

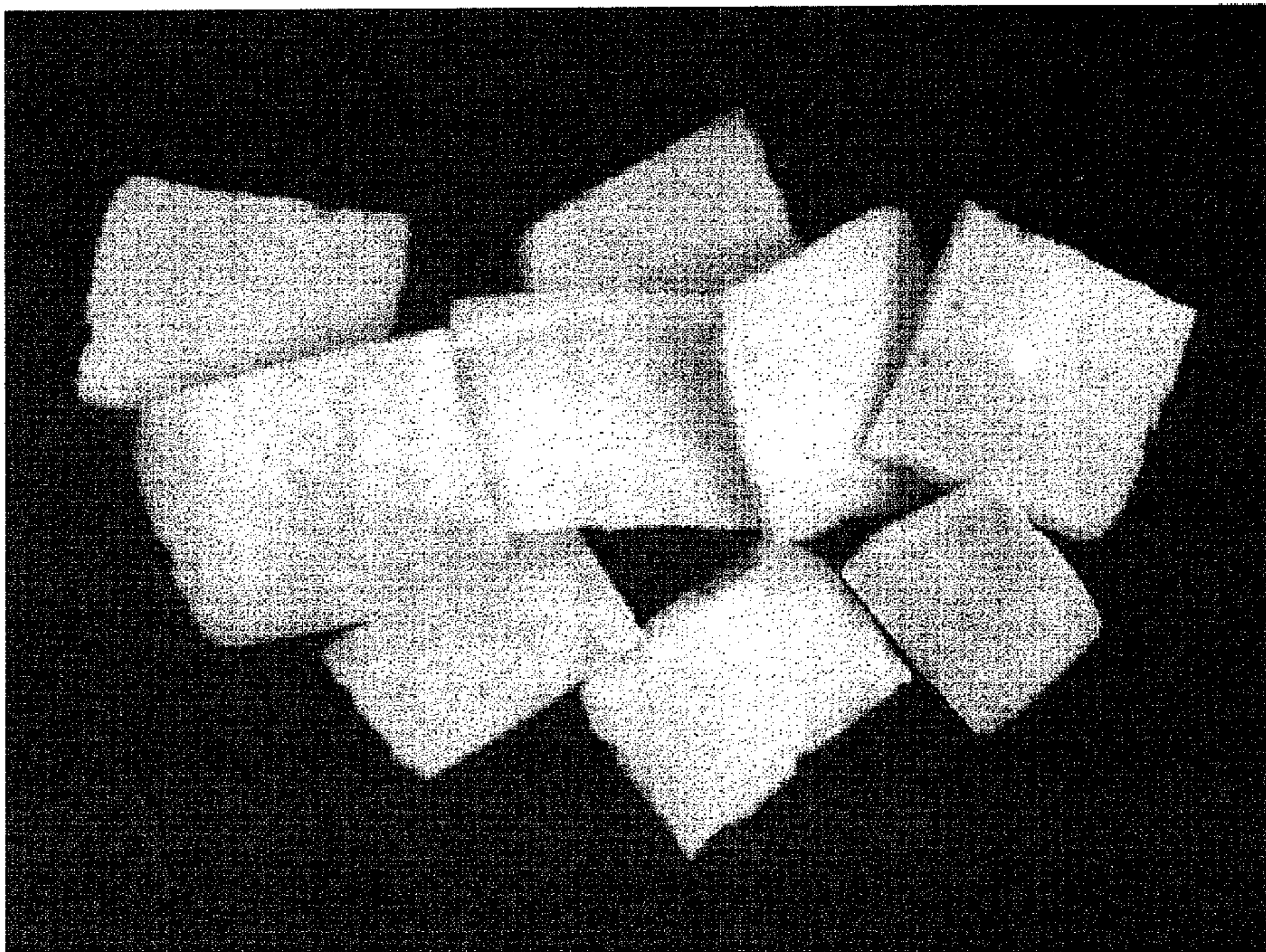


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(54) Title: A DRY FOOD PRODUCT CONTAINING LIVE PROBIOTIC

Examples of Probiotic Chips Produced as Described Herein



(57) Abrégé/Abstract:

The disclosure relates to a probiotic delivery system that can be consumed as a snack-food or added to a food product. In particular, the disclosure describes a crisp and tasty treat that comprises viable probiotic microorganisms preserved in a vacuum dried matrix of sugars, proteins, and polysaccharides. The probiotic remain viable within the treat for a longer time without the need for additional moisture barrier coating. The probiotic also remain viable in the animal gastrointestinal tract.

