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DESCRIPTION

FIELD OF THE INVENTION

[0001] The present invention relates to stable, durable granules with active agents. Specifically, the invention relates to thermostable, durable granules with active agents, the granules being particularly suitable for inclusion in steam treatment processes, including pelleting and tableting processes and steam processing of feed, without appreciable loss of active agent activity. The stable, durable granules have dissolution profiles suitable to release the active agent to provide efficacy for its intended purpose. The activity of the active agents is retained following storage in unpelleted mixtures and steam treatment.

BACKGROUND OF THE INVENTION

[0002] The use of active agents, such as enzymes, in animal feed is common. Enzymes are known to improve digestibility of feed, reduce antinutritional factors in feed, and improve animal productivity. It is known in the industry that acidic and basic feed ingredients as well as particular ingredients of animal feeds, including, but not limited to trace minerals, organic or inorganic acids or bases, reducing sugars, and hygroscopic substances, particularly choline chloride and sodium chloride, have an adverse effect upon active agents such as other vitamins, proteins, antimicrobials, prebiotics, probiotics and enzymes; and, it is also known that some feed production processes are detrimental to active agents.

[0003] A problem exists in the industry to provide protective formulations to make the active agents suitable for storage in unpelleted animal feed mixtures, such as base mixes and premixes, which may be acidic or basic and contain the ingredients that have an adverse effect upon the stability of the active agents. One mechanism for the adverse effect is said to be oxidation-reduction (redox) reactions that occur between oxidizing and reducing compounds in the premixes in the presence of water. A study reported in BASF Technical Bulletin NU0013 reports that two commercially available enzyme containing granules retained, respectively, 86% and 81% activity after storage for 3 weeks in a feed premix, and 55% and 33% activity, respectively, after storage in the feed premix for 6 weeks. Currently some manufacturers of enzymes for the feed industry recommend that enzymes be protected with barrier packaging if they are to be stored in premixes, or that they be stored separately from the premixes, or that they be stored in premixes for only a short period of time.

[0004] Additionally, many active agents used in food and feed are heat labile. Thermal stability of enzymes and their ability to survive heat processing steps in the manufacture of animal feed is a problem in the industry, particularly in the production of animal feed pellets. When compared with dry feed mixes, feed pellets have properties that are favored by the industry, such as improved feed quality, decreased pathogens, lower dust levels during manufacture, good handling, and more uniform ingredient dosing. Preferred industry pelleting processes utilize steam injection, in a process known as conditioning, which adds moisture and elevates the temperature prior to the pelleting step, which then forces the steam heated feed ingredients, or conditioned mash, through a die. The pelleting process temperatures may be from about 70°C to 95°C, or higher.

[0005] Enzymes are important feed ingredients and must be able to withstand increasingly higher processing temperatures used in pelleting processes, particularly those processes that employ expanders, while continuing to deliver *in vivo* efficacy.

[0006] Because of the steam, temperatures and compression forces used in pelleting processes, the stability of enzymes and other active agents is a problem that is illustrated by the fact that feed enzymes are often provided to the industry as stabilized liquid products that are added to feed pellets after the pelleting process to avoid enzyme inactivation. Homogeneous dosing is a problem when the enzyme is applied post pelleting, for instance, by spraying the enzyme onto the pellets, and the cost of the equipment to add enzyme post-pelleting is high. Alternatively, liquid enzyme formulations, or dry mix enzyme formulations are added to the mixer prior to pelleting. In certain instances, higher levels of enzymes than needed may be added in order to compensate for losses during pelleting.

[0007] Tablet forming processes also utilize compression forces and may or may not employ heat. Tablets are used in the household care industries, for instance, in laundry, dish and surface cleaning applications.

[0008] There is a need in the food, feed, and household care industries for stable, durable granules with active ingredients to serve as components in formulations that are subjected to steam treatment, for instance, pelleting and tableting processes,) without appreciable loss of active agent activity and with dissolution profiles suitable to release the active ingredients to provide efficacy for their intended purpose. There also is a need for stable, durable granules with active agents that retain their activity

when used as ingredients in animal feed formulations, such as unpelleted mixtures, that contain ingredients that adversely affect active agents.

[0009] WO 2006/034710 discloses salt-coated enzyme granules for use in animal feed.

SUMMARY OF THE INVENTION

[0010] The present invention relates to a granule for animal feed comprising:

a core;

an active agent;

and at least one coating comprising a moisture hydrating material that is an inorganic salt which is capable of taking up aqueous liquids by absorbing free water or by taking up water as crystalline water of hydration and which moisture hydrating material is at least 25% w/w of the granule,

wherein the water activity of the granule is less than 0.5,

and wherein the active agent of the granule retains at least 50% activity after conditions selected from one or more of:

1. a) a feed pelleting process,
2. b) a steam-heated feed pretreatment process,
3. c) storage,
4. d) storage as an ingredient in an unpelleted mixture, and
5. e) storage as an ingredient in a feed base mix or a feed premix comprising at least one compound selected from trace minerals, organic acids, reducing sugars, vitamins, choline chloride, and compounds which result in an acidic or basic feed base mix or feed premix.

[0011] In a process for producing a granule comprising an active agent for feed, the process comprises: preparing stable granules having a core, at least one active agent, and at least one coating; mixing the stable granules together with one or more of a) a diluent, b) a base mix, c) a premix, and d) a feed mixture for pelleting.

[0012] In another embodiment of the present invention, a process is provided for making a stable granule comprising enzyme for storage in a feed premix comprising choline chloride, the process comprising: providing a core material and enzyme, the enzyme distributed throughout the core material or layered over the core material; applying to the core material and enzyme a moisture hydrating material to form a layer that is at least 25% w/w of the stable granule; coating the layer with a moisture barrier material to form a coating that is at least 2% w/w of the granule, the applying and coating under conditions selected so that a water activity of the stable granule is less than 0.5.

[0013] Stable, durable granules of the above embodiments retain at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, and at least 95% active agent activity under conditions of steam heated pelleting or pretreatment processes that raise the temperature of the pelleted material to as high as 85°C to about 95°C for up to about several minutes. The stable, durable granules of the present invention are particularly suitable as ingredients in animal feed pellets.

[0014] In embodiments of the present invention, stable durable granules include a moisture hydrating material that constitutes at least about 55% w/w of the granule.

[0015] In other embodiments of the present invention, a stable durable granule has a moisture hydrating coating that comprises at least about 25%, about 34%, about 35%, about 44%, about 54%, about 55%, and about 60% w/w of the granule and a moisture barrier coating that comprises at least about 2% to about 40% w/w, about 2% to about 10%, about 2% to about 7%, and about 5% to about 15% of the granule.

[0016] In embodiments of the present invention, a stable durable granule comprises a silicate or clay core and an inherently thermal stable active agent within a matrix; an optional inorganic salt moisture hydrating protective coating; and an optional moisture barrier coating. In this embodiment, the optional coating layers constitute about 0% to about 15% w/w of the granule.

[0017] The present invention further includes methods for manufacturing the stable, durable granules as well as feed pellets and pet food, containing such stable, durable granules.

DETAILED DESCRIPTION OF THE INVENTION

[0018] The present invention provides stable, durable granules with active agents that withstand both high temperatures and compression forces when added to formulations subjected to, for instance, heated steam pretreatments, feed pelleting processes and tableting processes, while maintaining dissolution profiles suitable to release active agents that are able to provide efficacy for their intended purpose.

[0019] A first aspect of the present invention surprisingly provides stable durable granules with feed active agents that withstand steam pretreatments and feed pelleting steam-heated process temperatures and compression forces while maintaining dissolution profiles that release the feed active agents to provide intended efficacy. In this embodiment, granule components preferably are approved for use in feed.

[0020] A second aspect of the present invention provides stable, durable granules with enzymes that withstand steam-heated feed pretreatment and pelleting process temperatures and compression forces while maintaining dissolution profiles that release the enzyme to provide *in vivo* bioavailability efficacy. In this embodiment, granule components are edible by animals, and preferably are also digestible.

[0021] A third aspect of the present invention provides stable, durable granules with active agents for animal feed unpelleted mixtures, for example, premixes. The stable, durable granules retain efficacy when the unpelleted mixture is heat and steam pretreated prior to feeding to animals, or after the unpelleted mixture is pelleted. In this aspect of the invention, surprisingly, the active agents maintain activity when stored in unpelleted mixtures that contain ingredients that are detrimental to enzyme stability. In this aspect of the invention, without wishing to be bound by any particular theory, it is believed that a moisture hydrating material in the granules acts in combination with a moisture barrier material in the granules to protect the active ingredient from the detrimental ingredients in unpelleted mixtures. The moisture hydrating material retards or reduces the rate or extent of water migration into the area of the active agent, and the moisture barrier material excludes water. The combination of the moisture hydrating material and the moisture barrier material provides mechanical stability to further protect the active agent in the event that a layer of a moisture barrier material is damaged. Additionally, in some embodiments of the present invention, moisture barrier materials are used that oxidize only under extreme conditions, thereby in combination with moisture hydrating materials, the granules are chemically stable because it is believed that redox reactions are reduced during storage of the granules in unpelleted materials.

[0022] The stable, durable granules of the present invention are spherical or near spherical granules, although other shapes such as discs, ovals, cylindrical and oblong shapes may be used if desired. The granules have one or more protective layers surrounding the active agent.

[0023] The granules may be mixed together with dry ingredients, such as feed or household care formulations or unpelleted feed mixtures, for examples, pre-mix formulations, prior to use in a pelleting or tableting process, or used in dry feed mixtures and mash that are not pelleted. The granules are particularly suitable for use in feed pelleting manufacturing processes, and are also suitable for food, including pet food, and household care tablet manufacturing processes.

[0024] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention pertains. As used in the specification and claims, the singular "a", "an" and "the" include the plural references unless the context clearly dictates otherwise. For example, the term granule may include a plurality of granules.

[0025] For purposes of this disclosure the "active agent" may be any material that is to be added to a stable, durable granule. The active agent may be a biologically viable material, a food or feed ingredient, an antimicrobial agent, an antibiotic replacement agent, a prebiotic, a probiotic, an agrochemical ingredient, such as a pesticide, fertilizer or herbicide; a pharmaceutical ingredient or a household care active ingredient, or combinations thereof. In a preferred embodiment, the active ingredient is a protein, enzyme, peptide, polypeptide, amino acid; carbohydrate, lipid or oil, vitamin, co-vitamin, or hormone, or combinations thereof. In another embodiment, the active ingredient is an enzyme, bleach, bleach activator, perfume, or other biologically active ingredient. Inherently thermostable active agents are encompassed by the present invention and exhibit enhanced thermostability in the granules of the present invention. Alternatively, less coating materials may be used when providing granules with inherently

thermostable active agents, and the protective coating material amounts provided herein were selected primarily for active agents that were not inherently thermostable.

[0026] Most preferred active ingredients for food and feed applications are enzymes, peptides and polypeptides, amino acids, antimicrobials, gut health promoting agents, vitamins, and combinations thereof.

