIZER

**FIG - 4**

(57) Abstract: A method of forming an encapsulated fertilizer comprises the step of providing a core particle. The core particle comprises a fertilizer and has an outer surface. An example of the fertilizer is urea. The method further comprises the step of applying an isocyanate component directly to the outer surface of the core particle. The isocyanate component has isocyanate functional groups. An example of the isocyanate component is an isocyanate-terminated prepolymer. A specific example of the isocyanate-terminated prepolymer is one comprising the reaction product of a diphenylmethane diisocyanate (MDI) and a polyol. The method further comprises the step of reacting the isocyanate component with ambient moisture to form a layer disposed about the core particle to form the encapsulated fertilizer. The layer comprises polyurea linkages. The layer may further comprise polyurethane linkages. For example, the layer may be in the form of a polyurea/polyurethane layer when the isocyanate-terminated prepolymer is utilized.

## METHOD OF FORMING AN ENCAPSULATED FERTILIZER

### FIELD OF THE INVENTION

[0001] The present invention generally relates to a method of forming an encapsulated fertilizer and to an encapsulated fertilizer formed according to the method, specifically to a method of forming an encapsulated fertilizer comprising a core particle and a layer disposed about the core particle with the layer comprising the reaction product of an isocyanate component and ambient moisture.

### DESCRIPTION OF THE RELATED ART

[0002] Encapsulated fertilizers, such as controlled-release fertilizers, are known in the agricultural art. Such encapsulated fertilizers typically include one or more polyurethane layers disposed about a core particle, typically a fertilizer (e.g. urea). Thickness and integrity of the polyurethane layer limits dissolution rate of the encapsulated fertilizers, specifically, a rate that the core particle degrades releasing itself, e.g. a “payload”, into soil that includes moisture.

[0003] Unfortunately, many conventional encapsulated fertilizers include inconsistent thickness of the polyurethane layer, which results in accelerated dissolution rates. Such accelerated dissolution rates lead to waste and phytotoxicity, i.e., toxic effect of the fertilizer on plant growth. Further, many conventional encapsulated fertilizers have polyurethane layers with integrity problems, since the polyurethane layers include defects, such as cracks, pits, depressions, etc. When a polyurethane layer including such defects is disposed about the core particle, the cracks, pits, and/or depressions allow water and other liquids to permeate the polyurethane layer, thus prematurely contacting and dissolving the core particle. To cure such defects, multiple polyurethane layers must typically be disposed about the core particle resulting in a time consuming and expensive manufacturing process for forming the encapsulated fertilizers. Yet further, many conventional encapsulated fertilizers are prone to breakage, due to the polyurethane layer lacking adequate hardness and resiliency. Such breakage reduces shelf life of the encapsulated fertilizers, and results in waste during storage, handling, and use of the encapsulated fertilizers.

[0004] Conventional polyurethane layers are typically based on a two component ("2K") urethane system where one part is an isocyanate and the other part is a polyol. An excess of the isocyanate relative to the polyol is often used because of an isocyanate/water side-reaction from water that is present during the coating process.

Many processes also require the isocyanate and the polyol to be applied at different times or at the exact same time, which adds to inconsistencies during manufacture. There is also the issue of inconsistent moisture during application of the reactive components. If too much water is present during application of the reactive components, there may not be enough isocyanate left to react with all the polyol. This is because a polyurea reaction (between water and isocyanate groups) can be faster than a polyurethane reaction (between polyol and isocyanate groups). This can lead to inconsistent properties of the resulting layer, with unreacted polyol lowering performance of the encapsulated fertilizer.

[0005] Accordingly, there remains an opportunity to provide improved methods of forming encapsulated fertilizers. There also remains an opportunity to provide improved encapsulated fertilizers.

#### SUMMARY OF THE INVENTION AND ADVANTAGES

[0006] Disclosed is a method of forming an encapsulated fertilizer. The method comprises the step of providing a core particle. The core particle comprises a fertilizer and has an outer surface. The method further comprises the step of applying an isocyanate component directly to the outer surface of the core particle. The isocyanate component has isocyanate functional groups. The method further comprises the step of reacting the isocyanate component with ambient moisture to form a layer disposed about the core particle to form the encapsulated fertilizer. The layer comprises polyurea linkages. Optionally, the layer may further comprise polyurethane linkages.

[0007] Generally, using only one component (i.e., the isocyanate component) simplifies application and increases manufacturing output of the encapsulated fertilizer relative to systems utilizing two or more reactive components, e.g. an isocyanate and a polyol. A more consistent and better coating layer can be achieved since generally the only reaction occurring after the isocyanate component is applied is a polyurea reaction. Having a consistent coating layer also provides a more standardized time release of the fertilizer.

#### BRIEF DESCRIPTION OF THE DRAWINGS

[0008] Other advantages of the present invention will be readily appreciated, as the same becomes better understood by reference to the following detailed description when considered in connection with the accompanying drawings wherein:

[0009] Figure 1 is a cross-section of an encapsulated fertilizer having a core particle and a layer;

[0010] Figure 2 is a cross-section of another encapsulated fertilizer further having an intermediate layer;

[0011] Figure 3 is a cross-section of another encapsulated fertilizer further having a sealant layer;

[0012] Figure 4 is a cross-section of another encapsulated fertilizer having both intermediate and sealing layers; and

[0013] Figure 5 is a chart illustrating dissolution rates of example encapsulated fertilizers.

#### DETAILED DESCRIPTION OF THE INVENTION

[0014] Disclosed is a method of forming an encapsulated fertilizer. Also disclosed is an encapsulated fertilizer formed according to the method. Referring to the Figures, wherein like numerals indicate like parts throughout the several views, an encapsulated fertilizer formed according to the method is shown generally at **10**. The encapsulated fertilizer **10** is not drawn to scale.

[0015] The method comprises the step of providing a core particle **12**. The core particle **12** has an outer surface **14**. The method further comprises the step of applying an isocyanate component directly to the outer surface **14** of the core particle **12**. The method further comprises the step of reacting the isocyanate component with ambient moisture to form a layer **16** disposed about the core particle **12** to form the encapsulated fertilizer **10**. The layer **16** comprises polyurea linkages. In certain embodiments, the layer **16** further comprises polyurethane linkages.

[0016] The encapsulated fertilizer **10** comprises the core particle **12**. The core particle **12** comprises a fertilizer. In various embodiments, the outer surface **14** of the core particle **12** comprises the fertilizer. In specific embodiments, the entire core particle **12** comprises the fertilizer. In certain embodiments, the core particle **12** consists essentially of the fertilizer. In further embodiments, the core particle **12** consists of (i.e., is) the fertilizer.

[0017] In various embodiments, the fertilizer is selected from the group of nitrogen, phosphate, potash, sulfur, and combinations thereof. In certain embodiments, the fertilizer is nitrogen based. Examples of suitable nitrogen based fertilizers include anhydrous ammonia, urea, ammonium nitrate, urea ammonium nitrate, calcium

ammonium nitrate, and combinations thereof. In specific embodiments, the fertilizer comprises, consists essentially of, or consists of, urea. In other embodiments, the fertilizer is phosphate based. Examples of suitable phosphate based fertilizers include phosphoric acid, mono-ammonium phosphate, ammonium polyphosphate, ammonium phosphate sulfate, and combinations thereof. In yet other embodiments, the fertilizer is potash based. Examples of suitable potash based fertilizers include potash, ammonium nitrate, and combinations thereof. In yet other embodiments, the fertilizer is sulfur based. Examples of suitable sulfur based fertilizers include ammonium sulfate, sulfuric acid, and combinations thereof. Various combinations of fertilizers can be utilized as, or in, the core particle **12**. Suitable fertilizers are commercially available from a variety of suppliers, and this disclosure is not limited to any particular one.

**[0018]** Typically, the encapsulated fertilizer **10** includes one of the core particles **12**, e.g. as shown in the Figures. However, the encapsulated fertilizer **10** can also include any combination of two or more of the aforementioned core particles **12**. In such embodiments, the core particle **12** can comprise, but is not limited to, a blend of core particle compositions, individual sub-particles of the core particle **12**, and/or layers of different core particle compositions. For example, the core particle **12** can comprise an inner-core comprising sulfur with an outer-core disposed about the inner-core and comprising urea (not shown).

**[0019]** The core particle **12** can be of various sizes and shapes. Typically, the core particle **12** is substantially spherical, having an average diameter of from about 0.1 to about 5, about 0.1 to about 2.5, about 0.25 to about 1, millimeters (mm), or any average diameter between about 0.1 to about 5 mm. Other sizes and/or shapes of core particles **12** can also be used, such as irregular, oblong, and/or platelet shaped particles.