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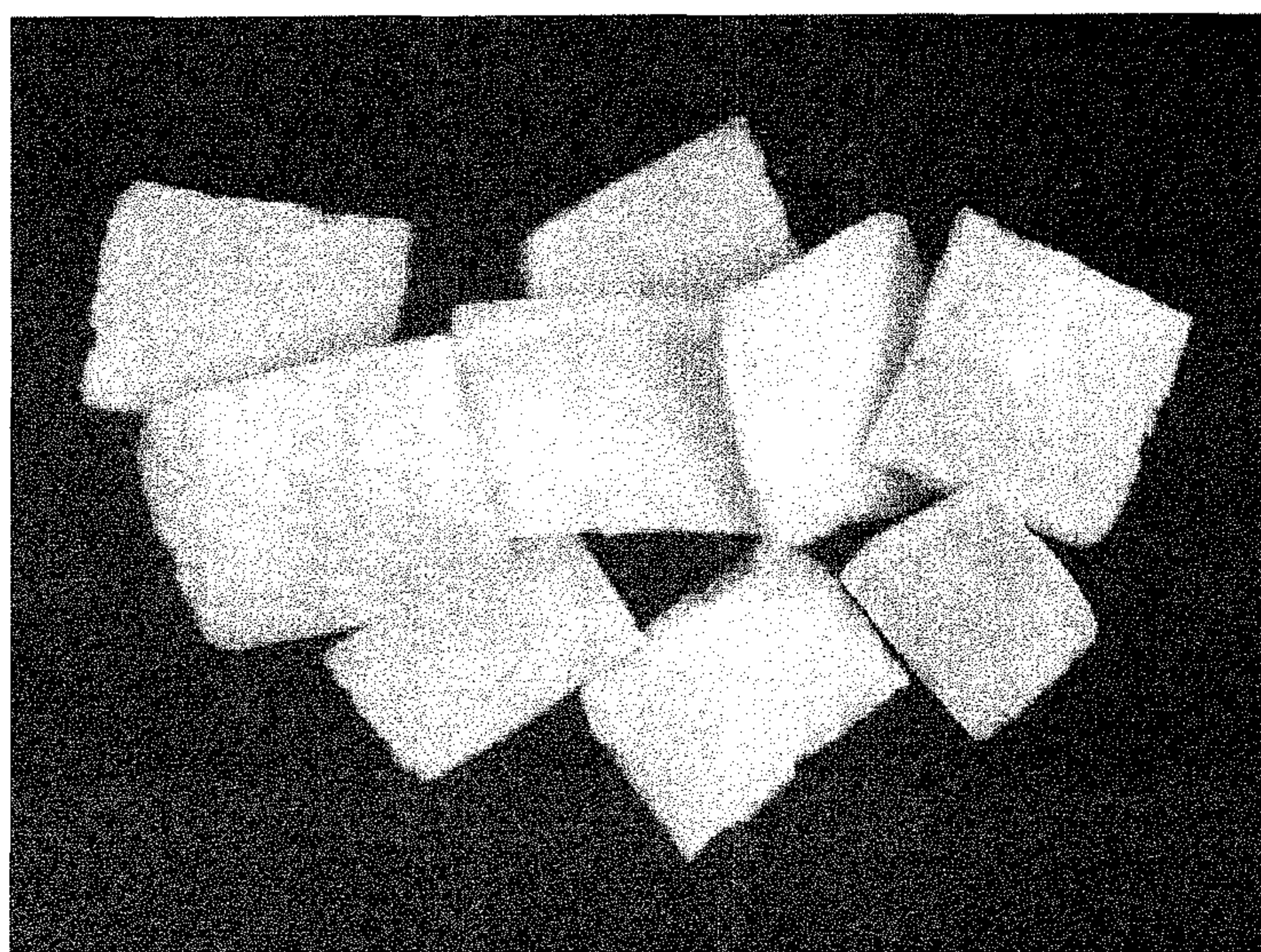
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(57) **Abstract:** The disclosure relates to a probiotic delivery system that can be consumed as a snack-food or added to a food product. In particular, the disclosure describes a crisp and tasty treat that comprises viable probiotic microorganisms preserved in a vacuum dried matrix of sugars, proteins, and polysaccharides. The probiotic remain viable within the treat for a longer time without the need for additional moisture barrier coating. The probiotic also remain viable in the animal gastrointestinal tract.

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TITLE OF THE DISCLOSURE

[0001] A Dry Food Product Containing Live Probiotic

BACKGROUND OF THE DISCLOSURE

5 [0002] The disclosure relates generally to the fields of probiotics and food.

[0003] The activity and long term stability of many biological materials, such as proteins, enzymes and microbial cells may be affected by a number of environmental factors; for example, temperature, pH, the presence of water and oxygen or oxidizing or reducing agents. Generally, biological materials must be dried before or during mixing with other foodstuff
10 ingredients. The drying process can often result in a significant loss in activity from mechanical, chemical, and osmotic stresses induced by the drying process. Loss of activity occurs at many distinct stages, including drying during initial manufacturing, feed preparation (high temperature and high pressure), transportation and long term storage (temperature and humid exposure), and after consumption and passage in the gastrointestinal (GI) track
15 (exposure to low pH, proteolytic enzymes and bile salts). Manufacturing food or feedstuffs with live cell organisms or probiotics is in particular challenging, because the probiotics are very sensitive to the drying process and to temperature and moisture conditions of the foodstuff. Another concern is the probiotic resistance in the acid environment in the stomach and its successful colonization of the intestine.

20 [0004] Probiotic microorganisms (probiotics) are living microorganisms, which upon ingestion in certain numbers, exert health benefits beyond basic nutrition. The beneficial effects that probiotics may induce are numerous. Few examples are; the reduction of lactose intolerance, the inhibition of pathogenic bacteria and parasites, the reduction of diarrhea, activity against *Helicobacter pylori*, the prevention of colon cancer, the improvement or
25 prevention of constipation, the in situ production of vitamins, the modulation of blood lipids, and the modulation of host immune functions. In domesticated and aquatic animals they also can improve growth, survival and stress resistance associated with diseases and unfavorable culture conditions. Therefore, there is considerable interest in including probiotics into human foodstuffs and into animal feed.

[0005] Many probiotics exhibit their beneficial effect mainly when they are alive. Hence, they need to survive the manufacturing process and shelf life of the food, and upon consumption of the food where they need to pass through the gastro-intestinal tract before reaching their place of colonization. Although many commercial probiotic products are available for animal and human consumptions, most of them lost their viability during the manufacture process, transport, storage and in the animal GI tract (see the viability studies of several probiotic products by (Hughes and Hillier 1990; Shah 2000). To compensate for such loss, an excessive quantity of probiotics is included in the product in anticipation that a portion will survive and reach their target. In addition to questionable shelf- life viability for these products, such practices are certainly not cost-effective. Alternatively, the probiotic microorganisms can be encapsulated in protective microenvironments. Generally, current microencapsulation and enteric coating techniques involve applying a film forming substance, usually by spraying liquids containing sugars or proteins onto the dry probiotics (Ko and Ping WO 02/058735). However, coating the microencapsulated probiotics with moisture protecting layers is an expensive process, and generally several layers must be added, to avoid water entering the microcapsules. In addition, it is extremely difficult to remove the added liquid in the coating substance without a corresponding decrease in shelf life.