[0027] Any enzyme may be used, and a nonlimiting list of enzymes include phytases, xylanases, β -glucanases, phosphatases, proteases, amylases (alpha or beta or glucoamylases) cellulases, lipases, cutinases, oxidases, transferases, reductases, hemicellulases, mannanases, esterases, isomerases, pectinases, lactases, peroxidases, laccases, other redox enzymes and mixtures thereof.

[0028] Particularly preferred enzymes include a xylanase from *Trichoderma reesei* and a variant xylanase from *Trichoderma reesei*, both available from Danisco A/S, Denmark and/or Genencor International, Inc., Palo Alto, California, or the inherently thermostable xylanase described in EP1222256B1, as well as other xylanases from *Aspergillus niger*, *Aspergillus kawachii*, *Aspergillus tubigenensis*, *Bacillus circulans*, *Bacillus pumilus*, *Bacillus subtilis*, *Neocallimastix patriciarum*, *Penicillium species*, *Streptomyces lividans*, *Streptomyces thermoviolaceus*, *Thermomonospora fusca*, *Trichoderma harzianum*, *Trichoderma reesei*, *Trichoderma viride*; examples of phytases are Finase L[®], a phytase from *Aspergillus* sp., available from AB Enzymes, Darmstadt, Germany; Phyzyme[™] XP, a phytase from *E. Coli*, available from Danisco, Copenhagen, Denmark, and other phytases from, for example, the following species: *Trichoderma*, *Penicillium*, *Fusarium*, *Buttiauxella*, *Citrobacter*, *Enterobacter*, *Penicillium*, *Humicola*, *Bacillus*, and *Peniophora*.

[0029] An example of a cellulase is Multifect[®] BGL, a cellulase (beta glucanase), available from Danisco A/S, Denmark, and other cellulases from species such as *Aspergillus*, *Trichoderma*, *Penicillium*, *Humicola*, *Bacillus*, *Cellulomonas*, *Penicillium*, *Thermomonospora*, *Clastridium*, and *Hypocrea*. The cellulases and endoglucanases described in US20060193897A1 also may be used. Amylases may be, for example, from species such as *Aspergillus*, *Trichoderma*, *Penicillium*, *Bacillus*, for instance, *B. subtilis*, *B. stearothermophilus*, *B. lentus*, *B. licheniformis*, *B. coagulans*, and *B. amyloliquefaciens*. Suitable fungal amylases are derived from *Aspergillus*, such as *A. oryzae* and *A. niger*. Proteases may be from *Bacillus amyloliquefaciens*, *Bacillus lentus*, *Bacillus subtilis*, *Bacillus licheniformis*, and *Aspergillus* and *Trichoderma* species.

[0030] Phytases, xylanases, phosphatases, proteases, amylases, esterases, redox enzymes, lipases, transferases, cellulases, and β -glucanases are enzymes frequently used for inclusion in animal feed. Enzymes suitable for inclusion into tablets for household care applications are similar, particularly proteases, amylases, lipases, hemicellulases, redox enzymes, peroxidases, transferases, and cellulases.

[0031] In particularly preferred aspects of the invention, the enzymes are selected from phytases, xylanases, beta glucanases, amylases, proteases, lipases, esterases, and mixtures thereof. In one embodiment of the present invention, two enzymes are provided in the granule, a xylanase and a beta-glucanase. The enzymes may be mixed together or applied to the granule separately. In another embodiment, three enzymes are provided in the granule, namely beta-glucanase, xylanase and phytase.

[0032] The above enzyme lists are examples only and are not meant to be exclusive. Any enzyme may be used in the durable granules of the present invention, including wild type, recombinant and variant enzymes of bacterial, fungal, yeast, plant, insect and animal sources, and acid, neutral or alkaline enzymes.

[0033] It will be recognized by those skilled in the art that the amount of enzyme used will depend, at least in part, upon the type and property of the selected enzyme and the intended use.

[0034] The durable granules of the invention include between about 0.0005 to about 20% on a dry weight basis of the enzyme component of the granule. For instance, the weight percent of enzyme in embodiments of the invention comprises at least 0.0005 to about 15%, at least 0.001 to about 15%, at least 0.01 to about 10%, at least 0.1 to about 10%, at least 1.0 to about 10%, at least 1.0 to about 8%, at least 1.0 to about 5%, and at least 2.0 to at least 5% in the granule. Typical doses of 25 to 400 grams of the stable, durable enzyme granules per ton of feed will deliver about 0.0001 to about 80 grams of active enzyme protein per ton of feed, and the enzyme granules may be dosed as high as 5000 grams per ton of feed. Dosages for other active ingredients typically are 0.001 to about 400 grams/ton of the feed, or higher. Dosages may be even higher when the thermostable granules of the present invention are used as a component of animal feed unpelleted mixtures, for example, pre-mixes, which may contain several active agents and are added to feed compositions. For example, the dosage of the enzyme granules added to a premix may be about 0.2% to 10% of the premix, or about 0.1 % to about 3% of a base mix. In an exemplary embodiment, the activity level of phytase containing stable, durable granules stored in a premix is about 500 U/g, or higher. Premixes typically are added

to the diet at about 0.5% to about 2%, and base mixes are added at about 2% to about 6%.

[0035] The portion of the durable granule, without the protective coating(s) of the invention, and comprising the active ingredient, including any processing solids, binders and other ingredients therein, may comprise less than about 70%, less than about 60%, less than about 50%, less than about 40%, less than about 30%, and less than about 20% by weight of the granule, with, generally, the weight percent being about 25% to about 50% w/w, about 40% to about 60%, or about 50% to about 60%.

[0036] The coating layer(s) may comprise more than about 30%, more than about 40%, more than about 50%, more than about 60%, more than about 70%, and more than about 80% by weight of the granule, with, generally, weight percent being about 50% to about 75% w/w, 40% to about 60%, or about 40% to about 50%, depending upon the type and number of coating layers.

[0037] The durable granules of the present invention may be sized as desired and are between about 300 um to about 1000 um in diameter, about 300 um to about 900 um, about 400 um to about 800 um, and about 400 um to about 600 um.

[0038] "Compression forces" for purposes of this disclosure refers to generally axially applied forces that cause the atoms of a material to compact. Compression forces, as used herein, occur in pelleting and tableting processes and may include an element of flexural stress where the forces applied are not completely symmetrical with respect to the longitudinal axis of the material.

[0039] An "inherently thermostable enzyme or protein" refers to enzymes and proteins having a melting point that is above about 60°C to about 65°C, for example, the inherently thermostable enzyme utilized in Table 2 for granules 28, 29 and 30, has a melting point of 69°C at a pH of 5.5 and a melting point of 72° C at a pH of 8.0. A number of the thermolabile enzymes referred to throughout the specification had, for example, a melting point of less than 60°C at pH values of 5.5 and 8.0.

[0040] "Moisture barrier materials" refers to materials that exclude, prevent or substantially retard water uptake. These materials typically are hydrophobic or amphiphilic, provide insulation against water and do not inherently absorb and/or bind water and include, but are not limited to, film-forming materials. Examples of moisture barrier materials include polymers, proteins, lipids, fats and oils, fatty acids and gums. Examples of film forming moisture barrier materials are natural and modified polymers, such as gum arabic, whey, whey protein concentrate, PVA, including modified PVA and synthetic polymers such as , latex, HPMC, and acid-thinned hydroxypropyl starch, for example; PureCote™, oxidized starch, and modified starch. Non-film forming moisture barrier materials include, for instance, waxes, fats, oils and lipids, and lecithin. Selected moisture barrier materials that do not readily oxidize are, for example, latex polymer and polymers such as gum arabic.

[0041] "Moisture hydrating materials" refers to materials that take up aqueous liquids, such as water, by one several mechanisms. In a first mechanism, the materials absorb free water. In a second mechanism, the materials take up bound water that generally is present as crystalline waters of hydration. Accordingly, the materials may be provided as partially or fully hydrated materials or as non-hydrated materials that will absorb or bind aqueous liquids and retard or reduce the rate or extent of migration of such liquids to the active agent. In a third mechanism, moisture hydrating materials thermally insulate the active agent by retarding heat transfer to the active agent within the granule and by maintaining the active agent at a lower temperature than the temperature at the exterior surface of the granule. Moisture hydrating materials include carbohydrates and inorganic salts, including hydrated salts, such as magnesium sulfate, sodium sulfate, and ammonium sulfate; maltodextrin; sugars, for example, sucrose; and cornstarch.

[0042] In the present invention, the moisture hydrating materials are inorganic salts which, when added to the granule at greater than about 25% w/w, are anhydrous or contain relatively low amounts of bound or free water so that the water activity of the completed granule is less than 0.5. Without wishing to be bound by any particular theory, when the granule is subjected to steam-heating processes, the inorganic salt moisture hydrating material will begin to take up water from the steam-heated treatment, the water moving into the moisture hydrating material in a kinetic process over the short time period of the steam-treatment to prevent the water from penetrating into the area of the granule bearing the active agent. For purposes of this disclosure, "hydrated", "partially hydrated", and "non-hydrated" refer to a material's hydration potential when the granule is at equilibrium prior to being subjected to steam-heating. A "hydrated" material refers to a material that contains water in a free or bound form, or a combination thereof. Water may be added either during or after coating processes and the degree of hydration of the granule is a function of the granule materials and the temperature, humidity and drying conditions under which it is applied.

[0043] "Pellets" and "Pelleting" refer to solid rounded, spherical and cylindrical tablets or pellets and the processes for forming such solid shapes, particularly feed pellets and solid, extruded animal feed. Known feed pelleting manufacturing processes generally include mixing together feed ingredients for about 1 to about 5 minutes, transferring the mixture to a surge bin, conveying the mixture to a steam conditioner, optionally transferring the steam conditioned mixture to an expander, transferring the mixture to the pellet mill or extruder, and finally transferring the pellets into a pellet cooler. Fairfield, D. 1994. Chapter 10,

Pelleting Cost Center. In Feed Manufacturing Technology IV. (McElhiney, editor), American Feed Industry Association, Arlington, VA, pp. 110-139.

[0044] The steam conditioner treats the mixture for about 20 to about 90 seconds, and up to several minutes, at about 85°C to about 95°C. The amount of steam may vary in accordance with the amount of moisture and the initial temperature of the feed mix. About 4% to about 6% added steam has been reported in pelleting processes, and the amount is selected to produce less than about 18% moisture in the mash prior to pelleting, or up to about 28% moisture in mash intended for extrusion.

[0045] An optional expander process occurs for about 4 to about 10 seconds at a temperature range of about 100°C to about 140°C. The pellet mill portion of the manufacturing process typically operates for about 3 to about 5 seconds at a temperature of about 85°C to about 95°C.