**[0020]** In various embodiments, the core particle **12** has a moisture content prior to the step of applying. The core particle **12** can have various moisture contents. Typically, the core particle **12** has a moisture content of at least about 1 weight percent (wt.%), alternatively a moisture content of from about 1 to about 20, about 1 to about 10, about 5 to about 10, wt.%, or any moisture content between about 1 and about 20 wt.%.

[0021] The moisture content may be imparted by purposeful addition (e.g. by soaking, spraying, coating, dipping, etc.), but is more typically innate. For example, certain fertilizers (e.g. urea), are hygroscopic. Hygroscopic fertilizers readily absorb moisture from the atmosphere. This can occur during routine manufacture, handling, and/or storage of the fertilizer. In other words, no purposeful addition of moisture/water is required. The moisture content of the core particle **12** is useful for establishing the ambient moisture as described further below. In addition, the moisture content of the core particle **12** can also provide surface moisture on or at the outer surface **14** of the core particle **12**. In these instances, the remainder of the moisture content is generally present throughout the core particle **12** and/or is generally present as a moisture content gradient.

[0022] In embodiments where the outer surface **14** of the core particle **12** comprises surface moisture, the step of reacting is further defined as reacting the isocyanate component with ambient moisture and the surface moisture to form the encapsulated fertilizer **10**. In this way, reaction occurs on both sides of the isocyanate component after application to the outer surface **14** of the core particle **12**. Specifically, isocyanate functional groups provided by the isocyanate component are free to react with surface moisture on the outer surface **14** of the core particle **12**, as well as with ambient moisture at the surface **20** of the layer **16** (during formation).

[0023] In other embodiments, the outer surface **14** of the core particle **12** comprises amine functional groups (i.e., “surface amines”). The surface amines can be provided by the core particle **12**, such as when the core particle **12** comprises urea as the fertilizer. In these embodiments, the step of reacting is further defined as reacting the isocyanate component with ambient moisture and the amine functional groups to form the encapsulated fertilizer **10**. In this way, reaction occurs on both sides of the isocyanate component after application to the outer surface **14** of the core particle **12**. Specifically, isocyanate functional groups provided by the isocyanate component are free to react with amine groups on the outer surface **14** of the core particle **12**, as well as with ambient moisture at the surface **20** of the layer **16** (during formation).

[0024] In related embodiments, the outer surface **14** of the core particle **12** comprises both surface moisture and surface amines. In these embodiments, the step of reacting is further defined as reacting the isocyanate component with ambient moisture, the surface moisture, and the amine functional groups to form the encapsulated fertilizer

**10.** This reaction scheme is a combination of the aforementioned embodiments. Specifically, isocyanate functional groups provided by the isocyanate component are free to react with the surface moisture and surface amines on the outer surface **14** of the core particle **12**, as well as with ambient moisture at the surface **20** of the layer **16** (during formation).

**[0025]** The core particle **12** can be present in the encapsulated fertilizer **10** in various amounts. Typically, the core particle **12** is present in the encapsulated fertilizer **10** in an amount of from about 75 to about 99, about 90 to about 99, about 96 to about 98, parts by weight (pbw), or any amount between about 75 and about 99 pbw, each based on 100 pbw of the encapsulated fertilizer **10**.

**[0026]** As shown in the Figures, the layer **16** is disposed about the core particle **12**. It is to be understood that as used herein, the terminology “disposed about” encompasses both partial and complete covering of the core particle **12**. Typically, the core particle **12** is completely covered by the layer **16**, thereby protecting the core particle **12** from exposure to environmental conditions (such as from premature exposure to moisture).

**[0027]** The layer **16** can be of various average thicknesses. The layer **16** typically has an average thickness that mimics thicknesses of conventional encapsulation layers, e.g. conventional fertilizer encapsulation layers. In various embodiments, the layer **16** has an average thickness of at least about 5, at least about 10, at least about 15, alternatively of from about 15 to about 300, about 50 to about 200, about 100 to 150, about 125 to about 150, microns, or any average thickness between about 5 and about 300 microns. The layer **16** can be formed into various thicknesses depending on one or more desired properties, such as dissolution rate of the encapsulated fertilizer **10**. Therefore, the average thicknesses can be less or greater than those specifically described herein. In certain embodiments, as in Figures 1 and 3, the layer **16** is in direct contact with the core particle **12**. In other words, there is no intervening layer (or portion thereof) between the layer **16** and the outer surface **14** of the core particle **12**.

**[0028]** With respect to various embodiments of the layer **16**, it is to be understood that the terminology “polyurea” encompasses chemistries that include both polyurea and/or urea linkages. In various embodiments, the layer **16** is defined as a polyurea layer **16**. In further embodiments, the layer **16** includes polyurea linkages as well as

additional linkages, e.g. polyurethane and/or urethane linkages. For example, in certain embodiments described below, the layer **16** is further defined as a polyurea/polyurethane layer **16**. As used herein, description of the layer **16** is generally interchangeable with description of the polyurea layer **16** or the polyurea/polyurethane layer **16**.

**[0029]** Typically, the layer **16** comprises the reaction product of the isocyanate component and water. Specifically, the layer **16** comprises the reaction product of the isocyanate component and water that is initially in the form of ambient moisture. In further embodiments, the layer **16** consists essentially of the reaction product of the isocyanate component and water. In yet further embodiments, the layer **16** consists of (i.e., is) the reaction product of the isocyanate component and water. In these embodiments, a portion of the water can also be imparted by surface moisture rather than entirely by ambient moisture. In addition or alternate to reaction with the surface moisture, the surface amines may also contribute to the reaction product.

**[0030]** The layer **16** can be present in the encapsulated fertilizer **10** in various amounts. Typically, the layer **16** is present in the encapsulated fertilizer **10** in an amount of from about 25 to about 1, about 10 to about 1, about 4 to about 2, pbw, or any amount between about 25 and about 1 pbw, each based on 100 pbw of the encapsulated fertilizer **10**. Generally, this disclosure provides for less material use, e.g. less isocyanate component use, relative to prior art encapsulated fertilizers.

**[0031]** The isocyanate component has isocyanate functional groups. The isocyanate component typically comprises a polyisocyanate having two or more isocyanate functional (NCO) groups. Suitable polyisocyanates include, but are not limited to, conventional aliphatic, cycloaliphatic, araliphatic and aromatic isocyanates. In certain embodiments, the isocyanate component is selected from the group of diphenylmethane diisocyanates (MDIs), polymeric diphenylmethane diisocyanates (PMDIs), and combinations thereof. Polymeric diphenylmethane diisocyanates are also referred to in the art as polymethylene polyphenylene polyisocyanates. In specific embodiments, the isocyanate component comprises a MDI. Examples of other suitable isocyanates include toluene diisocyanates (TDIs), hexamethylene diisocyanates (HDIIs), isophorone diisocyanates (IPDIIs), naphthalene diisocyanates (NDIIs), and combinations thereof.



[0032] In various embodiments, the isocyanate component comprises, consists essentially of, or consists of (i.e., is) an isocyanate-terminated prepolymer. The isocyanate-terminated prepolymer is typically the reaction product of an isocyanate and a polyol and/or a polyamine. The isocyanate may be any type of isocyanate, such as one or more of the aforementioned polyisocyanates. Various types of polyols and polyamines can be utilized.

[0033] Utilizing one or more polyols is useful for imparting polyurethane linkages in the isocyanate-terminated prepolymer and therefore, in the layer **16**. Similarly, utilizing one or more polyamines is useful for imparting polyurea linkages in the isocyanate-reactive component, and therefore, in the layer **16**. Reaction between isocyanate and hydroxyl groups to form polyurethane linkages is understood in the art. Moreover, reaction between isocyanate and amine groups to form polyurea linkages is understood in the art. One of skill in the art can readily envisage embodiments of the layer **16** having only polyurea linkages or both polyurea and polyurethane linkages depending on the particular isocyanate component utilized.

[0034] In certain embodiments, the isocyanate-terminated prepolymer comprises the reaction product of a MDI and a polyol having at least two hydroxyl functional groups. In these embodiments, the isocyanate-terminated prepolymer comprises polyurethane linkages such that the layer **16** is a polyurea/polyurethane layer **16**. Further, the isocyanate component may also include free isocyanate in addition to the isocyanate-terminated prepolymer. In these instances, the isocyanate-terminated prepolymer can be present in the isocyanate component in various amounts. Typically, the isocyanate-terminated prepolymer is present in an amount of from about 10 to about 90, about 25 to about 75, about 40 to about 60, about 45 to about 55, pbw, or any amount between about 25 and about 75 pbw, each based on 100 pbw of the isocyanate component.