[0006] Various protective agents have been used in the art, with varying degrees of success. These include proteins, certain polymers, skim milk, glycerol, polysaccharides, oligosaccharides and disaccharides. Disaccharides, such as sucrose and trehalose, are particularly attractive cryoprotectants because they are actually help plants and microbial cells to remain in a state of suspended animation during periods of drought. Trehalose has been shown to be an effective protectant for a variety of biological materials, both in ambient air-drying and freeze-drying (Crowe et al. 1998). However, there are some drawbacks associated with the use of sugars as the sole cryoprotectant. For example, large amounts of sugars (often greater than 60% by weight) must be used to preserve the biological materials during the drying process. This is costly. More serious problems associated with the use of sugars include their readiness to form crystals when the material is dried below its freezing point, and the low glass transition temperature which causes instability of the preserved biological materials at high temperatures, and/or in humid environments. Further, high concentration of sugars reduces the

solubility of other solutes in the system and at the same time renders the system extremely difficult to dry.

[0007] Accordingly, it has been proposed to dry sugar-based probiotic systems by foam formation in a very thin layer (Bronshtein WO2005117962), or to use combinations of sugars with a polymeric gelling agent, such as alginate, chitosan, carboxymethylcellulose or carboxyethylcellulose. Cavadini et al. (EP 0 862 863) provide a cereal product comprising a gelatinized starch matrix including a coating or a filling. The probiotic is included with the coating. According to that process, spray-dried probiotics are mixed with a carrier substrate, which may be water, fat or a protein digest. The mixture is then sprayed onto the cereal product and the whole product is dried again. Re-hydrating of the already dried bacteria and the additional coating/drying process is costly and damaging to the bacteria.

[0008] Kenneth and Liegh (US 6900173) describe the manufacturing of multivitamin protein and probiotic bar for promoting an anabolic state in a person. The dried probiotic bacteria are blended in sugar syrup and several other constituents, and the resultant mixture is then extruded and cut into bars. However, the document does not disclose any process or composition that will improve viability or long-term stability of probiotics in the nutritional bars and there is no indication that the bacteria even survive the process.

[0009] Ubbink et al. (US 2005/0153018) disclose the preservation of lactic acid bacteria in moist food. The spray-dried bacteria are added to a composition comprising fats, fermented milk powder and saccharides. That composition is then used as the filling of a confectionary product. The subject matter described in that document avoids the detrimental effects of water by embedding the probiotics in fat or oil rich matrix. However, fat based coating and preserving materials do not withstand long term exposure to humid conditions

[0010] Giffard and Kendall (US 2005/0079244) disclose a foodstuff in the form of a dried or semi-moist ready-to-eat kibble or powder mix, which contains a combination of a probiotic, prebiotic and a coating of colostrum. Prior to mixing in the food stuff, the probiotic is coated or encapsulated in a polysaccharide, fat, starch, protein or in a sugar matrix using standard encapsulation techniques. Similar to the above disclosure, the negative effects of water were avoided by embedding the probiotics in a matrix rich in fat or oil.

[0011] Farber and Farber (WO 03/088755) describe an oral delivery system for functional ingredients uniformly dispersed in a matrix. The matrix components include a sugar, a carbohydrate, a hydrocolloid a polyhydric alcohol and a source of mono-or divalent cations. The delivery system is extruded or molded into a final shape with a moisture content of
5 between 15% and 30% by weight. This type of matrix provides very little protection to the probiotics mostly under refrigerated conditions. No description or direction was provided as to how probiotic bacteria are stabilized during manufacturing or for prolonged storage at room temperatures.

[0012] Porubcan (US 2004/0175389) discloses a formulation for protecting probiotic
10 bacteria during passage through the stomach, whilst permitting their release in the intestine. The formulation has also a low water activity and correspondingly long shelf life. The capsule includes a water-free mixture of probiotic bacteria with monovalent alginate salts, and an enteric coating (e.g., gelatin or cellulose encapsulation). Upon contact with acidic environment, the outer shell of the capsule turned into a gel, which provides a protecting barrier against
15 proton influx into the capsule core. However, this composition is only useful for large particles such as tablets and capsules subjected to storage conditions of very low water activity and further requires storage in nitrogen-flushed or vacuum-sealed containers.

[0013] McGrath and Mchale (EP 1382241) describe a method of delivering a micro-organism to an animal. The micro-organism is suspended in a matrix of cross-linked alginate
20 and cryopreservant (trehalose or lactose, or a combination of both). The matrix is then freeze or vacuum dried to form dry beads containing live probiotics with a shelf-life stability up to 6 months but only under refrigerated conditions. Here again, no description or direction was provided as to how probiotic bacteria are stabilized during manufacturing or for prolonged storage at room temperatures and high humidity conditions.

[0014] None of the above compositions provide a mixture that can effectively protect the
25 probiotic in both drying processes and long-term storage at elevated temperatures and varying degrees of humidity. Therefore, there is an urgent need for such a composition that can effectively protect the probiotic bacteria during manufacturing, long-term storage at elevated temperatures and humidity and during gastrointestinal passage. There is a need also for a
30 drying process that is cost-effective and capable of entrapping and stabilizing probiotics in the

protective mixture with minimal viability loss at the end of the entire operation. There is a need for a protective mixture that provides protection in the animal stomach while allowing the release of the probiotic along the intestinal tract. There is also a need for a protective mixture that contains only approved ingredients generally regarded as safe (GRAS), and is less costly than those presently being used.