[0046] A "stable" granule refers to a granule in which the activity of the active agent(s) is substantially maintained after inclusion as an ingredient in a formulation subjected to steam heated pretreatment processes, steam heated pelleting processes, steam heated tableting processes, after storage alone and storage as an ingredient in unpelleted and untableted mixtures. Stability includes thermostability, shelf-life or storage stability, mechanical stability, and/or chemical stability when the active agent containing granules are stored in unpelleted or untableted mixtures and/or subjected to steam-heated and pressurized pelleting processes and tableting processes. Mechanical stability refers to the physical robustness or structural integrity of granules, which structural integrity includes resistance to microbials, resistance to dust production, and resistance to release of ingredients that may result in the production of odors. Chemical stability refers to the substantial maintenance of activity when the granules are stored in unpelleted or untableted mixtures with ingredient(s) harmful to the active agent(s). Thermostability is defined further herein, and generally refers to maintenance of activity following exposure to temperatures up to about 85°C to 95°C when the stable, durable granules are an ingredient of pellets, tablets and unpelleted and untableted mixtures.

[0047] "Tableting" refers to processes for forming solid slugs or tablets by compressing a mixture of ingredients in a tablet press as described in EP 1 257 631 B1, which is hereby incorporated by reference.

[0048] Tablets may be made by direct compression tableting of mixtures of active agents, fillers/binders, lubricants, and any other optional ingredients. The active agent component is mixed thoroughly with the other tablet ingredients prior to entering the tablet machine. Ingredients are blended in any suitable mixing device, such as a twin shell blender or similar apparatus, or using any mixing method that results in blending of the tablet ingredients.

[0049] The mixtures are then compressed into tablets, using any tableting device, such as a tablet press (Stokes Model R-4, Warminster, PA). Tablet presses generally have upper and lower shape-corresponding punches, which fit into a die from above and below the die. Mixed tablet material is filled into the die cavity and at least one of the punches, typically the upper punch, enters the die cavity. Pressure is applied to both the upper and lower punches. The action of the upper and lower punches moving toward each other, applies pressure to the material between the punches, thus forming a tablet.

[0050] A wide variety of tablet shapes can be made. Tablet shape is determined by the tooling of the punches. Compaction forces vary, depending on the punch geometry, type of instrument, and formulation used. Alternatively, tablets may be made using dry or wet granulation procedures as described in US Pat. No. 6,852,336, which is hereby incorporated by reference herein in its entirety. The 336 patent states that dry granulation procedures may be utilized where one of the components has sufficient cohesive properties to be tableted. The method mixes the ingredients with a lubricant, if required. The wet granulation procedure described mixes the dry ingredient using a twin shell blender or double-cone blender under shear mixing conditions and then adds solutions of a binding agent to the mixed powders to obtain a granulation.

[0051] A "thermostable" granule refers to a granule having an active agent that retains at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, and at least 95% of its activity, measured by a conventional assay specific for the selected active agent, following inclusion in a formulation, and exposure to heat or to heat and steam, such that the formulation reaches temperatures up to about 85°C to about 95°C for up to several minutes. Those skilled in the art will recognize that thermostable granules in formulations exposed to temperatures less than 85°C are stable, and test results conducted at 80°C establish that this is indeed the case. Accordingly, it will be understood that up to 85°C to 95°C is meant to encompass temperatures less than 85°C, which are included in the scope of this invention.

[0052] In one embodiment, "thermostable" refers to an active agent that retains at least 80% of its activity measured by a conventional assay specific for the selected active ingredient following inclusion in a formulation, exposed to steam, such that the formulation reaches temperatures of about 85°C-95°C for about 30 seconds. In yet another embodiment, thermostable refers to

an active agent that retains at least 50% of its activity measured by a conventional method that is specific for the selected active ingredient following inclusion in a formulation, exposed to steam, such that the formulation reaches temperatures of about 85° C-95°C for about 30 seconds.

[0053] "Unpelleted mixtures" refers to premixes or precursors, base mixes, mash, and diluents. Premixes typically contain vitamins and trace minerals. Base mixes typically contain food and feed ingredients such as dicalcium phosphate, limestone, salt and a vitamin and mineral premix, but not grains and protein ingredients. Diluents include, but are not limited to grains (for example wheat middlings and rice bran) and clays, such as phyllosilicates (the magnesium silicate sepiolite, bentonite, kaolin, montmorillonite, hectorite, saponite, beidellite, attapulgite, and stevensite). Clays also function as carriers and fluidizing agent, or diluent, for feed premixes. Mash typically comprises a complete animal diet.

[0054] The stable granules of the present invention may be added to these unpelleted mixtures, and to mash mixtures, which may be subsequently treated with steam and/or steam pelleted or dried.

[0055] "Water activity", symbolized as a_w refers to the fractional relative humidity of an atmosphere in equilibrium with a solid or liquid phase material, i.e., the ratio of the partial pressure of water vapor to that present above pure water at the same temperature. In all phases between which water distribution has reached equilibrium, it is by definition equal. The term "relative humidity" is generally used to describe the water in the atmosphere or gas phase in equilibrium with the solid, and is expressed as a percentage, with 100% as the relative humidity of pure water in a closed system. Thus, for any water activity value, there is a corresponding relative humidity given by $\%RH=100 \cdot a_w$. Water activity can be readily measured by methods known in the art, typically by placing a sample of the material inside the temperature-controlled chamber of a water activity meter, such as the water Activity System Model D2100 available from Rotronic Instrument Corp. (Huntington, N.Y.), and allowing the measurement to reach equilibrium as indicated on the display. The water activity referred to herein is that of the granule itself after application of all coatings.

[0056] The water activity for the granules of the present invention is less than 0.5.

[0057] The stable, durable granule of the present invention is a granule that may be prepared by any process that results in the application of one or more protective coatings to a core. The active agent may be a component of the core of the granule, or may be coated over a core material, and for the purposes of this disclosure, a core refers to all components of the durable granule except any protective coating(s) applied to the core of the granule.

[0058] The core may be made by any process known in the art, such as granulation, extrusion, pan coating, spheronization, drum granulation, high-shear agglomeration, fluid-bed spray coating, crystallization, precipitation, and prill processes. Such processes are known in the art and are described in US Pat. No. 4,689,297 and US Pat. No. 5,324,649 (fluid bed processing); EP656058B1 and US Pat. No. 454332 (extrusion process); US Pat. No. 6,248,706 (granulation, high-shear); and EP804532B1 and US Pat. No. 6,534,466 (combination processes utilizing a fluid bed core and mixer coating), all disclosures of which are incorporated by reference in their entirety.

[0059] The protective coatings of the present invention may be applied as described in the granulation, extrusion, fluid-bed and prill processes noted above. Additionally, the protective coatings may be applied by casting methods, including spinning disk casting methods, as described in WO 03/000625, which is hereby incorporated by reference in its entirety.

[0060] Additionally, in one embodiment, the process may include a heat annealing step, which heats the granule to above the glass transition temperature (T_g) of a moisture barrier component and then gradually reduces the temperature to cause the barrier material to harden, or become glassy.

CORES

[0061] The core is the inner nucleus of the granule, and as stated above, may include the active ingredient or the active ingredient may be coated around a core material. Suitable cores for use in the present invention are preferably a hydratable or porous material (i.e., a material which is dispersible or soluble in water) that is a feed grade material. The core material should either disperse in water (disintegrate when hydrated) or solubilize in water by going into a true aqueous solution. Clays (for example, the phyllosilicates bentonite, kaolin, montmorillonite, hectorite, saponite, beidellite, attapulgite, and stevensite, silicates, such as sand (sodium silicate), nonpareils and agglomerated potato starch or flour, or other starch granule sources such as wheat and corn cobs are considered dispersible. Nonpareils are spherical particles consisting of a seed crystal that has been built

onto and rounded into a spherical shape by binding layers of powder and solute to the seed crystal in a rotating spherical container. Nonpareils are typically made from a combination of a sugar such as sucrose, and a powder such as Cornstarch. In one embodiment of the present invention the core is a sodium chloride or sodium sulfate crystals sometimes referred to as a seed, or other inorganic salt crystal. In another embodiment of the present invention, the core is a sucrose crystal.

[0062] Particles composed of inorganic salts and/or sugars and/or small organic molecules may be used as the cores of the present invention. Suitable water soluble ingredients for incorporation into cores include: inorganic salts such as sodium chloride, ammonium sulfate, sodium sulfate, magnesium sulfate, zinc sulfate; or urea, citric acid, sugars such as sucrose, lactose and the like.

[0063] Cores of the present invention may further comprise one or more of the following: active agents, feed or food grade polymers, fillers, plasticizers, fibrous materials, extenders and other compounds known to be used in cores. Suitable polymers include polyvinyl alcohol (PVA), polyethylene glycol, polyethylene oxide, polyvinyl pyrrolidone, and carbohydrate polymers (such as starch, amylose, amylopectin, alpha and beta-glucans, pectin, glycogen), including mixtures and derivatives thereof.

[0064] Cores used in the examples include sugar crystals, inorganic salt crystals, corn cob cores, clays and silicate.

[0065] Suitable fillers useful in the cores include inert materials used to add bulk and reduce cost, or used for the purpose of adjusting the intended enzyme activity in the finished granule. Examples of such fillers include, but are not limited to, water soluble agents such as salts, sugars and water dispersible agents such as clays, talc, silicates, cellulose and starches, and cellulose and starch derivatives.

[0066] Suitable plasticizers useful in the cores of the present invention are low molecular weight organic compounds and are highly specific to the polymer being plasticized. Examples include, but are not limited to, sugars (such as, glucose, fructose and sucrose), sugar alcohols (such as, sorbitol, xylitol and maltitol and other glycols), polar low molecular weight organic compounds, such as urea, or other known plasticizers such as water or feed grade plasticizers.

[0067] Suitable fibrous materials useful in the cores of the present invention include, but are not limited to: cellulose, and cellulose derivatives such as HPMC (hydroxypropyl-methyl cellulose), CMC (carboxy-methyl cellulose), HEC (hydroxyethyl cellulose).

[0068] In one embodiment, particularly for feed application, of the present invention, the core is a water-soluble or dispersible corn cob material or sugar or salt crystal. In another embodiment particularly suitable for household cleansing applications, the core is a water-soluble or dispersible sugar or salt crystal or a non pareil.

[0069] Those skilled in the art will recognize that, for feed and food applications, the cores (and any polymers, fillers, plasticizers, fibrous materials, and extenders), are acceptable for food and/or feed applications.

[0070] The core of the granules of the present invention, including any coating having active ingredients therein, and excluding the protective coatings described below, preferably comprises less than about 70%, less than about 60%, less than about 50%, less than about 40%, less than about 30%, and less than about 20% w/w of the durable granule.

ACTIVE AGENT

[0071] The active agent or agents, particularly the enzymes discussed above, is obtained from a fermentation broth and may be a whole broth, lysed broth, or clarified fermentation broth recovered from the fermentation process. The enzyme may be in liquid, slurry, dried, lyophilized, or crystalline form as obtained using recovery processes from fermentation broths. The enzyme(s) optionally are mixed together with plasticizers, such as carbohydrates, and polymers such as modified, or native starch to form a matrix. A non-limiting list of plasticizers includes the sugars glucose, fructose and sucrose, dextrans, PVA, and sugar alcohols, such as, sorbitol, xylitol and maltitol, or any of the plasticizers listed above. Modified starches include, but are not limited to cornstarch and acid-thinned hydroxy propyl starch, for example, PureCote®, and oxidized starches. Alternatively, the active agent may be added to the core, or added to both the core and a layer surrounding the core.