[0035] It is thought that utilizing the isocyanate-terminated prepolymer provides one or more non-limiting benefits as follows. Using the isocyanate-terminated prepolymer can ensure a more consistent reaction between the polyol (and/or polyamine) and the isocyanate leading to a consistent layer **16** with desirable hydrophobic properties. Specifically, there are no competing reactions during the application/coating process (polyurea versus polyurethane reaction), which leads to a layer **16** with improved and more consistent properties (e.g. hardness, tack, surface tension, hydrophobicity,

toughness, etc.). Typically, there aren't issues of mix ratio for the two components (e.g. isocyanate and polyol) since only one component (i.e., the prepolymer) is being applied during formation of the encapsulated fertilizer **10**. Therefore, there is no need for complex or multiple pieces of application equipment. Applying one component instead of two components also leads to faster application times, improved layers **16**, and/or increased output of the encapsulated fertilizer **10**.

**[0036]** If utilized to make the isocyanate-terminated prepolymer, the polyol is typically selected from the group of ethylene glycol, diethylene glycol, propylene glycol, dipropylene glycol, butane diol, glycerol, trimethylolpropane, triethanolamine, pentaerythritol, sorbitol, and combinations thereof. In certain embodiments, the polyol utilized to form the isocyanate-terminated prepolymer is a graft polyol. In one embodiment, the graft polyol is a polymer polyol. In other embodiments, the graft polyol is selected from the group of polyharnstoff (PHD) polyols, polyisocyanate polyaddition (PIPA) polyols, and combinations thereof.

**[0037]** If utilized to make the isocyanate-terminated prepolymer, the polyamine is typically selected from the group of ethylene diamine, toluene diamine, diaminodiphenylmethane and polymethylene polyphenylene polyamines, aminoalcohols, and combinations thereof. Examples of suitable aminoalcohols include ethanolamine, diethanolamine, triethanolamine, and combinations thereof. The isocyanate-terminated prepolymer may be formed from a combination of two or more of the aforementioned polyols and/or polyamines.

**[0038]** The isocyanate component may also be a modified isocyanate, such as, carbodiimides, allophanates, isocyanurates, and biurets. Other suitable isocyanates and optional components include those described in U.S. Pat. Nos. 7,416,785 to Mente and 8,303,680 to Mente, in U.S. Pat. App. Pub. Nos. 2013/0305796 to Hudson et al. and 2013/0309499 to Neff et al., and in WO2012151506 to Neff et al., the disclosures of which are incorporated herein by reference in their entirety. Specific examples of suitable isocyanate components include those that are commercially available from BASF Corporation of Florham Park, NJ, under the trademark LUPRANATE<sup>®</sup>, such as LUPRANATE<sup>®</sup> M, LUPRANATE<sup>®</sup> MI, LUPRANATE<sup>®</sup> M20, LUPRANATE<sup>®</sup> M20SB, LUPRANATE<sup>®</sup> M20HB, LUPRANATE<sup>®</sup> M20FB, and LUPRANATE<sup>®</sup> MP102, isocyanates.

[0039] The isocyanate component can be of various viscosities. Typically, the isocyanate component has a viscosity which is suitable for specific applications of the isocyanate component to the core particle **12**, such as by spraying, fogging, and/or atomizing the isocyanate component to apply the isocyanate component to the outer surface **14** of the core particle **12**. In certain embodiments, the isocyanate component has a viscosity of from about 10 to about 5,000, about 25 to about 2,500, about 50 to about 1,500, about 100 to about 1,000, about 500 to about 1,000, cps at 25°C, or any viscosity between about 10 to about 5,000 cps at 25°C, according to ASTM D2196. Regardless of the application technique, the viscosity of the isocyanate component should be sufficient to adequately coat the core particle **12**.

[0040] The isocyanate component can be of various nominal functionalities and NCO contents. In certain embodiments, the isocyanate component has a nominal functionality of from about 1 to about 5, about 1.5 to about 4, about 2 to about 3, or any nominal functionality between about 1 and about 5. In certain embodiments, the isocyanate component has an NCO content of from about 1 to about 50, about 1 to about 40, about 1 to about 30, about 10 to about 30, about 20 to about 25, wt. %, or any NCO content between about 20 and about 50 wt.%, each based on 100 pbw of the isocyanate component. The aforementioned NCO contents generally provide a high molecular crosslink density of the isocyanate component that aids in the formation of a defect free layer **16**, and also provide the isocyanate component with more chemical bonds per unit of mass to improve cost efficiency. The isocyanate component may include any combination of the aforementioned isocyanates and/or isocyanate-terminated prepolymers.

[0041] The reaction product may further comprise an additive component. If utilized, the additive component is typically selected from the group of parting agents, catalysts, biocides, fillers, plasticizers, stabilizers, cross-linking agents, chain-extending agents, chain-terminating agents, air releasing agents, wetting agents, surface modifiers, moisture scavengers, desiccants, viscosity reducers, reinforcing agents, colorants, anti-oxidants, compatibility agents, ultraviolet light stabilizers, thixotropic agents, anti-aging agents, lubricants, coupling agents, solvents, rheology promoters, thickeners, anti-static agents, and combinations thereof. If utilized, the additive component may be present in customary amounts. The additive component may include any combination of the aforementioned additives. The additive

component may be reactive and/or inert with respect to the isocyanate component depending on the additive(s) utilized.

**[0042]** In certain embodiments, the additive component comprises a catalyst component. In other words, the layer **16** is formed in the presence of a catalyst. In one embodiment, the catalyst component comprises a tin catalyst. Suitable tin catalysts include tin(II) salts of organic carboxylic acids, e.g. tin(II) acetate, tin(II) octoate, tin(II) ethylhexanoate and tin(II) laurate. In one embodiment, the organometallic catalyst comprises dibutyltin dilaurate, which is a dialkyltin(IV) salt of an organic carboxylic acid. Specific examples of suitable organometallic catalyst, e.g. dibutyltin dilaurates, are commercially available from Air Products and Chemicals, Inc. of Allentown, PA, under the trademark DABCO<sup>®</sup>. The organometallic catalyst can also comprise other dialkyltin(IV) salts of organic carboxylic acids, such as dibutyltin diacetate, dibutyltin maleate and dioctyltin diacetate.

**[0043]** Examples of other suitable catalysts include iron(II) chloride; zinc chloride; lead octoate; tris(dialkylaminoalkyl)-s-hexahydrotriazines including tris(N,N-dimethylaminopropyl)-s-hexahydrotriazine; tetraalkylammonium hydroxides including tetramethylammonium hydroxide; alkali metal hydroxides including sodium hydroxide and potassium hydroxide; alkali metal alkoxides including sodium methoxide and potassium isopropoxide; and alkali metal salts of long-chain fatty acids having from 10 to 20 carbon atoms and/or lateral OH groups.

**[0044]** Further examples of other suitable catalysts, specifically trimerization catalysts, include N,N,N-dimethylaminopropylhexahydrotriazine, potassium, potassium acetate, N,N,N-trimethyl isopropyl amine/formate, and combinations thereof. A specific example of a suitable trimerization catalyst is commercially available from Air Products and Chemicals, Inc. under the trademark POLYCAT<sup>®</sup>.

**[0045]** Yet further examples of other suitable catalysts, specifically tertiary amine catalysts, include dimethylaminoethanol, dimethylaminoethoxyethanol, triethylamine, N,N,N',N'-tetramethylethylenediamine, N,N-dimethylaminopropylamine, N,N,N',N',N''-pentamethyldipropylenetriamine, tris(dimethylaminopropyl)amine, N,N-dimethylpiperazine, tetramethylimino-bis(propylamine), dimethylbenzylamine, trimethylamine, triethanolamine, N,N-diethyl ethanolamine, N-methylpyrrolidone, N-methylmorpholine, N-ethylmorpholine, bis(2-dimethylamino-ethyl)ether, N,N-dimethylcyclohexylamine (DMCHA), N,N,N',N',N''-pentamethyldiethylenetriamine,

1,2-dimethylimidazole, 3-(dimethylamino) propylimidazole, and combinations thereof. Specific examples of suitable tertiary amine catalysts are commercially available from Air Products and Chemicals, Inc. under the trademark POLYCAT<sup>®</sup>.