[0015] The subject matter described herein overcomes these needs and provides a composition and process for producing a composition that provides probiotic bacteria that are stable for long periods of time even at elevated temperatures and varying degrees of humidity.

10 [0016] It is, in particular, a purpose of the present disclosure to describe viable probiotic cultures that are substantially stable at room temperature and high humidity conditions thereby obviate the need for refrigeration or storage under vacuum or oxygen free environment.

15 BRIEF SUMMARY OF THE DISCLOSURE

[0016a] In one particular embodiment there is provided a probiotic substance comprising viable probiotic bacteria preserved in a dried matrix of at least one sugar wherein the total amount of sugar compound in the matrix is from about 10% to about 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix.

[0016b] In another particular embodiment there is provided a human food product comprising one or several probiotic substances comprising viable probiotic bacteria preserved in a dried matrix comprising at least one sugar, wherein the total amount of sugar compound in the matrix is from about 10% to about 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix, wherein the dried matrix has water activity below 0.2.

[0016c] In yet another particular embodiment there is provided an animal feed product comprising one or several probiotic substances comprising viable probiotic bacteria preserved in a dried matrix comprising at least one sugar wherein the total amount of sugar compound in the matrix is from about 10% to about 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix.

[0016d] In still yet another particular embodiment there is provided a method of making probiotic substance or food product according to any of the above claims, the method comprising: a) mixing a preparation of probiotic micro-organisms with at least one sugar, at least one protein, and at least one polysaccharide to obtain a smooth and uniform slurry or gel in a desired shape and form; and b) vacuum drying the preformed solid gel to a water activity below 0.1, wherein the vacuum drying comprises: setting the temperature of the above the freezing temperature of the probiotic substance; and vacuum drying at reduced pressure of about 5000 mTOR at a shelf temperature of from 5-50°C and then at a pressure of about 100 mTOR while maintaining the shelf temperature.

[0017] It was unexpectedly found that probiotic bacteria are protected for an extended period of time in high temperature and humid conditions when preserved in a certain protective mixture. Additional qualities of the protecting mixture are, a fast and cost effective drying process and gastric protection. The mixture comprises: (a) at least one sugar compound, where the total amount of sugar compound in the mixture is from about 10% to about 60% by weight of the mixture (b) proteins, where the total amount of proteins in the medium is from about 2% to about 20% by weight of the mixture and (c) polysaccharides, where the total amount of polysaccharides in the medium is from about 0.5% to about 5% by weight of the mixture. This aqueous protective mixture can be used in a multiplicity of preservation processes, including freezing, freeze-drying, spray-drying, vacuum-drying, or ambient air-drying, to provide a stable and preserved composition of probiotics. The probiotic substance is stable for extended periods of time at superambient temperatures and/or relative humidity. Further, the aqueous protective

mixture containing the probiotics can be molded into a desirable shape or form and vacuum-dried to produce a crisp and tasty probiotic treat that can be added to food stuff or consumed on its own by humans or animals.

[0018] Therefore, the present disclosure also describes a method for preparing a
5 preserved probiotic substance containing the above-noted protective mixture.

[0019] Preferably, the probiotic substance is provided in a dry form that is substantially free of water. The probiotic substance may be freeze-dried, vacuum dried or air dried, or otherwise dried by methods known in the art. Accordingly, the probiotic substance preferably comprises a protective mixture capable of maintaining the viability of the probiotic micro-organisms for
5 extended periods of time in ambient temperature and humidity conditions.

[0020] Preferably, the sugar in the protective mixture is a disaccharide, most preferably trehalose or sucrose or lactose or a combination thereof. The protective mixture preferably comprises trehalose at 20% w/v to 60% w/v trehalose, preferably 20%, 30% or 40%.

[0021] Preferably, the protein in the protective mixture of the subject matter described
10 herein is egg albumen or soy protein isolate or hydrolysate and a mixture thereof. The protective mixture preferably comprises proteins at 2% w/v to 20% w/v proteins, preferably 5%, 10% or 20%.

[0022] Preferably, the polysaccharides in the protective mixture described herein can form a firm gel or viscous solution with the other ingredients in the mixture, most preferably a
15 combination of alginates with different viscosities, agarose, pectin or chitosan. The protective mixture preferably comprises alginates at 0.5% w/v to 10% w/v alginates, preferably 1%, 2% or 4%.

[0023] In accordance with the subject matter described herein, there is provided the use of the probiotic substance described herein for the manufacture of a probiotic product or a
20 probiotic food or feed product for the consumption by humans, domestic animals, aquatic animals and pets.

[0024] Remarkably, it was found that by adding a mixture containing 30% trehalose (w/v), 10% soy protein isolate (w/v) and 2% sodium alginate to a probiotic bacteria concentrate, forming viscous solution or hydrogel and vacuum drying it at temperature above the freezing
25 point of the mixture an excellent process recovery, prolonged stability over storage time in ambient conditions and gastric protection are obtained.

[0025] Optionally, the probiotic substance may be coated with a moisture barrier component. In principle, any food-grade substance having water repelling or impermeable properties may be selected. Suitable moisture barriers may be, for example, a mixture of oil
30 based substances.

[0026] Consequently, in a first aspect, the subject matter described herein includes a probiotic product comprising a dry micro-matrix particle of the above mixture, wherein the micro-matrix particle comprises viable microorganisms and has a size between 10 and 2000 microns.