PROTECTIVE COATINGS

[0072] The protective coatings of the present invention generally are applied as one or more layers surrounding the core when the active agent is not inherently thermostable. Embodiments include one, two, three or four protective coating layers.

[0073] Suitable protective coating materials are polymers, carbohydrates, proteins, lipids, fats and oils, fatty acids, inorganic salts, and gums and mixtures thereof.

[0074] The protective coatings include optional moisture barrier coatings and a moisture hydrating coating as defined in claim 1. The optional moisture barrier coatings function by excluding moisture, for instance by forming a shell layer that typically does not absorb moisture and prevents or retards the rate of moisture migration into the granule. Moisture hydrating coatings on the granule absorb or bind moisture as either free water or water of hydration, thereby acting to impede or retard the extent or rate of transport of external moisture into the granule. The moisture hydrating coatings typically constitute at least about 35% w/w of the granule. The moisture hydrating materials in the coatings thermally insulate the active agents and will absorb a certain amount of moisture and retain it within the hydrating material without allowing it to pass through into the portion of the granule having the active agent. For moisture hydrating coatings on stable, durable granules that do not contain appreciable amounts of water prior to steam treatment, such coatings may constitute about 25% w/w of the granule.

[0075] Moisture barrier coatings typically comprise hydrophobic materials, such as hydrophobic polymers, for example PVA, HPMC, acid-thinned hydroxypropyl starches and oxidized starch; proteins, for example whey and whey protein concentrates; lipids, for example, lecithin; fats and oils, fatty acids, latex and gums, for example, gum arabic. Certain moisture barrier coatings, such as PVA and gum arabic are not readily oxidized and find particularly applicability in providing chemical stability when the granules of the invention are stored in unpelleted or untableted mixtures, for instance, in premixes that contain choline chloride. Moisture hydrating coating materials typically are hydrophilic materials, such as carbohydrates and inorganic salts, including hydrated salts. Examples of moisture hydrating materials are magnesium sulfate, sodium sulfate, maltodextrin, ammonium sulfate, sugars, for example, sucrose, and native cornstarch.

[0076] Polymers used for the protective coatings are polyvinyl alcohol (PVA), polyethylene glycol, polyvinyl pyrrolidone, polyacrylates, polyethylene oxides (PEO), polylactic acid, polyvinylchloride, polyvinylacetate, polyvinyl pyrrolidones (PVP), cellulose ethers, alginates, gelatin, modified starches and substituted derivatives, hydrolysates and copolymers thereof, such as acid-thinned hydroxypropyl starch, such as, Pure Cote™, hydroxypropyl methyl cellulose (HPMC), methyl cellulose (MC), carboxymethyl cellulose (CMC), and ethyl cellulose. Most preferred polymer for the protective coatings are PVA, modified PVA, as described in U.S. Pat. NO. 6,872,696, and modified cellulose, such as methyl cellulose and hydroxypropylmethyl cellulose, as described in PCT Publication No. WO 99/51210, both of which are incorporated by reference wherein.

[0077] Carbohydrates used for the protective coatings are maltodextrin hydroxymethyl cellulose, modified or native starches made from corn, sorghum, arrowroot, rice, wheat, rye, barley, oat, potato, yam, tapioca, cassava, sago, and sugars including sucrose, corn syrup solids, molasses, glucose, fructose, and lactose.

[0078] Proteins used for the protective coatings are whey powder, whey protein concentrate, whey protein isolate, caseinates, soy protein concentrate and isolate, zein, albumin and gelatin.

[0079] Simple, compound and derived lipids that may be used in the protective coatings are waxes (for example, vegetable, mineral and synthetic, such as carnauba, candelilla, beeswax, cerumen, carnuba, shellac, paraffin, and microcrystalline waxes); lecithin (for example mono- and diglycerides); fatty acids (for example stearic, palmitic, linoleic, oleic, butyric, and arachidonic fatty acids and their salts of sodium, potassium, calcium and zinc); and fats and oils (for example, hydrogenated or partially hydrogenated fats and oils, such as soy, corn, cottonseed, tallow, canola, and linseed oil). A preferred lipid for the protective coatings is lecithin.

[0080] Inorganic salts used for the protective coatings include salts of sulfate, citrate, chloride, carbonate, sulfite, phosphate, phosphonate, and bicarbonate salts of sodium, ammonium, potassium, calcium, magnesium and zinc. Preferred salts are magnesium, sodium and ammonium sulfates.

[0081] Gums that may be used in the protective coatings include gum arabic, guar gum, agar, gum tragacanth, kary gum, locust bean gum, carageenan, xanthan gum, and alginates.

[0082] The protective coatings of the present invention further may include plasticizers, lubricants, pigments and powders, such as talc, bentonite, kaolin, cornstarch, magnesium silicate, calcium carbonate, and chitosan.

[0083] Certain embodiments of the present invention typically have a single layer of a moisture hydrating material that is

approximately at least 55% w/w of the granule. Because the capacity of moisture hydrating coatings to take up and sequester water has a limit, relatively high levels of single layer coatings are applied. Alternatively, moisture hydrating material(s) may be applied in two layers. Other embodiments of the present invention have protective coatings utilizing both moisture hydrating materials and moisture barrier materials. In these embodiments, the amount of moisture hydrating material may be lower, at least about 25% w/w of the granule and the moisture barrier material is about 2% to 25% w/w of the granule. Using both moisture hydrating materials and moisture barrier materials combines protective mechanisms and typically reduces cost, particularly of the moisture barrier materials. Moisture barrier materials, particularly film-forming materials may be subject to mechanical damage which, if these materials are used alone as a thin coating may lead to loss of protection for the active agent. The combination allows for the use of less of both materials than would be required if the materials were used alone. The combination allows for some damage to the moisture barrier layer in view of the presence of the moisture hydrating material. As discussed above, the combination of the materials is particularly suitable to provide chemical stability by maintaining activity when the granules are stored in unpelleted mixtures.

[0084] Processes of the present invention, in addition to the processes described for manufacture of the stable, durable granules, include mixing of the granules with unpelleted feed mixtures and storage of the resulting mixtures. Additional processes include steam-treating the resulting mixtures and/or pelleting the resulting mixtures. Of course the stable, durable granules of the present invention also may be stored alone and mixed with feed, or pelleted, when desired.

EXAMPLES

[0085] Table 1 below is a list of ingredients and abbreviations used in the following examples.

Table 1

Ingredient	Name & Product No.	Supplier
Sucrose	Domino Pure Cane Extra Fine Granulated Sugar	Tate & Lyle North American Sugars, Baltimore, MD
Cornstarch	Corn Starch	Cargill Foods, Minneapolis, MN
MgSO ₄	Magnesium sulfate heptahydrate, MgSO ₄ 7H ₂ O (Epsom salts)	PQ Corporation, Berwyn, PA
HPMC	Hydroxypropyl Methylcellulose, Brandname: METHOCEL	Dow Chemical
Carnauba wax	Michem Lube 160HS (carnauba emulsion, 50% solids)	Michelman Inc. Cincinnati, OH
Modified starch	PURE-COTE®	Grain Processing Corporation, Muscatine, IA
Lecithin	ULTRALEC® P Soy lecithin	ADM Corp., Decatur, IL
PVA	polyvinyl alcohol, Elvano® 51-05	DuPont, Wilmington, DE
Sodium sulfate	Sodium Sulfate Anhydrous	Cooper Natural, Tulsa, Oklahoma
Ammonium sulfate	Ammonium Sulfate CAS # 7783-20-2	General Alum & Chemical Corp. Searaport, Maine
Talc	NYTAL 4000 TALC	RT Vanderbilt Co, Inc. Norwalk, CT
Whey powder	PC42010	Leprino Foods, Denver CO
Whey protein	Proliant instantized whey protein concentrate, #8010, lot H4212	Hilmar Cheese Company, Hilmar, CA (WPC formally sold through Proliant)
Linseed oil	ASTM Raw Linseed Oil	Cargill Industrial Oils, Chicago, IL
Maltodextrin	Maltodextrin M150	Grain Processing Corp (GPC) Muscatine, IA
Gum Arabic	TIC PRETESTED Gum Arabic FT PreHydrated	TIC Gums, Belcamp, MD
Canola oil	Canola Oil	Safeway
Latex	Aquacoat ECD (Ethylcellulose dispersion 30%)	FMC BioPolymer, Philadelphia, PA
Sand	Sodium silicate	Sigma-Aldrich Chemical Co
Corn cobs	Ground corn cobs or corn pith	ICBP Independence, IA

[0086] Table 2 illustrates the composition of a number of exemplary stable, durable granules and several granules that did not demonstrate at least 50% recovered active agent activity under particular pelleting conditions. None of these granules is a granule of the present invention.

Table 2

Formulation #1*		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	36.9
Active		
Agent	Enzyme	7.4
	sucrose	12.6
	Cornstarch	12.6
1st coating	Sucrose	10.5
	Cornstarch	10.5
2nd Coating	carnauba wax	6.0
3rd coating	HPMC	3.5
	*Heat annealing step	
Formulation #2		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	17.5
Active		
Agent	enzyme	3.5
	sucrose	6.0
	cornstarch	6.0
1st Coating	MgSO ₄	67.0
Formulation #3		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	16.5
Active		
Agent	enzyme	4.8
	sucrose	4.0
	cornstarch	8.0
1st Coating	MgSO ₄	66.7
Formulation #4		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	17.5
Active		
Agent	enzyme	3.5
	sucrose	4.0
	cornstarch	8.0

Formulation #4		
	Component	% (w/w) dry basis
1st Coating	maltodextrin	67
Formulation #5		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	14.8
Active		
Agent	enzyme	2.9
	sucrose	3.3
	cornstarch	6.6
1 st Coating	MgSO4	55.6
2nd Coating	gum arabic,	17.0
Formulation #6		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	14.8
Active		
Agent	enzyme	2.9
	sucrose	3.3
	cornstarch	6.6
1st Coating	MgSO4	55.6
2nd Coating	whey	16.7
Formulation #7		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	14.8
Active		
Agent	enzyme	2.9
	sucrose	3.3
	cornstarch	6.6
1st Coating	MgSO4	55.6
2nd Costing	protein concentrate	17.0
Formulation #8		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	13.4
Active		
Agent	enzyme	2.6
	sucrose	3.0
	cornstarch	6.0
1st	PureCote® starch	15.0
2nd Coating	lecithin	10.0
3rd Coating	MgSO4	50.0