[0046] If utilized, the catalyst component can be employed in customary amounts. The catalyst component may include any combination of the aforementioned catalysts. In certain embodiments, the encapsulated fertilizer **10** is formed in the absence of a catalyst.

[0047] In embodiments utilizing one or more catalysts, the catalyst may be purposefully added for reaction between the isocyanate component and ambient moisture. Alternatively, or in addition, one or more catalysts may already be present in the isocyanate component prior to reaction with ambient moisture. For example, in instances where the isocyanate-terminated prepolymer is utilized, one or more catalysts may have been used to initially form the prepolymer (e.g. a tertiary amine catalyst) and such catalysts can still be present during formation of the layer **16**.

[0048] In certain embodiments, the additive component comprises a colorant, such as a dye and/or a pigment, for coloring the layer **16**. The colorant allows completeness of the layer **16** to be visually evaluated and can provide various marketing advantages. If utilized, the colorant can be utilized in customary amounts.

[0049] In certain embodiments, the outer surface **14** of the core particle **12** is at least partially coated, alternatively is completely coated, with a compound other than fertilizer. For example, as shown in Figures 2 and 4, the encapsulated fertilizer **10** includes an intermediate layer **18** disposed between the core particle **12** and layer **16**. Specifically, the intermediate layer **18** is disposed about the core particle **12** and the layer **16** is disposed about the intermediate layer **18**.

[0050] The intermediate layer **18** may be a partial or a complete covering. The intermediate layer **18** is typically formed from the compound. Typically, the intermediate layer **18** is already present on the core particle **12** prior to formation of the layer **16**. For example, the intermediate layer **18** can be a protective layer for the core particle **12**.

[0051] In various embodiments, the compound can be referred to in the art as a dust suppressant. Dust suppressants are useful for handling the core particle **12**, and may be present for other purposes in addition or alternate to dust suppression (e.g. to prevent agglomeration, skin contact, moisture absorption, etc.). Dust suppressants are

typically liquids, such as oils, but can be solids, such as waxes. Particular examples of dust suppressants are petroleum residue, hydrogenated mineral oil, and wax. Suitable examples of waxes which may be used include, but are not limited to, organic waxes, thermoplastic polymers, mineral oils, or combinations thereof. More particularly, the wax may comprise paraffinic oil, paraffinic wax, vegetable wax, triglyceride, microcrystalline wax, petrolatum, olefin, polyethylene, petroleum wax, and combinations thereof. In one embodiment, the wax comprises a petroleum wax. In another embodiment, the wax comprises a petroleum wax and at least one other wax.

[0052] While optional, the dust suppressant can be utilized in various amounts. Typically, the intermediate layer **18** is present in an amount of from about 0.1 to about 10, about 0.1 to about 5, about 0.5 to about 3, pbw, or any amount between about 0.1 and about 10 pbw, each based on 100 pbw of the core particle **12**. The intermediate layer **18** can be of various average thicknesses, such as those above for the layer **16**.

[0053] Typically, the layer **16** is substantially free of water after formation. By “substantially free”, it is meant that in these embodiments, water is present in an amount no greater than about 5, no greater than about 2.5, no greater than about 1, or approaching or equaling 0, pbw, each based on 100 pbw of the layer **16**. If the layer **16** is completely free of water, the layer **16** is classified as anhydrous. It is generally preferred that the layer **16** has little to no water to prevent the encapsulated fertilizer **10** from prematurely degrading, such as by softening, rupturing, or sticking/agglomerating to other adjacent encapsulated fertilizers **10** or surfaces. Such degradation can cause premature release of the core particle **12**, and waste.

[0054] In certain embodiments, the layer **16** is substantially free of a wax, such as paraffin. By “substantially free”, it is meant that in these embodiments, wax is present in an amount no greater than about 5, no greater than about 2.5, no greater than about 1, or approaching or equaling 0, pbw, each based on 100 pbw of the layer **16**. Typically, the layer **16** has little to no wax to allow for a uniform layer **16**, which can prevent water from permeating the layer **16**. However, wax can be utilized in the layer **16** if compatible with the specific components utilized, e.g. the isocyanate component. Customary waxes and amounts thereof can be utilized. In certain embodiments, the encapsulated fertilizer **10** is formed in the absence of wax.

[0055] In certain embodiments, the layer **16** is substantially free of sulfur. By “substantially free”, it is meant that in these embodiments, sulfur is present in an

amount no greater than about 5, no greater than about 2.5, no greater than about 1, or approaching or equaling 0, pbw, each based on 100 pbw of the layer **16**. The method will now be described in further detail.

**[0056]** Referring to Figures 3 and 4, the encapsulated fertilizer **10** may include a sealant layer **22** disposed about the layer **16**. The sealant layer **22** may seal imperfections in the layer **16**, may improve the flow characteristics of the encapsulated fertilizer **10**, and may provide improved moisture barrier properties. The sealant layer **22** may comprise organic waxes, such as paraffinic wax, vegetable wax, triglycerides, microcrystalline thermoplastic polymers, mineral oils, petroleum waxes or combinations thereof. More particularly, the sealant layer **22** may comprise paraffinic oil, paraffinic wax, vegetable wax, triglyceride, microcrystalline wax, petrolatum, olefin, polyethylene, and combinations thereof. Further suitable sealants for forming the sealant layer **22** are described in U.S. Pat. App. Pub. No. 2013/0305796 to Hudson et al.

**[0057]** The sealant layer **22** may comprise other materials other than those described above. The sealant layer **22** may include various conventional additives. For example, the sealant layer **22** may include various polymers to improve the abrasion resistance and barrier properties, anti-blocking adjuvants to improve the handling characteristics, dyes, and other known adjuvants.

**[0058]** Generally, the sealant layer **22** has a melting point below the melting point of the core particle **12** and the layer **16**. For example, the sealant layer **22** may have a melting point below about 80 °C and flow at the temperature of its application process. The sealant layer **22** may be non-tacky after application to the layer **16** at temperatures below about 60, 50, or 40 °C. Furthermore, the sealant layer **22** may have a melt viscosity less than about 1,500 cP or less than about 1,000 cP at its application temperature to allow for good flowability on the core particle **12** having the layer **16**.

**[0059]** While optional, the sealant can be utilized in various amounts. For example, the encapsulated fertilizer **10** may include the sealant layer **22** in various amounts depending on the desired release rate of the encapsulated fertilizer **10**. Typically, the sealant layer **22** is present in an amount of from about 0.1 to about 10, about 0.1 to about 5, about 0.5 to about 3, pbw, or any amount between about 0.1 and about 10

pbw, each based on 100 pbw of the encapsulated fertilizer **10**. The sealant layer **22** can be of various average thicknesses, such as those above for the layer **16**.

**[0060]** As introduced above, the core particle **12** is provided. The core particle **12** can be provided in various manners, such as in a free flowing particulate form. Typically, the core particle **12** is disposed in a vessel to retain the core particle **12** and assist in formation of the encapsulated fertilizer **10**. The vessel can be of various sizes and shapes, and should be closable or sealable to retain components therein during manufacture of the encapsulated fertilizer **10**. The core particle **12** can be provided with or without the intermediate layer **18**.

**[0061]** In various embodiments, the method further comprises the step of heating the core particle **12**. The core particle **12** can be heated via direct and/or indirect heat. For example, the vessel can be heated (e.g. via a heat exchanger), the air in the vessel can be heated, the core particle **12** can be heated, etc., and combinations thereof. While ambient air in the vessel may be relied upon to provide ambient moisture for reaction with the isocyanate component, heating the core particle **12** is useful to establish at least a portion of the ambient moisture, alternatively a majority (to all) of the ambient moisture, for reaction with the isocyanate component. This is especially the case when the fertilizer of the core particle **12** is hygroscopic (e.g. the fertilizer comprises urea). Heating the core particle **12** can also be useful for indirectly heating the isocyanate component, facilitate reaction, etc.

**[0062]** In certain embodiments, the core particle **12** is heated to a temperature of at least about 20 °C, alternatively to a temperature of from about 15 to about 45, about 40 to about 90, about 60 to about 120, °C, or any temperature between about 0 and 200 °C. The core particle **12** can be heated for various periods of time. To retain/establish ambient moisture, and especially ambient moisture provided by the core particle **12**, the vessel should be closed. Closing the vessel is also useful for maintaining temperature. In specific embodiments, the vessel is heated to heat the core particle **12** disposed therein. In further embodiments, the vessel is a tumbler, which allows for uniform mixing, heating, etc.