5 [0027] In a second aspect, the subject matter described herein includes a food product containing probiotic flakes or treats, wherein the flakes or treats are the probiotic substance described herein, characterized in that the flake or treat has a desirable shape and size between 2-50 millimeters.

[0028] In a third aspect, the subject matter described herein includes a process for obtaining
10 micro-matrix particles, to supplement a food product with viable micro-organisms. The process comprises the steps of mixing micro-organisms concentrate and further protective components, forming viscous solution or hydrogel, drying the mixture by freeze-drying, spray-drying, vacuum-drying, or ambient air-drying, and, if necessary, grinding the dry material to obtain micro-matrix particles comprising a size between 50-2000 microns.

15 [0029] In a fourth aspect, the subject matter described herein includes a process for obtaining flakes or treats as supplement or stand-alone food product with viable micro-organisms. The process comprises the steps of mixing micro-organisms concentrate and further protective components, Forming a hydrogel in a desirable shape and size and drying by vacuum-drying, to obtain crispy and tasty flakes or treats comprising a size between 2-50
20 millimeters.

[0030] One major advantage of the subject matter described herein is that it provides a significant improvement over other drying methods of sugar based substances and production methods of stable probiotic micro-organisms in semi-dry and/or humid particulate foodstuffs.

[0031] Another advantage of the subject matter described herein is that the drying process
25 is easy to upscale and straightforward with no need of additional coating or several drying stages.

[0032] Yet another advantage of the subject matter described herein is that the probiotic substance provides gastric protection and a release mechanism of the probiotics along the intestinal tract at their site of action.

[0033] Yet another advantage of the subject matter described herein is that it provides a suitable delivery vehicle for further functional ingredients, in particular non-digestible sugars, natural proteins and prebiotic fibers, which in turn may improve the physico-chemical characteristics of the probiotic substance, as described herein.

5

BRIEF SUMMARY OF THE SEVERAL VIEWS OF THE DRAWINGS

[0034] Figure 1 is a table that shows drying loss after vacuum drying of *L. rhamnosus* probiotic in the MicroMatrix product described in Example 1.

[0035] Figure 2 is a graph that illustrates storage stability (recovered cfu/g {colony-forming units per gram} versus storage time) of dry *L. rhamnosus* probiotic substance stored at 40° C and 33% relative humidity.

[0036] Figure 3 is a bar chart that illustrates gastric stability (recovered cfu/g) of probiotic substances incubated in simulated gastric juice (pH=1.2) at 37° C for 2 hours.

[0037] Figure 4 is an image of examples of probiotic “chips” produced as described herein.

15

DETAILED DESCRIPTION

[0038] The disclosure relates to food products including a probiotic component.

[0039] The present disclosure relates to a food or feed product comprising viable microorganisms, a mixture that protects probiotics against high temperature, humidity and low pH, the use of the probiotic substance in a food product and a process for obtaining a probiotic substance to supplement food or feed products. The disclosure further relates to a food product comprising the probiotic substance.

[0040] The subject matter described herein relates generally to a composition for preserving probiotic microorganisms, and to the production and drying methods of the substance. More specifically, the subject matter described herein includes a dry probiotic substance with long-term shelf life under high temperature and humid conditions.

[0041] Definitions

[0042] As used herein, each of the following terms has the meaning associated with it in this section.

30

[0043] The term "food product" is intended to encompass any consumable matter of either plant or animal origin or of synthetic sources that contain a body of nutrients such as a carbohydrate, protein, fat vitamin, mineral, etc. The product is intended for the consumption by humans or by animals, such as domesticated animals, for example cattle, horses, pigs, sheep, goats, and the like. Pets such as dogs, cats, rabbits, guinea pigs, mice, rats, birds (for example chickens or parrots), reptiles and fish (for example salmon, tilapia or goldfish) and crustaceans (for example shrimp). Preferable, the subject matter described herein includes standard food products pelleted feeds, and pet food (for example a snack bar, crunchy treat, cereal bar, snack, biscuit, pet chew, pet food, and pelleted or flaked feed for aquatic animals).

[0044] The word "probiotic" is intended to refer to any consumable microorganism owing to any beneficial effect it may have on its consumer.

[0045] The term "probiotic substance" is a dry consumable substance in any shape or form that contains probiotics. More specifically, a probiotic substance comprises live probiotics embedded in a matrix of sugars, proteins and polysaccharides. Hence, it may be a food product on its own.

[0046] The term "micro-matrix" particles may assume any dry powder form of the probiotic substance. The micro-matrix particles may serve as a carrier for the probiotics and comprise a size from 10 micron up to 2000 micron.

[0047] The term "flake or treat" is not intended to refer to specific form or shape of the probiotic substance. A flake or treat may assume any form obtained by molding shaping or slicing a hydrogel. For example, a flake or treat may have the form of a sphere, cube, pyramid, tablet, cereal or any complex three-dimensional form that comprise a size of at least 2 millimeters. For example, if the treats are used as a probiotic delivery system for pet-food, they may have the form of bones, rods, rings or other kibble forms.

[0048] The word "hydrogel" may refer to any moist food-grade substance that has the property of a solid or viscous gel. A hydrogel, either anionic or cationic, can be formed by one or more hydrophilic polymers, polysaccharides, gums, resins, or hydrolyzed proteins, either alone or in combination, in which the microorganisms are disposed. Preferably, the hydrogel compounds include agarose, alginate, chitosan or any other compound, which preferably can present characteristics of a solid gel.

[0049] Detailed Description

[0050] In order to prepare the probiotic substance as described herein, a single or a mixture of several micro-organisms may be selected. As a probiotic micro-organism, any micro-organism may be selected. Preferably, a micro-organism exerting beneficial effects on health and welfare on humans or animals is used.