Formulation #9		
	Component	% (w/w) dry basis
core		
Material	sodium sulfate	9.1
Active		
Agent	enzyme	1.8
	sucrose	2.0
	cornstarch	4.1
1st Coating	MgSO4	83.0
Formulation #10 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	26.5
Active		
Agent	enzyme	5.3
	sucrose	9.1
	cornstarch	9.1
1st Coating	carnuaba wax	50
Formulation #11		
	Component	% (w/w) dry basis
Core		
Material Active	sodium sulfate	10.7
Agent	enzyme	2.1
	sucrose	2.4
	cornstarch	4.8
1st Coating	PVA	4
	talc	36
2nd	Coating MgSO4	40
Formulation #12 (Test granule with no protective coating)		
	component	% (w/w) dry basis
Core		
Material Active	sucrose 35-50 mesh	53.1
Agent	enzyme	10.6
	sucrose	18.1
	cornstarch	18.1
Formulation #13 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	44.2
Active		
Agent	enzyme	8.9
	sucrose	15.1
	cornstarch	15.1
1st Coating	sucrose & starch (1:1 ratio)	16.7

Formulation #14 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	48.3
Active		
Agent1st		
coating	enzyme	9.7
	sucrose	16.5
	cornstarch	16.5
1 st Coating	caruaba wax	9.1
Formulation #15 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	49.6
Active		
Agent	enzyme	9.9
	sucrose	16.9
	cornstarch	16.9
1 st Coating	ethylcellulose	11.5
Formulation # 16 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	48.3
Active		
Agent	enzyme	9.7
	sucrose	16.5
	cornstarch	16.5
1 st Coating	hydroxypropylmethylcellulose	9.1
Formulation #17		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	10.7
Active		
Agent	enzyme	2.1
	sucrose	2.4
	cornstarch	4.8
1 st Coating	MgSO4	40
2nd Coating	PVA	4
	talc	36
Formulation #18		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	13.4
Active		

Formulation #18		
	Component	% (w/w) dry basis
Agent	enzyme	2.6
	sucrose	3.0
	cornstarch	6.0
1st Coating	MgSO4	50.0
2nd Coating	lecithin	10.0
	PureCote® starch	15.0
Formulation #19		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	16.6
Active		
Agent	enzyme	4.8
	sucrose	4.0
	cornstarch	0.0
	talc	8.0
1st Coating	MgSO4	66.7
Formulation #20		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	14.8
Active		
Agent	enzyme	2.9
	sucrose	3.3
	cornstarch	6.6
1st Coating	gum arabic	17.0
2nd Coating	MgSO4	55.6
Formulation #21 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sucrose 35-50 mesh	51.9
Active		
Agent	enzyme	10.4
	sucrose	17.7
	cornstarch	17.7
1st Coating	canola oil	2.2
Formulation #22		
	Component	% (w/w) dry basis
Core		
Material	ground corncob	17.5
Active		
Agent	enzyme	3.5
	sucrose	6.0

Formulation #22		
	Component	% (w/w) dry basis
	cornstarch	6.0
1st Coating	MgSO4	67
Formulation #23		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	31.1
Active		
agent	enzyme	2.4
	cornstarch	9.4
1st Coating	MgSO4	43.7
2nd Coating	Gum Arabic	3.4
	maltodextrin	10.1
Formulation #24 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	29.9
Active agent	enzyme	3.3
	cornstarch	9.3
1st Coating	PureCote	12.0
2nd Coating	Lecithin	8.0
3rd Coating	MgSO4	37.5
Formulation #25 (Thermolabile)		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	47.8
Active		
agent	Enzymes	5.3
	cornstarch	14.9
1st Coating	PureCote	19.2
2nd Coating	Lecithin	12.8
Formulation #26		
	Component	% (w/w) dry basis
Core		
Material	sodium sulfate	31.0
Active		
agent	Enzyme	2.4
	Cornstarch	9.4
1st Coating	sodium sulfate	43.7
2nd Coating	gum Arabic	13.5

Formulation #27		
	Component	% (w/w) dry basis
Core		
Material sodium	sulfate	31.0
Active		
agent	Enzyme	2.4
	Cornstarch	9.4
1st Coating	MgSO4	43.7
2nd Coating	Whey	13.5
Formulation # 28**		
	Component	% (w/w) Dry Basis
Core Material	sodium sulfate	32.7
Active Agent #1	enzyme #1	9.1
	sucrose	4.1
	corn starch	8.5
Active Agent #2	enzyme #2	1.9
	sucrose	0.9
	corn starch	1.8
Coating #1	sodium sulfate	33.6
Coating #2	PVA 51-05	2.5
	talc	5.0
**Inherently thermostable enzyme		
Formulation #29**		
	Component	% (w/w) Dry Basis
	sodium	
Core Material	sulfate	32.7
Active Agent #1	enzyme #1	
and #2 Mixed	and #2	10.9
	sucrose	5.0
	corn starch	10.3
	sodium	
Coating #1	sulfate	33.6
Coating #2	PVA 51-05	2.5
	talc	5.0
**Inherently thermostable enzyme		
Formulation #30**		
	Component	% (w/w) Dry Basis
Core Material	Sand	84.80
Active Agent #1	enzyme #1	3.52
	sucrose Whole	1.62
	Ground	
	Wheat	3.31
Coating #1	sodium sulfate	5.50
Coating #2	PVA 51-05	0.42

Formulation #30**		
	Component	% (w/w) Dry Basis
	talc	0.83
**Inherently thermostable enzyme		
Formulation #31		
	Component	% (w/w) Dry Basis
Core Material	sodium sulfate	40.0%
Spray 1	enzyme	5.0%
	PVA	1.0%
	corn starch	5.0%
Spray 2	sodium sulfate	40.0%
Spray 3	PVA	3.0%
talc 6.0%		
Formulation #32		
	Component	% (w/w) Dry Basis
Core Material	sodium sulfate	40.0%
Spray 1	enzyme	5.0%
	corn starch	9.0%
Spray 2	sodium sulfate	39.0%
Spray 3 gum arabic 7.0%		
Formulation #33		
	Component	% (w/w) Dry Basis
Core Material	Sodium sulfate	54.8
Active agents	Mixed enzymes	11.8
(2)	sucrose	5.4
	corn starch	11.1
Coating 1	Sodium sulfate	26.0
Coating 2	PVA	2.5
	talc	4.9
Formulation #34		
	Component	% (w/w) Dry Basis
Core Material	Sodium sulfate	54.8
Active agents (2)	E enzyme mixture	11.8
Coating 1		26.0
Coating 2	Sodium sulfate	2.5
	PVA	4.9
	talc	

REFERENCE EXAMPLE 1. PREPARATION OF GRANULES IN TABLE 2.

[0087] All of the granules in Table 2, except granule number 1, are granules that were prepared using a fluid bed process as described in US Pat. No. 5,324,649. The fluid bed process fluidized the core materials in a Vector FL-1 processor (made by Vector Corp, Marion, IA, USA), a Glatt 3, or a Uniglatt processor (both made by Glatt Air Techniques, Binzen, Germany). An enzyme/sugar/starch mixture was spray coated onto the core material. Then, any protective coating(s) were sprayed sequentially

onto the enzyme layer and allowed to dry.

[0088] For example, the formulation #3 granule was prepared as follows:

In a Glatt 3 top spray suit bed coater, sodium sulfate crystals screened to - 45/+140 mesh were charged and fluidized using a heated bed temperature. A xylanase ultra filtration concentrate from *Trichoderma reesei* was mixed with corn starch and sucrose and sprayed onto the crystals. The solution was about 33% dry solids. The final batch weight was 4000 grams.

[0089] In a vector FL-1 top spray coater, moisture hydrating material, a magnesium sulfate solution, was sprayed onto 833 grams of the material described above. The bed temperature was 50°. The final batch weighed 2500 grams.

[0090] Preparation of Granule number 1 further included a heat annealing step in which the granule was heated to a temperature sufficient to exceed the glass transition temperature (T_g) of the carnauba wax material followed by slow cooling which, produced a hardened, glassy wax layer.

REFERENCE EXAMPLE 2: PREPARATION OF MASH SAMPLES WITH GRANULES AND PELLETING

[0091] Three different feed formulations and pelleting processes were used to prepare pellets with the granules listed in Table 2. Relatively high dosages of granules were added to the feed formulations to optimize the active agent remaining activity assays.

Mill #1:

Selected granules from Table 2 were mixed together with a feed formulation.

The composition of the feed formulations was as follows:

75% (w/w) cornmeal (enriched yellow degerminated cornmeal, no 50956, General Mills Operations, Minneapolis, MN); and

25% (w/w) soybean meal (Pro Soybean meal, Cargill Oilseed Co., Cedar Rapids, IA).

12 kg of the above feed mixture was combined with each sample granule (dosed at 5 g/kg of feed formulation), and blended in a large Hobart blender (model D-300T, Troy OH), for 8 minutes. Approximately 150 g of each sample was retained as the mash sample, or unpelleted feed mixture. Each batch was then split into three 4 kg sub-batches, and pelleted in triplicate pellet mill runs.

[0092] The pellet mill used was CPM model CL5 (California Pullet Mill Co., Crawfordsville, IN). The temperature in the steam conditioner was controlled by the amount of injected steam, at 1.36 atm. Conditioning temperature of the mash, measured immediately prior to entering the die of the pellet mill, was targeted to 89-90°C. Temperature measurement was done using a thermometer and a "J" type thermocouple (OMEGA Engineering, Inc. Stamford, CT). The residence time in the conditioner was approximately 5 seconds. Die dimensions were 12.7 cm inner diameter 17.8 cm outer diameter, 4.7 mm hole diameter. The pellet mill was run continuously, and the samples were processed through the mill sequentially, separated by running approximately 2 kg of a cornmeal/soy mix between samples. Approximately 1 minute after the test mash was added to the conditioner, and the conditioner temperature had stabilized to target temperature, pellets were collected over a 30 second interval, allowing approximately 1 kg of pellets to be collected for each sample. The collected pellets were held under ambient conditions, for 30 seconds, and then cooled with an air cooler for 2-3 min to room temperature.

Mill #2:

Prepared durable granules from Table 2 were pelleted with a corn and soy feed formulations. The exact composition of the feed formulations was as follows:

61.2% (w/w) cornmeal, 31.6% (w/w) soybean meal, 3.0% meat and bone meal, 2.5% soybean oil, 1.3% limestone, 0.28 % Salt, and 0.08% methionine. For each granule tested, 370 to 2000 grams of the granules, depending on enzyme type and activity, were combined with 450 kg of the feed without oil, and blended in a plow mixer for 2 minutes. Then soybean oil was added and the sample was mixed for an additional 3 minutes. The pellet mill was the Master model, manufactured by California Pellet Mill. Die pellet hole diameter was 4.5 mm. The typical feed rate was 780 kg per hour. The temperature in the steam conditioner was controlled manually, and was measured at the feed outlet from the conditioner. Two conditioning temperatures were used; 85°C and 95°C. The residence time in the conditioner was approximately 30 seconds. When the target temperature was reached, the system was run for approximately 5 minutes before sampling took place. Samples of approximately 5 kg of pelleted feed were

taken, and were cooled by spreading on screened trays.