**[0063]** In certain embodiments, the method further comprises the step of heating the isocyanate component. This can be in addition or alternate to heating of the core particle **12**. The isocyanate component can be heated via direct and/or indirect heat. For example, the isocyanate component can be preheated (e.g. in a drum and/or



inline) before disposing the isocyanate component into the vessel. In certain embodiments, the isocyanate component is heated to a temperature of at least about 0 °C, alternatively to a temperature of from about 20 to about 40, about 25 to about 60, about 15 to about 80, °C, or any temperature between about 0 and 150 °C.

**[0064]** The isocyanate component is applied to the core particle **12**. Upon contact (and while under reaction conditions understood in the art), the isocyanate component and ambient moisture begin to react to form the layer **16** disposed about the core particle **12**. Specifically, it is thought that after application of the isocyanate component to the outer surface **14** of the core particle **12**, ambient moisture (i.e., water in the gaseous form) begins to react with free isocyanate groups provided by the isocyanate component to form polyurea linkages and therefore, the layer **16**. If the applied isocyanate component is at a temperature lower than that of the ambient air/moisture in the vessel, it is also thought that a portion of the ambient moisture condenses to liquid form on the applied isocyanate component, which further facilitates formation of the layer **16**. Reaction can also be occurring between the surface moisture and/or the surface amines (if present) and the isocyanate component.

**[0065]** The isocyanate component can be applied by various methods understood in the art. Examples of suitable methods of encapsulation include bulk coating, tumbling, sheet coating, pan coating, fluidized-bed coating, co-extrusion, spraying and spinning disk encapsulation, etc. These methods and other encapsulation methods are understood by those of ordinary skill in the art.

**[0066]** In certain embodiments, the isocyanate component is sprayed, atomized, and/or fogged onto the core particle **12**, typically a number of core particles **12**, while the core particles **12** are being agitated in suitable equipment. Spraying, atomizing, and/or fogging the isocyanate component onto the core particle **12** generally results in a uniform, complete, and defect-free layer **16** disposed about the core particle **12**. Spraying, atomizing, and/or fogging the isocyanate component also results in a thinner and less expensive layer **16** disposed about the core particle **12**. Several application steps can be used to build-up the thickness of the layer **16**. Spraying, atomizing and fogging of the isocyanate component can occur via use of one or more nozzles in the vessel. In specific embodiments, the isocyanate component is sprayed onto the core particle **12**.

[0067] In various embodiments to maximize coverage of the core particles **12**, the isocyanate component is generally applied by spraying droplets or atomizing or fogging particles of the isocyanate component onto the core particles **12** as the core particles **12** are being tumbled in a rotary blender or similar apparatus (i.e., as the vessel). As another example, the core particles **12** can be coated with the isocyanate component in a rotary drum blender equipped with at least one spinning disk atomizer. Tumblers, drums, or rollers including baffles can also be used as the vessel. Alternatively, the core particles **12** can be provided directly to a carrier, e.g. a screened conveyor belt and the isocyanate component can be applied to the core particles **12**, e.g. by spraying or sheeting, to form the encapsulated fertilizers **10**. Prior to the layer **16** fully curing, the encapsulated fertilizer **10** is typically kept moving to avoid agglomeration. Optionally, the sealant can later be applied in a similar manner as the isocyanate component above to form the sealant layer **22**.

[0068] The amount of the isocyanate component to be applied and mixed with the core particles **12** is dependant upon several variables including the specific isocyanate component, the size and type of core particle **12**, the intended end use of the encapsulated fertilizer **10**, the desired thickness of the layer **16**, the desired properties of the encapsulated fertilizer **10**, etc.

[0069] Typically, the method of this disclosure is free of the step of applying an isocyanate-reactive component to the core particle **12** to form the encapsulated fertilizer **10**. In other words, no isocyanate-reactive component is present on the outer surface **14** of the core particle **12**. The isocyanate-reactive component is typically selected from the group of polyols, polyamines, water, and combinations thereof. As described above, the isocyanate component is applied directly to the outer surface **14** of the core particle **12**. Therefore, no intervening layer formed from a purposefully applied isocyanate-reactive component is present. Some amount of surface moisture can be present, but this is outside the scope of purposefully applying water to the core particle **12**.

[0070] The encapsulated fertilizer **10** can be of various sizes and shapes. Typically, the encapsulated fertilizer **10** is substantially spherical, having an average diameter of from about 0.5 to about 7.5, about 0.5 to about 3, about 1 to about 2, mm, or any average diameter between about 0.5 and about 7.5 mm. However, the encapsulated

fertilizer **10** can be of other shapes, such as irregular, pocked, oblong or platelet shaped.

[0071] The encapsulated fertilizer **10** can be supplied to consumers in various manners. Typically, the encapsulated fertilizer **10** is supplied in bags, such as paper and/or plastic bags containing about 1 to about 80 pounds of the encapsulated fertilizer **10** for ease of handling and use. Such bags can include other nutrients and/or fillers in addition to the encapsulated fertilizer **10**, such as those typically included along with conventional fertilizers. In certain embodiments, the bags can include both encapsulated and non-encapsulated fertilizers. The bags can include different grades of encapsulated fertilizer **10**, such as ones having different dissolution rates, which can be imparted by different layer **16** thicknesses.

[0072] Various types of the encapsulated fertilizer **10** can be supplied to consumers in blends, such as encapsulated fertilizer **10** having different core particles **12**, e.g. different types of fertilizers, such as nitrogen and potash, nitrogen and sulfur, etc. The encapsulated fertilizer **10** can also be of different physical properties, such as different sizes and shapes, different dissolution rates, different hardness, etc. Such properties are described below. Generally, the encapsulated fertilizer **10** does not agglomerate, such that the encapsulated fertilizer **10** is pourable for ease of use, i.e., is free flowing, and thereby does not clog apparatuses used to supply and/or spread the encapsulated fertilizer **10**.

[0073] The encapsulated fertilizer **10** can be of various hardness (or “crush” strength) depending on several variables including the specific components and amounts thereof utilized, size and shape of the core particle **12**, size and shape of the encapsulated fertilizer **10**, and thickness of the layer **16**. Typically, the encapsulated fertilizer **10** has an excellent hardness relative to prior art encapsulated fertilizers. Hardness of the encapsulated fertilizer **10** provides for longer life of the encapsulated fertilizer **10**, such that the encapsulated fertilizer **10** has excellent handling and storage/shelf life. Specifically, the layer **16** provides for increased resiliency of the encapsulated fertilizer **10**, relative to conventional encapsulated fertilizers. Typically, the encapsulated fertilizer **10** has a shelf life of at least about 1, at least about 5, or at least about 10, months. The encapsulated fertilizer **10** can have a near infinite shelf life, more typically 20 or more years, based in part upon exposure (or lack thereof) to moisture.

[0074] The encapsulated fertilizer **10** is also useful for providing the core particle **12**, e.g. fertilizer, in a time-released manner. Time release, i.e., time required for the layer **16** to be perforated such that the core particle **12** can be released to the surrounding environment, depends on several variables including the specific components and amounts thereof utilized, size and shape of the core particle **12**, size and shape of the encapsulated fertilizer **10**, and thickness of the layer **16**. Such a determination can be easily tested by placing the encapsulated fertilizer **10** in a beaker full of water, such that the encapsulated fertilizer **10** is completely covered by the water. Perforations of the layer **16** can be evaluated upon visual inspection of the encapsulated fertilizer **10**, and/or by observing changes in clarity of the water. Perforations can range from pin-holing in or sloughing off of the layer **16**. Time release only applies if the layer **16** fully envelops the core particle **12**; otherwise, the core particle **12** is already at least partially exposed to the surrounding environment.

[0075] Time release of the encapsulated fertilizer **10** is also associated with dissolution rate of the encapsulated fertilizer **10**, as introduced above. Dissolution rate is generally a measurement of how much of the core particle **12** is dispersed into the surrounding environment. For example, the amount of fertilizer released to the surrounding soil over time once the encapsulated fertilizer **10** is exposed to moisture, e.g. from rain or irrigation. The encapsulated fertilizer **10** has excellent dissolution rates relative to prior art encapsulated fertilizers. Generally, the encapsulated fertilizer **10** of this disclosure has a uniform dissolution rate, i.e., a near steady-state release of the core particle **12** over a period of time. Alternatively, or in addition to, the encapsulated fertilizer **10** of this disclosure has a time-delayed release, e.g. the core particle **12** will begin to dissolve and therefore release some period of time after the encapsulated fertilizer **10** is first exposed to a sufficient amount of moisture to permeate the layer **16**.