[0051] Examples of suitable probiotic micro-organisms include yeasts such as *Saccharomyces cerevisiae*, molds such as *Aspergillus*, *Rhizopus*, *Mucor*, and *Penicillium*, bacteria such as the genera *Bifidobacterium*, *Clostridium*, *Bacillus* and *Lactobacillus*. Specific examples of suitable probiotic micro-organisms are: *Aspergillus niger*, *A. oryzae*, *Bacillus coagulans*, *B. lentus*, *B. licheniformis*, *B. mesentericus*, *B. pumilus*, *B. subtilis*, *B. natto*, *Bifidobacterium adolescentis*, *B. animalis*, *B. breve*, *B. bifidum*, *B. infantis*, *B. lactis*, *B. longum*, *B. pseudolongum*, *B. thermophilum*, *Candida pintolepesii*, *Clostridium butyricum*, *Enterococcus cremoris*, *E. diacetylactis*, *E. faecium*, *E. intermedius*, *E. lactis*, *E. muntii*, *E. thermophilus*, *Lactobacillus acidophilus*, *L. alimentarius*, *L. amylovorus*, *L. crispatus*, *L. brevis*, *L. casei*, *L. curvatus*, *L. cellobiosus*, *L. delbrueckii* ss. *bulgaricus*, *L. farciminis*, *L. fermentum*, *L. gasseri*, *L. helveticus*, *L. lactis*, *L. plantarum*, *L. johnsonii*, *L. reuteri*, *L. rhamnosus*, *L. sakei*, and *L. salivarius*.

[0052] The probiotic micro-organisms are preferably mixed in a concentrated wet paste form or frozen paste form (for example a probiotic paste of >10% solids) with the other protective substances. Microorganisms may also be mixed, directly after fermentation, with the protective components described herein followed by hydrogel formation and a drying process thereafter. For example, probiotic micro-organisms are mixed with the protective materials such as a saccharide, for example trehalose, sucrose, lactose or maltodextrin, a protein, for example egg albumen, soy protein isolate or hydrolysate either alone or in combination and a polysaccharide, for example, agarose, alginate or chitosan either alone or in combination. A hydrogel is then formed in a desired shape and size or sliced after hardening the gel according to established procedures known to persons skilled in the art. If micro-matrix particles are required, then the hydrogel can be sliced or extruded and then dried using a variety of drying techniques, for example fluidized bed drying, freeze drying, air drying, convention oven drying

or another adequate drying process. The dry probiotic substance is then ground and sieved to preferred sizes. If flakes or treats are required, then the molded or otherwise pre-shaped or sliced hydrogel is preferably dried in a vacuum drier or freeze drier at a temperature above the freezing point of the hydrogel. The pre-shaped dried flake or treat is then ready for packaging alone or in combination with other food products.

[0053] Preferably, the probiotic substance comprises significant amounts of the protective composition, in which the micro-organisms are embedded. Preferably, the probiotic substance comprises, in percent by weight of total dry matter, 1-50%, preferably 5-25%, more preferably 10-20% of probiotic microorganisms in the protective composition.

[0054] In one embodiment, the probiotic substance described herein comprises 10^6 to 10^{12} viable micro-organisms (cfu) per gram dry weight. Preferably, it comprises 10^7 to 10^{11} cfu per gram dry weight.

[0055] In another embodiment, the dried probiotic substance is characterized by a water activity below 0.2. Preferably, the water activity is below 0.1, for example, the water activity is in the range of 0.01 to 0.09.

[0056] The probiotic substance in accordance with a preferred embodiment comprises of sugar, proteins and polysaccharides. Wherein the protein is selected from natural proteins including albumen, arginine/lysine polypeptide, collagen and hydrolyzed collagen, gelatin and hydrolyzed gelatin, glycoproteins, milk protein, casein, whey protein, serum albumin, meat, fish, seafood, poultry, egg proteins, silk, soybean, corn, peanut, cottonseed, sunflower, pea, wheat protein, wheat germ protein, gluten-protein, zein and any isolate or hydrolyzed of any vegetable protein, and the like.

[0057] Preferably, the polysaccharide components of the probiotic substance may be selected in a way that a formation of a solid gel is possible. Generally, this may be achieved by cross linking the polysaccharide (for example by mixing divalent cations with alginate or by cooling the gel (for example agarose).

[0058] Additional functional ingredients may be selected to provide further benefits to the probiotic substance described herein. For example fructo-oligosaccharides (FOS) and polyfructoses, for example, inulin, pectin, 6-glucans, resistant starches, for example high amylose starch, dextrans, acacia gum, guar and locust bean gum, agar, carrageenans, xanthan

and maltodextrins, and mixtures thereof. Additional functional ingredients may also comprise trace elements, minerals, vitamins, antioxidants, sterols, antioxidants and/or other functional molecules. However, the effect of any additional components on the protective characters of the probiotic substance should be evaluated first. Examples of vitamins and/or antioxidants are
5 carotenoids, such as lycopene, beta-carotene, lutein, xanthophylls, vitamin A, tocopherols, vitamin C, and mixtures thereof.

[0059] Since one of the objectives of the compositions and processes described herein is to add the probiotic substance to a food product, it is an advantage that the probiotic substance may be shaped like the desired food product. For example, if the probiotic substance is to be
10 added to a pet food, the probiotic substance may be shaped like a pellet, kibble or bone. Accordingly, if the probiotic substance is added to breakfast-cereals, it may be shaped like cereals. Or, the probiotic substance may be added to a snack as chips. Additionally, the probiotic substance may be added with flavors that used to prepare the food product.