Mill #3:

The prepared durable granules were pelleted with a wheat and soy feed formulation or a wheat and barley formulation. The composition of the feed formulation was either 60 % wheat, 31.5% soybean meal, 4% soybean oil, 1.5% dicalcium phosphate, 1.23% vitamin mineral premix, 1.2% limestone, 0.4% salt, and 0.2% DL methionine; or 60% wheat, 30% barley, with two conditioning temperatures, 90° C and 95° C used. For each granule tested, 200 to 500 grams of granules, depending on enzyme type and activity, was combined with 160 kg of the feed, and blended in a horizontal ribbon mixer, for approximately 15 minutes. The pellet mill was a Simon Heesen, mono roll type, fitted with a 17.3 cm inner diameter die, with a pellet hole diameter of 3 mm. Die speed was 500 rpm and was driven by a 7.5 kW motor. The typical feed rate was 300 kg per hour. The temperature in the steam conditioner was kept at +/- 0.1 degrees Celcius, measured at the feed outlet from the conditioner. The conditioner had a cascade type mixer system. Three conditioning temperatures were used; 85°C 90 °C, and 95°C. Steam inlet pressure was 2 atm, and the temperature in the conditioner was controlled by manual adjustment of three valves that regulate the steam deliver. The residence time in the conditioner was approximately 30 seconds. When the target temperature was reached, the system was run for approximately 5 to 10 minutes before sampling took place. Samples were taken for 1-1.5 minute periods, corresponding to 5-7.5 kg of pelleted feed, and were immediately placed in a cooling box with a perforated bottom and air flow of 1500 cubic meters per hour. After cooling for 15 minutes, the samples were downsized twice using a sample divider, and 1 kg was taken for lab tests.

REFERENCE EXAMPLE 3: ENZYME ACTIVITY MEASUREMENTS

Determination of enzyme activity

[0093] To determine the enzyme activity after pelleting, the mash and pelleted samples were then ground for 30 seconds in a kitchen coffee grinder (model 203-42, Krups North America Inc., Medford MA), and assayed for enzyme activity as described below. Alternatively, samples were ground in a ZM-200 centrifugal mill, fitted with a 1 mm sieve (Retsch GmbH, Germany).

[0094] Calculation of Percent Recovered Activity:

For each test sample, both mash and the corresponding pelleted samples were assayed for activity. The percent recovered activity was calculated as follows:

$$\% \text{ recovered activity} = \frac{\text{activity in pellet} \times 100}{\text{activity in the mash}}$$

[0095] The phase enzyme assay was conducted according to AOAC (Association of Analytical Chemists) Official Method 2000.12, as described in "Determination of phytase activity in feed by a colorimetric enzymatic method collaborative interlaboratory study". Engelen AJ, van der Heeft FC, Randsdorp PH, Somers WA, Schaefer J, van der Vat BJ. JAOAC Int. 2001 May-Jun; 84(3):629-33. Briefly, the ground samples extracted in 220 mM sodium acetate trihydrate, 68.4 mM calcium chloride dihydrate, 0.01% Tween 20, pH 5.5. The supernatant is then assayed. The assay measures the release of inorganic phosphate from rice phytate, at pH 5.5, for 60 min at 37C. The assay is stopped with acidic molybdate/vanadate reagent, and phosphate is quantified by intensity of yellow colored complex of the vanadomolybdophosphor.

[0096] The xylanase enzyme assay was performed using the Xylanase Assay Kit (Xylazyme AX Format), Cat No: K-XYLS Megazyme International Ireland Ltd., Wicklow, Ireland. Materials for the essay include 16 x 125 mm disposable glass culture tubes; Extraction buffer: 100 mM MES sodium salt buffer, pH 6.0; Assay buffer: 25 mM sodium phosphate buffer, pH 6.0; Stopping solution: 20 g NaPO₄12H₂O in 1 liter MilliQ water; Polystyrene Tissue Culture Flasks, 225 ml (Coming Incorporated - Life Sciences Big Flats, New York). 6-12 g of ground feed is extracted in 120 ml of extraction buffer, in a culture flask, for 1 hour at 20-25° C, while shaking on an orbital gel shaker set to 100 rpm. Samples are then centrifuged at 2000 g for 1 min, and the supernatant is assayed. For the assay, 20 to 100 ul of the feed supernatant is diluted into 500 ul using assay buffer, in culture tubes, and equilibrated in a water bath at 40°C for 10 min. Then, one substrate tablet is placed into each culture tube, and samples are incubated for an additional 10 minutes at 40C. 10 ml of stopping solution is then added to each tube. The samples are vortexed briefly, and then filtered through Whatman #1 filter paper. The absorbance of the filtrate is read with a spectrophotometer at a wavelength of 590 nm. The spectrophotometer is first zeroed with a blank, made by combining 20 to 100 ul of the feed supernatant,

500 ul assay buffer, and 10 ml of stopping solution. One substrate tablet, is then added, followed by, vortexing, and filtering in the same manner as the samples. Interference from some feed components may affect assay response. In order to correct for any interference, standard curves with feed background were prepared. Uncoated xylanase granules were added to blank mash, and to blank pellets, at several levels. The spiked mash and pellet samples were then ground, and extracted exactly as described above. From this series of extracts, standard curves for both the mash and the pellets were generated.

[0097] The beta-glucanase enzyme assay was performed using a Beta-glucanase Assay Kit (Beta-Gluczyme Tablet format), Cat No: T-BGZ200, Megazyme International Ireland Ltd., Wicklow, Ireland. Materials for the assay include 16 x 125 mm disposable glass culture tubes; Extraction buffer and assay buffer: 25 mM sodium acetate buffer; Stopping solution: 2% NaPO₄12H₂O in 1 liter MilliQ water; 140 ml glass beakers. 10 g of ground feed was extracted in 100 ml of extraction buffer, by mixing in a beaker for 1 hour at 20-25°C. Samples are then centrifuged at 2000 g for 1. min, and the supernatant is assayed. For the assay, 20 to 100 ul of the feed supernatant is diluted into 500 ul using assay buffer, in culture tubes, and equilibrated in a water bath at 40°C for 10 min. Then, one substrate tablet is placed into each culture tube, and samples are incubated for an additional 10 minutes at 40°C. 10 ml of stopping solution is then added to each tube. The samples are vortexed briefly, and then filtered through Whatman #1 filter paper. The absorbance of the filtrate is read with a spectrophotometer at a wavelength of 590 nm. The spectrophotometer is first zeroed with a blank, made by combining 20 to 100 ul of the feed supernatant, 500 ul assay bluffer, and 10 ml of stopping solution. One substrate tablet, is then added, followed by, vortexing, and filtering in the same manner as the samples. A standard curve is prepared by assaying a series of enzyme diluted to appropriate levels in assay buffer.

RETAINED ENZYME ACTIVITY RESULTS

[0098]

Table 3: Pelleting Results - Percent Recovered Activity after pelleting

Formulation	Mill #1, 90°C	Mill #2, 3, or 4 at 85-95°C
10	< 30% xylanase	
12	< 30% xylanase	
13	< 30% xylanase	
14	< 30% xylanase	
15	< 30% xylanase	
16	< 30% xylanase	
21	< 30% xylanase	
25	< 30% xylanase	
1	> 50% xylanase	
2	> 50% xylanase	
3	> 70% xylanase	> 70% xylanase
5	> 50% xylanase, > 70% e. coli phytase	> 50% xylanase, > 80% e. coli phytase
6	> 50% xylanase	
7	> 50% xylanase	
9	> 50% xylanase	
11		> 50% xylanase, > 70% Aspergillus phytase
30		>60% thermostable xylanase
22	> 50% xylanase	
23	> 60% e. coli phytase	> 50% xylanase, > 80% e. coli phytase
27	> 50% xylanase	> 80% e. coli phytase
33		>90% BGL, >70% thermostable xylanase
4	> 70% xylanase	
8	> 70% xylanase	
17	> 70% xylanase	

Formulation	Mill #1, 90°C	Mill #2, 3, or 4 at 85-95°C
18	> 70% xylanase	
19	> 70% xylanase	
20	> 70% xylanase	
26	> 70% xylanase	> 90% e. coli phytase
28		> 90% BGL, > 90% thermostable xylanase
29		> 70% BGL, > 90% thermostable xylanase
31		> 90% e. coli phytase
32		> 90% e. coli phytase

[0099] The pellet mills used can be broken into two groups. Mill #1 is a lab scale mill, which is much harsher than what is typically encountered in commercial practice. The harshness is attributed to the large amount of steam which is delivered to a relatively small portion of mash, coupled with the fact that the feed mixture does not contain oil, which typically lubricates the feed while traveling through the die. Mills #2 through #4 are considered to be more representative of commercial conditions.

[0100] The retained activities of the tested granules are shown in Table 3. Granules of the present invention that are not considered stable, durable granules were numbers 10, 12, 13, 14, 15, 16, 21 and 25, all of which demonstrated less than 30% recovered activity. These granules have a core coated with an enzyme matrix layer wherein the coated core is greater than about 50 w/w% of the entire granule. Most of these non-stable granules had only one protective coating layer of a moisture barrier material, such as carnauba wax, or ethylcellulose, or HPMC, or canola oil. Without wishing to be bound by any particular theory, and recognizing that these granules may show greater than 50% retained active agent activity at 70-85°C, the poor thermostability of these granules under these test conditions, relative to stable, durable granules, may be due to the provision of coating materials as relatively thin layers, comprising, in five examples, about 2.0 to about 17.0% w/w of the granule, and/or the particular coating agent employed. For instance, granule number 10 has a single layer of coating material that is 50 w/w% moisture barrier material (carnauba wax); and granule number 25 has two moisture barrier protective coating layers that together constitute 32% w/w of the granule.

[0101] Particularly stable, durable granules of the present invention were numbers 4, 8, 17, 18, 19, 20, 26, 28, 29, 31 and 32, all of which have greater than 70% recovered xylanase activity when pelleted at about 90° C for about 5 seconds. Additionally, granules 26, 31 and 32 demonstrated greater than 90% recovered phytase activity, when pelleted between 85° C and 95° C for about 30 seconds. Granules 28 and 29 demonstrated greater than 90% retained activity of one inherently thermostable xylanase in the enzyme mixture and 70%-90% retained activity of a thermolabile beta-glucanase when pelleted between 85° C and 95° C for 30 seconds. Those stable durable granules having an active agent that is not inherently thermostable are either; 1) coated with a single, thick protective layer of a moisture hydrating material, or 2) coated with two to three protective layers, where at least one of the layers is a moisture hydrating material and one layer is a moisture barrier material.

[0102] In general, when moisture hydrating materials are applied as individual layers, the coating must comprise from about at least about 55% of the granule, to provide suitable thermo protection; and, when the amount of moisture hydrating material was lowered to less than 55% using an individual layer, percent retained activity dropped to less than 50%. However, when moisture hydrating materials are used in combination with moisture barrier materials, the water hydrating protective layer may be lowered to at least about 25% w/w of the granule, while the moisture barrier material may be about 2% to 40% w/w of the granule.