[0076] One way in which such dissolution works is by the layer **16** becoming permeated such that that water can enter into the encapsulated fertilizer **10**. After entry of water, the water can interact with the core particle **12**. For example, if the core particle **12** comprises a nitrogen based fertilizer, after water migrates through the layer **16**, the nitrogen based fertilizer dissolves into solution inside the encapsulated fertilizer **10**, and the solution then migrates out through the layer **16** into surrounding soil to deliver the nitrogen. When the core particle **12** is the fertilizer, water and/or

other liquids generally cannot permeate the layer **16** too quickly, thus preventing the water and/or other liquids from rapidly dissolving the core particle **12**, thus preventing phytotoxicity, and providing for more predictable and controlled dissolution rates.

[0077] The following examples, illustrating the method and encapsulated fertilizer of this disclosure, are intended to illustrate and not to limit the present invention.

#### EXAMPLES

[0078] Encapsulated fertilizer is prepared in a lab-scale tumbler (or mixer). Core particles, specifically urea fertilizer spheres, are added to the tumbler. The tumbler is closed. The tumbler is heated to establish ambient moisture therein. Specifically, the core particles are heated to about 60 °C for 90 minutes. The core particles have a moisture content of from about 5 to about 10 wt.% prior to heating. Moisture content can be determined via a moisture balance.

[0079] An isocyanate component is heated to about 35 °C. The isocyanate component is a liquid modified MDI, specifically an isocyanate-terminated prepolymer comprising the reaction product of excess 4,4'-MDI and at least one polyol. The isocyanate component has an NCO content of 23 wt.%, a viscosity of 700 cps at 25 °C, and is commercially available from BASF Corporation.

[0080] The tumbler is opened and the isocyanate component is sprayed into the tumbler and onto the core particles. The tumbler is closed and rotated to ensure complete contact between the core particles and the isocyanate component. Complete contact results in a layer disposed about the core particles. The layer is a polyurea/polyurethane layer. Rotation continues to minimize agglomeration resulting in a free flowing group of encapsulated fertilizers, i.e., encapsulated urea spheres. Overall, the layer is applied at 3 wt% based on total core particle weight.

[0081] Referring to Figure 5, dissolution rates of various encapsulated fertilizers can be appreciated. "1 Component" encapsulated fertilizer (i.e., invention example) is the example described above having a 3 wt.% layer. "2 Component System A" and "2 Component System B" encapsulated fertilizers (i.e., comparative examples) are each formed by separately applying isocyanate and isocyanate-reactive components to the core particles. Each of "A" and "B" has a 3 wt% layer including 2.5 wt.% urethane/urea and 0.5% wax. The comparative examples are generally formed according to the disclosure of U.S. Pat. No. 7,416,785 to Mente. As illustrated in the

chart, it is thought that dissolution rate of the invention example is generally more uniform over time relative to the comparative examples.

**[0082]** It is to be understood that the appended claims are not limited to express any particular compounds, compositions, or methods described in the detailed description, which may vary between particular embodiments which fall within the scope of the appended claims. With respect to any Markush groups relied upon herein for describing particular features or aspects of various embodiments, it is to be appreciated that different, special, and/or unexpected results may be obtained from each member of the respective Markush group independent from all other Markush members. Each member of a Markush group may be relied upon individually and or in combination and provides adequate support for specific embodiments within the scope of the appended claims.

**[0083]** It is also to be understood that any ranges and subranges relied upon in describing various embodiments of the present invention independently and collectively fall within the scope of the appended claims, and are understood to describe and contemplate all ranges including whole and/or fractional values therein, even if such values are not expressly written herein. One of skill in the art readily recognizes that the enumerated ranges and subranges sufficiently describe and enable various embodiments of the present invention, and such ranges and subranges may be further delineated into relevant halves, thirds, quarters, fifths, and so on. As just one example, a range “of from 0.1 to 0.9” may be further delineated into a lower third, i.e., from 0.1 to 0.3, a middle third, i.e., from 0.4 to 0.6, and an upper third, i.e., from 0.7 to 0.9, which individually and collectively are within the scope of the appended claims, and may be relied upon individually and/or collectively and provide adequate support for specific embodiments within the scope of the appended claims. In addition, with respect to the language which defines or modifies a range, such as “at least,” “greater than,” “less than,” “no more than,” and the like, it is to be understood that such language includes subranges and/or an upper or lower limit. As another example, a range of “at least 10” inherently includes a subrange of from at least 10 to 35, a subrange of from at least 10 to 25, a subrange of from 25 to 35, and so on, and each subrange may be relied upon individually and/or collectively and provides adequate support for specific embodiments within the scope of the appended claims. Finally, an individual number within a disclosed range may be relied upon and

provides adequate support for specific embodiments within the scope of the appended claims. For example, a range “of from 1 to 9” includes various individual integers, such as 3, as well as individual numbers including a decimal point (or fraction), such as 4.1, which may be relied upon and provide adequate support for specific embodiments within the scope of the appended claims.

**[0084]** The present invention has been described herein in an illustrative manner, and it is to be understood that the terminology which has been used is intended to be in the nature of words of description rather than of limitation. Many modifications and variations of the present invention are possible in light of the above teachings. The present invention may be practiced otherwise than as specifically described within the scope of the appended claims. The subject matter of all combinations of independent and dependent claims, both single and multiple dependent, is herein expressly contemplated.

## CLAIMS

What is claimed is:

1. A method of forming an encapsulated fertilizer, said method comprising the steps of:

providing a core particle comprising a fertilizer and having an outer surface;

applying an isocyanate component having isocyanate functional groups directly to the outer surface of the core particle; and

reacting the isocyanate component with ambient moisture to form a layer disposed about the core particle to form the encapsulated fertilizer;

wherein the layer comprises polyurea linkages, and optionally, polyurethane linkages.

2. The method as set forth in claim 1, wherein no isocyanate-reactive component is present on the outer surface of the core particle.

3. The method as set forth in claim 1 or 2, wherein the core particle is at least partially coated with a compound other than fertilizer and inert with respect to the isocyanate component prior to the step of applying.

4. The method as set forth in any one of the preceding claims, wherein the isocyanate component comprises an isocyanate-terminated prepolymer, with the isocyanate-terminated prepolymer comprising polyurethane linkages such that the layer is further defined as a polyurea/polyurethane layer.

5. The method as set forth in claim 4, wherein the isocyanate-terminated prepolymer:

i) comprises the reaction product of a diphenylmethane diisocyanate (MDI) and a polyol having at least two hydroxyl functional groups;

ii) is present in an amount of from about 10 to about 90, alternatively from about 25 to about 75 parts by weight, alternatively of from about 40 to about 60 parts by weight, based on 100 parts by weight of the isocyanate component; or

iii) both i) and ii).

6. The method as set forth in any one of the preceding claims, wherein the isocyanate component:

i) is selected from the group of polymeric diphenylmethane diisocyanates (PMDIs), diphenylmethane diisocyanates (MDIs), and combinations thereof;



- ii) has an NCO content of from about 1 to about 50 wt.%, alternatively about 1 to about 40 wt.%; or
  - iii) both i) and ii).
7. The method as set forth in any one of the preceding claims, wherein the core particle has a moisture content of at least about 1 wt.%, alternatively from about 1 to about 20 wt.%, prior to the step of applying.
8. The method as set forth in claim 7, further comprising the step of heating the core particle to establish at least a portion of the ambient moisture, alternatively a majority of the ambient moisture, for reaction with the isocyanate component.
9. The method as set forth in claim 8, wherein the core particle is heated to a temperature of at least about 0 °C, alternatively to a temperature of from about 0 to about 200 °C.
10. The method as set forth in any one of the preceding claims, wherein the step of applying is further defined as spraying.
11. The method as set forth in any one of the preceding claims, further comprising the step of heating the isocyanate component.
12. The method as set forth in any one of the preceding claims, wherein the outer surface of the core particle comprises surface moisture, and wherein the step of reacting is further defined as reacting the isocyanate component with ambient moisture and the surface moisture to form the encapsulated fertilizer.
13. The method as set forth in any one of claims 1 through 11, wherein the outer surface of the core particle comprises amine functional groups, and wherein the step of reacting is further defined as reacting the isocyanate component with ambient moisture and the amine functional groups to form the encapsulated fertilizer.
14. The method as set forth in any one of the claims 1 through 11, wherein the outer surface of the core particle comprises surface moisture and amine functional groups, and wherein the step of reacting is further defined as reacting the isocyanate component with ambient moisture, the surface moisture, and the amine functional groups to form the encapsulated fertilizer.
15. The method as set forth in any one of the preceding claims, wherein the fertilizer comprises urea.
16. The method as set forth in any one of the preceding claims, wherein the layer:
- i) is in direct contact with the core particle; and/or

- ii) is substantially free of sulfur; and/or
- iii) is formed in the presence of a catalyst; and/or
- iv) has an average thickness of at least about 5, alternatively of from about 15 to about 300, alternatively of from about 50 to about 200, microns.