[0060] In one embodiment, the probiotic substance may be coated with a moisture barrier
15 component. In principle, any food-grade substance having water repelling or impermeable properties may be selected. Suitable moisture barriers may be, for example, waxes (paraffin wax, beeswax, carnauba wax, candellila wax, microcrystalline wax, rice bran wax, shellac, lanolin, hydrogenated castor oil, jojoba oil), fatty acids (for example, oleic acid, stearic acid, palmitic acid, lauric acid), monoglycerides, diglycerides and triglycerides (for example, MCT
20 oil, triglycerides based on coconut/palm kernel oil), vegetable oils and fats (for example, rapeseed, sesame, cornseed, nut, cottonseed, peanut, sunflower, linseed, olive, soy bean, cocoa butter), hydrogenated or hardened vegetable oils and fats, oils and fats of animal origin (for example, beef, poultry, pork, for example, beef tallow, lard and fish oil), hydrogenated or hardened oils and fats of animal origin, dairy fats (for example, milkfat, butterfat), proteins (for
25 example, gluten, zein, sodium and calcium caseinate), phospholipids (for example, lecithin), carbohydrates, for example, cellulose and cellulose derivatives (for example, hydroxypropyl methylcellulose, ethylcellulose, methylcellulose, carboxymethyl cellulose), carrageenans, sorbitan esters (for example, mono-oleate, -palmitate, -stearate, trioleate) and mineral oils and fats (for example, paraffin).

[0061] The preparation of the probiotic substance after selection of the probiotic micro-organism and further components of the protecting matrix may be performed in any suitable way. A few principle steps of preparation of the probiotic substance may usually comprise the steps; concentrating the probiotic yield from a fermentor to a solid content of at least 10% and
5 cfu counts of at least 5×10 cfu/g paste, wet mixing of the probiotic concentrate with the other protective components, hydrogel formation in a desirable shape, drying and packaging. If probiotic micro matrix particles are required then a grinding and sieving steps are added. If additional moisture barrier coating is required then the dry probiotic substance is coated immediately after the completion of the drying step.

10 [0062] Most of these steps, for example "wet mixing" and "drying", may be subdivided, for example "mixing only few of the ingredients, drying them by one drying method, adding other ingredients to the mixture, mixing, and drying again with the same or different drying method.

[0063] For example, a wet mix of the micro-organisms and further components, is prepared by mixing all components in water, the slurry is then poured into molds of a desired shape and
15 let to harden by adding a cross-linking chemical or by cooling the slurry, or hardening the gel slurry first and then slicing, chopping or shaping the hydrogel to a desired shape. Then the molded chopped, or shaped hydrogel may be dried to water activity (A_w) below 0.2, preferably below 0.1. Possible drying processes include air dryers or convection ovens, belt dryers, vacuum dryers, fluidized bed dryers, rotary dryers, just to mention a few. Alternatively, all
20 components can be added to the wet probiotic concentrate without the addition of more water. For example, trehalose is first mixed in a probiotic concentrate (that contains about 10-20% solids), and then egg albumen or soy protein isolate and polysaccharides are added. Additional amount of proteins may be added to obtain semi-dry powder that can be further granulated, dried and sieved to specific size range of probiotic granulated-matrix particles.

25 [0064] The preparation of probiotic substances in a shape of crispy flakes or treats involves a vacuum drying process where the product temperature is set above the freezing temperature of the probiotic substance. In general, vacuum drying are performed in two steps. The first step involves moderately reduced pressure (ca. 5000 mTOR) and high shelf temperatures (ca. 5-50°C), whereas the second step involves lower pressures (e.g., higher vacuum such as not more
30 than 100 mTOR) while maintaining higher shelf temperature (up to about 50°C).

[0065] This process can be achieved using a programmable control system for vacuum pressure and product temperature. The vacuum and temperature conditions for the first drying step is adjusted empirically according to the size of the drier, heat transfer capacity, and the product load, but the goal is to keep the product above its freezing temperature while maximizing the water evaporation rate. In one embodiment, the temperature is initially maintained at about 40°C for about 6 hours, until most of the free water evaporated from the material, followed by gradually increasing the vacuum from the initial set up of ca. 5000 mTOR up to 100 mTOR, then maintaining these drying conditions until the water activity of the probiotic substance is sufficiently low (see values given above). Following this protocol, the drying procedure is completed within 24 hours without substantially compromising the probiotic viability. The large surface area of the shaped, sliced or chopped hydrogel greatly increases evaporation rate without the need to “boil” or foam the product in thin layers as indicated by other disclosures, thus eliminating inconsistent drying conditions and splattering of the foamed product solution within the vacuum chamber. Additionally, the disclosed composition and method of drying enable higher loading capacity of product in the vacuum or freeze drier as compared to other drying methods of materials with high sugar contents (i.e., foam formation).

[0066] The flake or treat may have the shapes as indicated above and be of any suitable, adequate or desired form. For example, they may have the form of spheres, cubes, pyramids, tablets, long tubes or any complex three-dimensional form. Furthermore they may have a form that corresponds to the food product to which they are added. For example, if the treats are added to a pet food for dogs, they may have the form of bones, animals, cats or other forms that fit with the food product.

[0067] In still another embodiment, the dried probiotic substance may be coated to further protect the micro-organism from deleterious effect of subsequent absorption of water during the shelf-life of the food product. The coating may be done by any suitable coating technique, for example, spraying, melt or solvent coating equipment, fluidized bed coater, drum coater or pan coater, just to mention a few.