[0103] Other stable, durable granules of the present invention were numbers 1, 2, 3, 5, 6, 7, 9, 11, 22, 23, 27, 30 and 33 all of which demonstrated greater than 50% recovered xylanase or phytase activity when pelleted at 90°C for 5 seconds or at 85° C to 95° C for 30 seconds. As can be seen from these results, inorganic salt outer or inner coatings alone provide good protection for the enzyme and enhanced protection when combined with moisture barrier coatings. As noted earlier, granule number 1 was made using a heat annealing processing step and it is believed that addition of a heat annealing step enhances barrier properties of the moisture barrier materials, such as polymers, proteins, and lipids thereby reducing the amount of moisture hydrating material needed to provide thermostability. When a granule otherwise identical to granule #1 was made without heat annealing, the recovered activity was 35%. Granules 2, 3, and 22 show that different core materials can be used without compromising granule stability.

[0104] Granule 3 demonstrated greater than 70% recovered enzyme activity at both 5 seconds and 30 seconds pelleting time periods. Granule 11 demonstrated greater than 50% recovered xylanase activity when pelleted at 85° C to 95° C for about 30

seconds, and greater than 70% recovered phytase activity when pelleted at 85° C to 95° C for about 30 seconds. Granule #23 demonstrated greater than 60% recovered activity of phytase when pelleted for 5 seconds at 90° C; greater than 50% recovered activity of xylanase when pelleted at 85° C to 95° C for 30 seconds; and, greater than 80% recovered activity of phytase when pelleted at 85° C to 95° C for 30 seconds.

REFERENCE EXAMPLE 4: DIRECT STEAMING TEST

[0105] 100 gram portions of mash comprised of 75% cornmeal and 25% soy were dosed with granules, containing *E. coli* phytase, at an inclusion rate of 1.25 g of granules per 1 kilogram of mash. Mash samples were then wrapped in a cheese cloth bag, and steamed for 20 seconds by placing the bags into a funnel with 30 psi steam applied to the bottom of the funnel. After steaming, samples were cooled to ambient temperature, allowed to dry overnight, and then assayed for phytase activity. The percent recovered activity of the steamed samples, relative to unheated mash, was reported.

[0106] A granule that was not considered suitably steam resistant under these test conditions was number # 25, which had less than 35% recovered activity after steaming. Stable, durable, granules of the present invention were numbers #5 and #23, which had greater than 50 % recovered activity after steaming.

REFERENCE EXAMPLE 5: BIOAVAILABILITY TEST OF PELLETTED GRANULE

[0107] The stable, durable granules of the present invention may be tested for bioavailability of the active agent enzyme using known bioavailability tests, such as the phytase bioavailability study disclosed in WO 00/47060; bioavailability tests described in Enzymes in Animal Nutrition, Proceedings of the 1 st Symposium, Kartause Ittingen, Switzerland, October 13016, 1993; the bioavailability tests described in Chemgen patents.

[0108] Stable granules of the present invention were assessed after pelleting to determine bioavailability in broiler chicks fed a commercial diet.

Materials and Methods

[0109] Tested Enzymes and Granules and Dietary Treatments included Phyzyme[®] XP (6-phytase; E.C. 3.1.3.26) enzyme alone, and both PVA-coated and Gum Arabic (GA) coated stable granules of the present invention containing Phyzyme XP enzyme. (See Table 4). The tested enzymes and granules were fed to the broiler chickens in either a mash diet or a pellet (prepared at 90°C) diet. The experimental diets were: positive control (commercial diet), negative control, negative control + 500 U/kg Phyzym[®] XP, negative control + 500 U/kg PVA-coated-phytase, negative control + 500 U/kg GA-coated-phytase, negative control + 500 U/kg PVA-coated-phytase pelleted at 90°C, and negative control + 500 U/kg GA-coated-phytase pelleted at 90° C. Except for calcium and phosphorus, all the diets were isocaloric and isonitrogenous. (See Table 6). Calcium and available phosphorus in the positive and negative control diets were 0.90% and 0.78%, and 0.37% and 0.26%, respectively. Titanium dioxide (0.10%) was added to the diets as an indigestible marker to help estimate nutrient digestibility. The experimental diets were manufactured by ADAS Gleadthorpe to a commercial specification and retained at ADAS Gleadthorpe and stored under refrigeration. Samples of the enzyme additive mixes (25g), cereal (250g), soybean meal (250g) and a sample of all the diets (250g) were sent to Danisco Animal Nutrition Enzymes Feed Services, Edwin Rahrs Vej 38, DK-8220 Brabrand, Denmark. All feed samples were assayed for phytase prior to feeding.

Experimental Animals and Design:

[0110] The chicks were male Ross 308 broilers aged 0 to 21 days using eight replicate pens per treatment with 30 birds per pen. The chicks were placed randomly into boxes before being weighed and randomly allocated to the treatment pens. Each pen contained one tube feeder and birds had free access to water via nipple drinkers (4 per pen) and feed at all times. The height of the drinkers was adjusted regularly to keep level with the top of the back of the birds. Litter was provided in the form of clean wood shavings to a depth of 5cm. The house was heated by a warm air brooding system and the brooding target temperature regime was 31 °C at day old, reducing by 1°C every other day until 21°C was reached on day 21. The minimum ventilation rate was automatically calculated to supply $1.9 \times 10^4 \text{ m}^3$ of air per second per $\text{kg}^{0.75}$ bodyweight and this rate was supplied by one

610mm fan controlled by a Farm-Ex Diacam control panel. Lighting programme was 23 hours light and 1 hour dark. Light intensity was reduced from maximum attainable at day old (40-60 lux) to an intensity of approximately 10 lux by 10 days of age. Relative humidity was monitored on a daily basis using a Tinytalk® data logger. Chicks were vaccinated with IB H120 & 50% Avinue (ND) at Hatchery.

Feed Intake/Body Weight

[0111] Feed usage was measured between 0 and 21 days. The feed in each pen was measured by weighing back the amount of feed remaining at the end of the period and deducting it from the quantity offered. The birds were weighed in their pen groups at day old. At 21 days all the birds in all of the pens were weighed in batches. The total weight of all the birds in each pen was determined and the mean calculated. All birds that died were weighed and the details recorded, and any lame birds or birds unable to reach feed and water were culled from the study and the reason for culling was listed.

Tibia ash

[0112] At 21 days of age 2 birds from each pen were culled using cervical dislocation and the left leg removed. The tibia was dissected out and the bone sent to Eurofins laboratories, Woodthorne, Wolverhampton for ashing on a plot basis. All the data were analyzed using general linear models procedure of SAS (SAS Inst. Inc., Cary, NC). Where significant differences were found, Duncan's test was used to compare individual treatments means.

Results and Discussion

[0113] Analyzed phytase activity in the experimental diets were <50, <50, 563, 467, 558 FTU/kg (PVA-coated-phytase), and 554 and 440 (GA-coated-phytase), for positive control, negative control, negative control + 500 U/kg Phyzyme® XP, negative control + 500 U/kg PVA-coated-phytase and GA-coated-phytase, and negative control + 500 U/kg PVA-coated-phytase and GA-coated-phytase pelleted at 90°C., respectively (Table 5), indicating that activity of both the PVA-coated-phytase and the GA-coated-phytase was not destroyed after pelleting at 90°C.

[0114] Birds fed diets supplemented with 500 U/kg PVA-coated-phytase and GA-coated-phytase, pelleted at 90°C, were heavier at day 21 than birds fed the positive control, negative control and negative control + 500 U/kg Phyzyme® XP diets (Table 7). Birds fed negative control diet consumed less feed than birds fed diet supplemented with 500 U/kg PVA-coated-phytase and GA-coated-phytase, pelleted at 90°C (Table 8). Birds fed the mash diets supplemented with 500 U/kg Phyzyme® and PVA-coated-phytase and GA-coated-phytase showed no statistical difference with the positive control. Those fed diets supplemented with 500 U/kg PVA-coated-phytase and GA-coated-phytase, pelleted at 90°C, had better feed conversion ratio at day 21 than birds fed negative control + 500 U/kg Phyzyme® XP and positive control diets (Table 9). Tibia ash was lowest for birds fed negative control diet compared to all other treatments. Birds fed positive control and phytase-supplemented diets had similar tibia ash (Table 10).

[0115] The phytase tested, either as Phyzyme® XP or coated with PVA or GA, performed as well as the positive control diet when fed as mash. When PVA or GA-coated phytase was added to commercial diets pelleted at 90°C, recovery of the product was maintained at non-pelleted levels and the performance was at least as good as or better than the positive control diet. The results indicate that coating enzyme with PVA or GA improved its thermostability at pelleting temperature of 90°C without losing bioefficacy in broilers fed commercial diets.

Table 4 Dietary treatments

Dietary treatment	Enzyme, U/Kg	Inclusion (g/T)
Positive control	0, mash diet	0
Negative control	0, mash diet	0
Phyzyme XP Mash diet	500 U/kg Phyzyme XP, mash diet	100
PVA-coated Mash diet	500 U/kg PVA-coated phytase, mash diet	100
PVA coated pellet diet	500 U/kg PVA-coated phytase (Pellet at 90°C)	100
GA-coated Mash diet	500 U/kg GA-coated phytase, mash diet	100

Dietary treatment	Enzyme, U/Kg	Inclusion (g/T)
GA coated pellet diet	500 U/kg GA-coated phytase (Pellet at 40°C)	100

Table 5 Phytase activity in experimental diets

Dietary treatment	Expected	Observed	Observed, %
	FTU/Kg		
Positive control, Mash	<50	<50	-
Negative control, Mash	<50	<50	-
Phyzyme XP, Mash diet	500	563	112.6
PVA-coated, Mash diet	500	467	93.4
PVA coated Pelleted at 90 °C	500	558	111.6
GA-coated, Mash diet			
GA coated Pelleted at 90° C	500	440	88.0

Table 6 Experimental Diets

Ingredients: %	Positive control	Negative control
Maize	58.56	59.49
Soybean meal -48	34.65	34.55
Soya oil	2.82	2.48
salt	0.30	0.30
Sodium Bicarbonate	0.20	0.20
Dicalcium Phos.	1.59	0.83
Limestone	0.95	1.12
Vitamin premix	0.50	0.50
Lysine-HCl	0.10	0.10
DL-Methionine	0.23	0.23
Titanium dioxide	0.10	0.10
Enzyme premix carrier (corn)	0	0.10
<i>Nutrient Composition</i>		
Crude Protein, %	21.68	21.68
Poultry metabolisable energy, MJ/Kg	12.8	12.8
Calcium, %	0.90	0.78
Phosphorus, %	0.67	0.54
Available phosphorus, %	0.37	0.26
Methionine+Cystine, %	0.92	0.92
Methionine, %	0.56	0.56
Lysine, %	1.25	1.25
Threonine, %	0.82	0.82
Tryptophan, %	0.25	0.25