17. An encapsulated fertilizer formed according to the method as set forth in any one of the preceding claims.

18. An encapsulated fertilizer comprising:

a core particle comprising a fertilizer and having an outer surface; and

a layer disposed about said core particle and comprising the reaction product of

an isocyanate component having isocyanate functional groups, and  
ambient moisture;

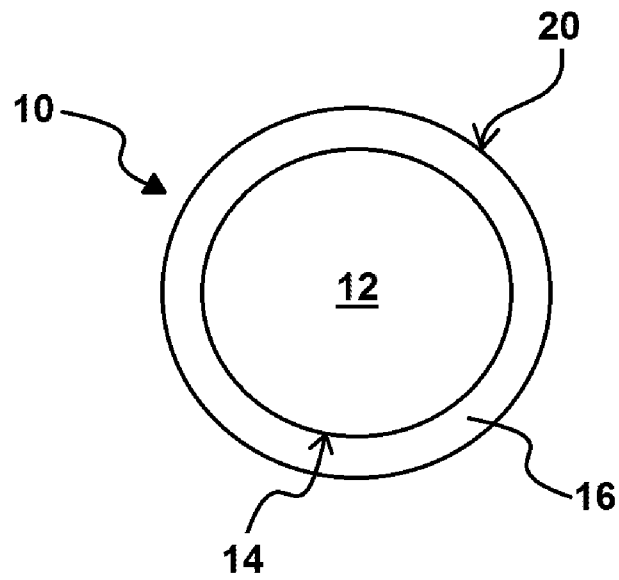
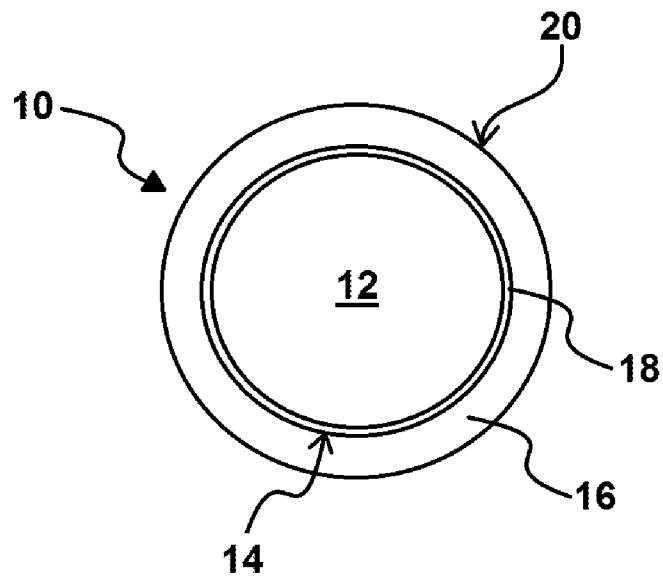
said encapsulated fertilizer formed according to a method comprising the steps of  
providing said core particle,

applying said isocyanate component directly to said outer surface of said core  
particle, and

reacting said isocyanate component with ambient moisture to form said layer  
disposed about said core particle to form said encapsulated fertilizer;  
wherein said layer comprises polyurea linkages, and optionally, polyurethane  
linkages.

19. The encapsulated fertilizer as set forth in claim 18, wherein said isocyanate  
component comprises an isocyanate-terminated prepolymer, with said isocyanate-  
terminated prepolymer comprising polyurethane linkages such that said layer is  
further defined as a polyurea/polyurethane layer.

20. The encapsulated fertilizer as set forth in claim 18 or 19, formed according to  
the method as set forth in any one of claims 2 through 16.

**FIG - 1****FIG - 2**

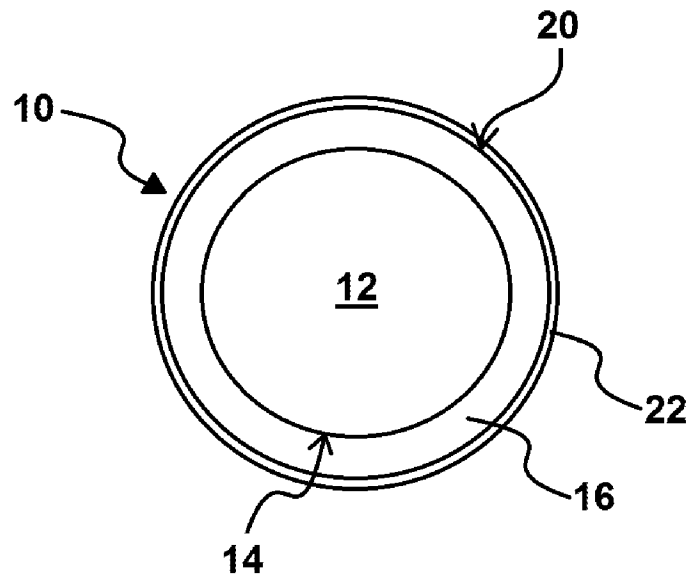
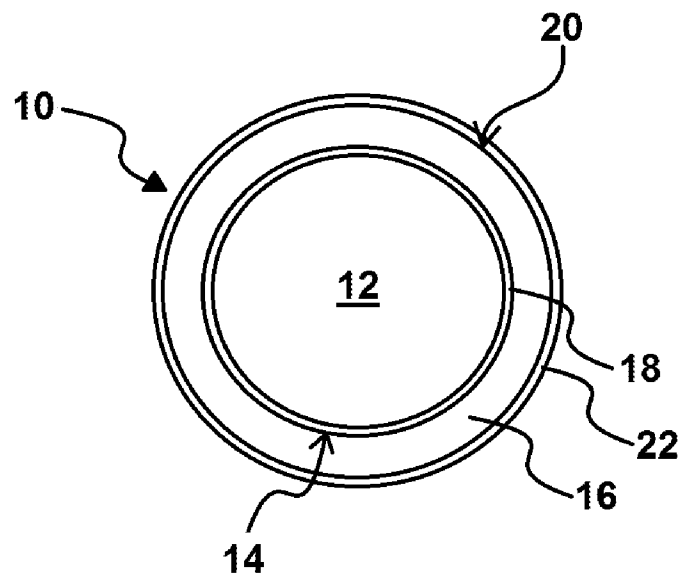
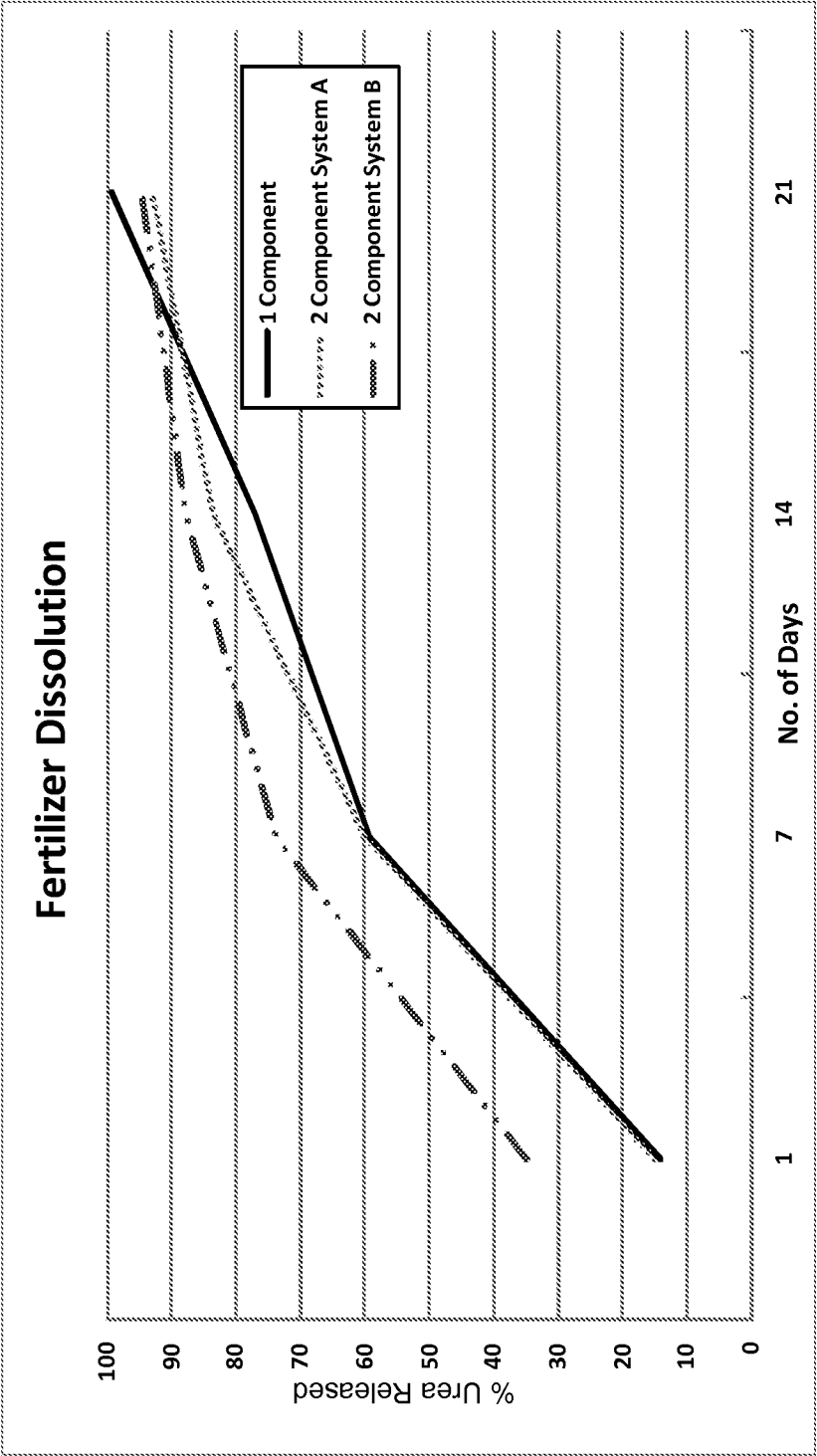
**FIG - 3****FIG - 4**