Table 7 Initial and Final Bodyweight of Broilers fed Experimental Diets

Dietary treatment	Bodyweight, g	
	Day 0	Day 21
Positive control, Mash	44.43	789b
Negative control Mash	43.55	680d
500 U/kg Phyzyme XP, Mash	43.72	753c
500 U/kg PVA-coated phytase, Mash	43.75	740c

Dietary treatment	Bodyweight, g	
500 U/kg GA-coated phytase, Mash		
500 U/kg PVA-coated phytase, Pelleted at 90 °C	43.70	8992
500 U/kg GA-coated phase, Pelleted at 90 °C	43.67	934a
Standard error of difference between means	0.261 (PVA)	6.015 (PVA)
	0.228 (GA)	7.741 (GA)
P value	0.165 (PVA)	<0.001 (PVA)
	0.053 (GA)	<0.001 (GA)
Differing superscripts in the same column indicate significant difference		

Table 8 Overall Feed Intake of Broilers fed Experimental Diets

Dietary treatment	Feed intake (g/bird/day)
Positive control Mash	56.69b
Negative control, Mash	46.00a
500 U/kg Phyzyme XP, Mash	59.03b)
500 U/kg PVA-coated phytase, Mash	56.70b
500 U/kg PVA-coated phytase, Pelleted at 90 °C	57.18b
500 U/kg GA-coated phytase, Pelleted at 90° C	64.33b
Standard error of difference between means	2.321 (PVA)
	2.260 (GA)
P value	0.004 (PVA)
	0.001 (GA)
Differing superscripts in the same column indicate significant difference	

Table 9 Feed Conversion Ratio of Broilers fed Experimental Diets

Dietary treatment	Feed Conversion Ratio
Positive control, Mash	1.617ab
Negative control, Mash	1.548ab
500 U/kg Phyzyme XP, Mash	1.752a
500 U/kg PVA-coated phytase, Mash	1.715a
500 U/kg PVA-coated phytase, Pelleted at 90°C	1.412b
500 U/kg GA-coated phytase, Pelleted at 90° C	1.537
Standard error of difference between means	0.715 (PVA)
	0.069 (GA)
P value	0.016 (PVA)
	0.061 (GA)
Differing superscripts in the same column indicate significant difference	

Table 10 Tibia Ash Content of Broilers fed Experimental Diets

Dietary treatment	Tibia ash content (g/100g)
Positive control, Mash	14.03b
Negative control, Mash	10.86a
500 U/kg Phyzyme XP, Mash	13.93b
500 U/kg PVA-coated phytase, Mash	13.24b
500 U/kg PVA-coated phytase, pelleted at 90 °C	13.50b

Dietary treatment	Tibia ash content (g/100g)
500 U/kg GA-coated phytase, pelleted at 90° C	13.84a
Standard error of difference between means	0.403 (PVA)
	0.302 (GA)
P value	<0.001 (PVA)
	<0.00 (GA)
Differing superscripts in the same column indicate significant difference	

REFERENCE EXAMPLE 6: ENZYME PREMIX & PELLETING STABILITY

[0116] Granules containing phytase, made according to formulations in Table 2 having either a PVA or a gum arabic coating and a moisture hydrating coating, were mixed with the clay sepiolite, or a standard broiler vitamin mineral premix with and without choline chloride. Blending ratios were 100 grams of granules added to 900 grams of clay (sepiolite), or 100 grams of granules added to 500 grams of vitamin mineral premix. Samples were stored in sealed containers at 35°C for 3 weeks, and then subjected to pelleting, as described above in Mill #3. The experimental control was granules that were not stored in a premix, and were held at ambient conditions for 3 weeks.

[0117] Tables 11 and 12 show the percent-recovered phytase activity of these mixtures after pelleting at 90°C and 95°C. Granules mixed with sepiolite or the vitamin mineral premixes were found to have significantly increased recovered activity after pelleting.

Table 11				
PVA Formulation Granules				
Premix	Percent recovered activity after pelleting, relative to mash		Percent increase in recovered activity, relative to control granules	
	90°C	95°C	90°C	95°C
granule alone (control)	71%	57%		
Sepiolite + granules	95%	88%	34%	53%
Vitamin mineral premix, no choline chloride + granules	93%	80%	32%	40%
Vitamin mineral premix, with choline chloride + granules	98%	85%	39%	48%
Table 12				
GA Formulation Granules				
Premix	Percent recovered activity after pelleting, relative to mash		Percent increase in recovered activity, relative to control granules	
	90°C	95°C	90°C	95°C
granules alone (control)	92%	81%		
Sepiolite + granules	99%	88%	8%	9%
Vitamin mineral premix, no choline chloride + granules	100%	81%	8%	0%
Vitamin mineral premix, with choline chloride + granules	97%	80%	5%	-2%

[0118] Without wishing to be bound by any particular theory, it is believed that the clay and the vitamin mineral premix have a capacity to take up water and, during storage with stable granules, they absorb residual moisture from the granules. To illustrate this effect, Table 13 shows the results of an experiment in which granules and sepiolite are stored in open containers, positioned side-by-side, within a closed chamber. The water activity of the granules, and of the sepiolite, was measured before and after 7 days of storage at 25 °C. During storage, the sepiolite absorber water from the granules until the system reached an equilibrium.

After storage, the granules show a decrease in water activity, while the sepiolite shows an increase in water activity.

Table 13				
	Initial water activity		Water activity after 7 days of storage together	
	Sepiolite	granules	Sepiolite	granules
Formulation GA granule	0.312	0.558	0.359	0.373
Formulation PVA granule	0.312	0.516	0.354	0.365

REFERENCE EXAMPLE 7: STORAGE STABILITY WITH CHOLINE CHLORIDE

[0119] Choline chloride, or N-(2-Hydroxyethyl) trimethylammonium chloride, is an important feed additive, a vitamin nutrient, in poultry, pig and other animal feeds. Choline chloride is a reactive molecule and has a well-known destructive effect on other vitamins, and enzymes. Choline chloride is often included in premixes and base mixes. Maximum levels used in premixes are 74,800 mg/kg for swine, and 150,000 mg/kg for poultry. And typical levels for swine base mixes are about 966 to 1282.9 mg/kg.

[0120] The stable, durable granules of the present invention were shown to preserve enzyme activity when stored in the presence of choline chloride. Granules containing phytase, made with formulations with either PVA or gum arabic and a moisture hydrating material were mixed with a standard broiler vitamin mineral premix, with and without choline chloride. The blending ratio was 100 grams granules added to 500 grams of vitamin mineral premix. Samples were stored in sealed containers, at 35 °C, for 3 weeks, and then assayed for activity, using the phytase assay protocol described above. The experiment control was granules that were not stored in a premix, and were held at 35 °C for 3 weeks. Tables 14 and 15 show the measured activity of the mixtures before and after storage, and the percent change in activity. None of the samples show an appreciable loss of activity after storage. The error of this assay, including error of sampling, extraction, and activity assay is approximately 15%.

Table 14			
PVA Formulation Granule			
Sample	Initial activity (FTU/g)	Activity after storage 3 weeks at 35°C (FTU/g)	percent change in activity
granules only (control)	11,600	11,858	2%
granules + vitamin mineral premix without choline chloride	1,933	1,727	-12%
granules + vitamin mineral premix with choline chloride	1,933	1,720	-12%

Table 15			
GA Formulation Granule			
Sample	Initial activity (FTU/g)	Activity after storage 3 weeks at 35 °C (FTU/g)	percent change in activity
granules only (control)	10,373	10,528	1%
granules + vitamin mineral premix without choline chloride	1,729	1,543	-12%
granules + vitamin mineral premix with choline chloride	1,729	1,622	-7%

EXAMPLE 8: WATER ACTIVITY EFFECT ON PELLETING STABILITY

[0121] Phytase granules were prepared according to a formulation having a PVA coating and/or an inorganic salt layer coating, in a fluid bed process, as described above. An additional drying step can be used if the water activity of the granule is greater than 0.5 after processing, so that in a fluid bed coater, or other suitable process; the granules may be dried until a water activity of <0.5 is achieved. The results are shown in Table 16 and demonstrate that retained activity after pelleting is enhanced when the water activity is less than 0.5.

Table 16					
PVA Formulation Granule					
Granule	Water activity	Percent recovered activity after pelleting, relative to mash		Percent increase in recovered activity, relative to control granules	
		90°C	95°C	90°C	95°C
A (control)	0.57	81%	69%		
B	0.41	98%	87%	21%	26%
C	0.49	110%	97%	36%	41%

REFERENCES CITED IN THE DESCRIPTION

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STABILE, HOLDBARE GRANULATER MED AKTIVE STOFFER

PATENTKRAV

1. Dyrefodergranulat, hvilket granulat omfatter:
 - en kerne;
 - 5 et aktivt stof;
 - og mindst en belægning omfattende et fugthydrerende materiale, der er et uorganisk salt, som er i stand til at optage vandige væsker ved at absorbere frit vand eller ved at optage vand som krystallinsk hydreringsvand, hvilket fugthydrerende materiale udgør mindst 25 % vægt/vægt af granulatet,
 - hvor granulatets vandaktivitet er mindre end 0,5,
 - 10 og hvor granulatets aktive stof bevarer mindst 50 % aktivitet efter betingelser udvalgt fra én eller flere af:
 - a) en foderpelletteringsproces,
 - b) en dampopvarmet foderforbehandlingsproces,
 - c) oplagring,
 - 15 d) oplagring som en ingrediens i en ikke-pelletteret blanding og
 - e) oplagring som en ingrediens i en foderbaseblanding eller en foder-præmix, der mindst omfatter én forbindelse, der er udvalgt fra spormineraler, organiske syrer, reducerende sukker, vitaminer, cholinchlorid og forbindelser, der resulterer i en sur eller basisk foderbaseblanding eller et foder-præmix.
2. Granulat ifølge krav 1, hvor det fugthydrerende materiale er natriumsulfat.
- 20 3. Granulat ifølge et hvilket som helst af kravene 1-2, hvor det fugthydrerende materiale udgør mindst 55 % vægt/vægt af granulatet.
4. Granulat ifølge krav 1, hvor det aktive stof er én eller flere enzymer.
5. Granulat ifølge krav 4, hvor det aktive stof er en Citrobacter-phytase.
6. Granulat ifølge krav 1, hvor granulatets aktive stof bevarer mindst 80 % aktivitet efter en
 - 25 dampopvarmet pelletteringsproces, der udføres ved mellem 85 °C og 95 °C i op til adskillige minutter, hvor granulatet er en ingrediens.
7. Granulat ifølge krav 1, hvor granulatet er blandet med et præmix omfattende cholinchlorid.
8. Dyrefodersammensætning omfattende granulatet ifølge et hvilket som helst af de foregående krav.
9. Dyrefodersammensætning ifølge krav 8, hvilket foder er udvalgt fra et
 - 30 dyrefoderfortyndingsmiddel, en dyrefoderbaseblanding, et dyrefoder-præmix, en dyrefoderblanding og en dyrefoderpellet.
10. Dampbehandlet pelletteret fodersammensætning omfattende granulater ifølge et hvilket som helst af kravene 1-7.