FIG - 5



## INTERNATIONAL SEARCH REPORT

International application No  
PCT/US2015/027725

## A. CLASSIFICATION OF SUBJECT MATTER

INV. C05G3/00 C05C9/00 C05C1/02 C05C11/00  
ADD.

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)

C05G C05C

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)

EPO-Internal, WPI Data, BIOSIS, COMPENDEX

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category* | Citation of document, with indication, where appropriate, of the relevant passages                                                                                                                                                                  | Relevant to claim No. |
|-----------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------|
| X         | WO 98/29359 A1 (BAYER AG [US])<br>9 July 1998 (1998-07-09)<br>page 5, line 21 - page 6, line 18<br>page 17, line 3 - line 21<br>claims 8-11<br>page 7, line 28 - page 8, line 5<br>page 7, line 11 - line 27<br>page 15, line 12 - line 15<br>----- | 1-20                  |
| X         | EP 0 974 568 A2 (BAYER AG [US])<br>26 January 2000 (2000-01-26)<br>paragraph [0001]<br>paragraph [0017]<br>paragraph [0020]<br>paragraph [0027]<br>paragraph [0031]<br>claims<br>-----<br>-/-                                                       | 1-20                  |



Further documents are listed in the continuation of Box C.



See patent family annex.

\* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

29 June 2015

Date of mailing of the international search report

07/07/2015

Name and mailing address of the ISA/

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Authorized officer

Rodriguez Fontao, M

## INTERNATIONAL SEARCH REPORT

International application No

PCT/US2015/027725

| C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT |                                                                                                                                                                                                                                                                  |                       |
|------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------|
| Category*                                            | Citation of document, with indication, where appropriate, of the relevant passages                                                                                                                                                                               | Relevant to claim No. |
| X                                                    | EP 0 974 569 A2 (BAYER AG [US])<br>26 January 2000 (2000-01-26)<br>paragraph [0024]<br>paragraph [0027] - paragraph [0034]<br>paragraph [0041]<br>paragraph [0048]<br>claims<br>paragraph [0044]<br>-----                                                        | 1-20                  |
| X                                                    | EP 0 974 609 A1 (BAYER AG [US])<br>26 January 2000 (2000-01-26)<br>paragraph [0027] - paragraph [0028]<br>paragraph [0022] - paragraph [0023]<br>paragraph [0036]<br>claims<br>-----                                                                             | 1-20                  |
| X                                                    | WO 91/10362 A1 (DU PONT [US])<br>25 July 1991 (1991-07-25)<br>claims<br>page 2, line 23 - line 31<br>page 5, line 33 - line 35<br>page 7, line 36 - page 8, line 2<br>page 9, line 1 - line 8<br>page 9, line 22 - line 32<br>page 10, line 1 - line 15<br>----- | 1-20                  |

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2015/027725

| Patent document<br>cited in search report | Publication<br>date | Patent family<br>member(s) | Publication<br>date     |
|-------------------------------------------|---------------------|----------------------------|-------------------------|
| WO 9829359                                | A1                  | 09-07-1998                 | AU 5605198 A 31-07-1998 |
|                                           |                     | CA 2275887 A1 09-07-1998   |                         |
|                                           |                     | DE 69713147 D1 11-07-2002  |                         |
|                                           |                     | DE 69713147 T2 14-11-2002  |                         |
|                                           |                     | EP 0948474 A1 13-10-1999   |                         |
|                                           |                     | ES 2178031 T3 16-12-2002   |                         |
|                                           |                     | IL 130493 A 23-12-2001     |                         |
|                                           |                     | US 5851261 A 22-12-1998    |                         |
|                                           |                     | WO 9829359 A1 09-07-1998   |                         |
| EP 0974568                                | A2                  | 26-01-2000                 | AT 293585 T 15-05-2005  |
|                                           |                     | CA 2277513 A1 23-01-2000   |                         |
|                                           |                     | DE 69924791 D1 25-05-2005  |                         |
|                                           |                     | DE 69924791 T2 09-03-2006  |                         |
|                                           |                     | EP 0974568 A2 26-01-2000   |                         |
|                                           |                     | ES 2241213 T3 16-10-2005   |                         |
|                                           |                     | IL 130988 A 13-09-2001     |                         |
|                                           |                     | US 6165550 A 26-12-2000    |                         |
| EP 0974569                                | A2                  | 26-01-2000                 | AT 294768 T 15-05-2005  |
|                                           |                     | CA 2273768 A1 23-01-2000   |                         |
|                                           |                     | DE 69925074 D1 09-06-2005  |                         |
|                                           |                     | DE 69925074 T2 19-01-2006  |                         |
|                                           |                     | EP 0974569 A2 26-01-2000   |                         |
|                                           |                     | ES 2242332 T3 01-11-2005   |                         |
|                                           |                     | IL 130987 A 23-12-2001     |                         |
|                                           |                     | US 6001147 A 14-12-1999    |                         |
| EP 0974609                                | A1                  | 26-01-2000                 | AT 219113 T 15-06-2002  |
|                                           |                     | CA 2274200 A1 23-01-2000   |                         |
|                                           |                     | DE 69901761 D1 18-07-2002  |                         |
|                                           |                     | DE 69901761 T2 21-11-2002  |                         |
|                                           |                     | EP 0974609 A1 26-01-2000   |                         |
|                                           |                     | ES 2178322 T3 16-12-2002   |                         |
|                                           |                     | IL 130986 A 31-08-2005     |                         |
|                                           |                     | US 6152981 A 28-11-2000    |                         |
| WO 9110362                                | A1                  | 25-07-1991                 | AU 7175091 A 05-08-1991 |
|                                           |                     | CA 2073689 A1 13-07-1991   |                         |
|                                           |                     | CS 9100058 A3 17-06-1992   |                         |
|                                           |                     | EP 0513027 A1 19-11-1992   |                         |
|                                           |                     | JP H05503697 A 17-06-1993  |                         |
|                                           |                     | NZ 236736 A 26-03-1993     |                         |
|                                           |                     | PT 96469 A 15-10-1991      |                         |
|                                           |                     | WO 9110362 A1 25-07-1991   |                         |
|                                           |                     | ZA 9100237 A 30-09-1992    |                         |