[0068] The amount of coating depends on the size and form of the probiotic substance. Generally, the amount of coating compound is higher for smaller size particles. Accordingly,

the amount of the coating compound for probiotic micro-matrix or granulated matrix particles is from 10 to 50%, more preferably 20 to 40% of the total weight of the coated probiotic particles. The amount of the coating compound for flake or treat shapes probiotic substance is from 5 to 30%, more preferably 10 to 20% of the total weight of the coated probiotic treat. It is also understood that, for the purposes described herein, the coating process can result in a single layer of one compound or a mixture of compounds or to multiple layers of one or more compounds.

[0069] In a further embodiment, the probiotic substance provides gastric protection to the probiotic microorganisms. While the sugar component in the protective matrix immediately dissolves upon intact with the animal stomach juice, the polysaccharide protein matrix retains its form in the acid environment, thereby protecting the embedded microorganisms from digestive incursions. The matrix, however, slowly disintegrate in the alkali-phosphate environment of the gut and hence liberating the intact probiotic microorganisms to colonize the animal gut.

[0070] In additional embodiment, the probiotic product may be used exclusively as an entire food product, for example as a treat or a supplement, or added and mixed with a food product, or be used in its powder form to coat an existing food product, for example for top-coating pelleted or extruded feeds.

[0071] The following examples are given by way of illustration only and in no way should be construed as limiting the subject matter of the present application.

[0072] Examples

[0073] The subject matter of this disclosure is now described with reference to the following Examples. These Examples are provided for the purpose of illustration only, and the subject matter is not limited to these Examples, but rather encompasses all variations which are evident as a result of the teaching provided herein.

[0074] Example 1

[0075] Preparation of dry and stable probiotic substance

[0076] Basic formulation

[0077] 300 g of trehalose (Cargill Minneapolis, MN) was added to 1000 ml water and

5 allowed to completely dissolve. Soy protein isolate (100g, Fearn Natural Foods, Mequon, WI) and soy hydrolysate (20 g, Sigma-Aldrich, St. Louis, MO) were added under vigorous mixing using a standard household blender. Sodium alginate (20 g) was then mixed into the slurry and allowed to cool down to room temperature. *Lactobacillus paracasei* (100 g frozen concentrate direct from fermentation harvest) was then added to the slurry under vigorous mixing until a
10 smooth and uniform thick gel was achieved. The composition of the hydrogel is provided in Table 1.

Table 1. Hydrogel composition (g dry weight/100 ml water)

15	Trehalose	30 g
	Soy protein isolate	10 g
	Soy protein hydrolysate	2 g
	Sodium Alginate	2 g
	<i>L. paracasei</i> paste	10 g

20

[0078] Production of probiotic flakes

[0079] Five gram of calcium phosphate dibasic was mixed in the basic formula followed by 5 g of gluconolactone and the slurry was allowed to harden (solid hydrogel) at room temperature over the next 4 hours. The firm gel was sliced to thin and long leafs, through
25 cheese grinder. The thin leafs were loaded on a tray (13 x 10 inch) and placed in a freeze drier (Virtis Advantage, Virtis, Gardiner, NY). The condenser was set to -70°C and shelf temperature was set to $+40^{\circ}\text{C}$. The vacuum was then initiated and controlled at about 5000 mTOR with an external vacuum controller (Thyr-Cont, Electronic, GmbH). The wet product temperature fell to and stabilized at about -5 to 0°C . The chamber atmospheric pressure was
30 then gradually decreased as the product temperature started to warm up (measured by a pair of temperature sensors plugged in the wet material), until full vacuum pressure of 100 mTOR was established. Over this time period of increasing vacuum, the product temperature was carefully maintained between -5°C and $+5^{\circ}\text{C}$. Twenty four hours after establishing full vacuum, the

dried product was taken out of the freeze drier. The water activity (A_w) of the probiotic substance after the drying protocol was 0.05 (Measured by HygroPalm Aw1, R tonic Huntington, NY).

[0080] Production of probiotic treats

5 [0081] The above basic formulation slurry was poured into muffin plates having molds of small hearts or stars shapes and allowed to harden (solid hydrogel) at room temperature over the next 4 hours. The plates containing the molded hydrogel was placed in a freeze drier and allowed to dry as described above.

[0082] Production of probiotic micro matrix particles

10 [0083] The above basic formula was extruded or dripped into a 1000 ml bath (held at 0-5°C) containing 5 g CaCl_2 and 300 g trehalose using a syringe equipped with 18 G needle. The CaCl_2 bath was gently stirred while injecting the slurry. The matrix strings or drops were allowed to cross-link for 30 minutes and then harvested and blotted on paper towel. The strings or drops were first dried in a convection oven at 35 degree C until water activity of the material
15 reduced to 0.5 then they were transferred to a freeze drier for final drying of about 24 hours. The dry drops or strings ($A_w=0.06$) then ground to fine powder using standard coffee grinder and sieved through 50-250 micron screens.

[0084] Alternatively, probiotic micromatrix particles can be obtained by grinding and sieving already dried probiotic flakes or treats as described above.

20

[0085] Production of probiotic granulated matrix particles

[0086] 600 g of trehalose (Cargill Minneapolis, MN) was added to 1200 ml of concentrated paste of *Lactobacillus acidophilus* (20% solid concentrate direct from fermentation harvest) and allowed to completely dissolve. Soy protein isolate (200g, Fearn Natural Foods, Mequon, WI)
25 and soy hydrolysate (20 g, Sigma-Aldrich, St. Louis, MO) were added under vigorous mixing using a standard household mixer. Sodium alginate (40 g) was then mixed into the slurry and allowed to cool down to room temperature. After a smooth, thick and uniform gel was achieved, 5 g of calcium carbonate was added. Immediately after, 1000 g of egg white (Sigma-Aldrich, St. Louis, MO) was slowly added under vigorous mixing until a semi-moist free
30 flowing powder was obtained. The granulated probiotic powder was dried in a convection oven

at 40 degree C for 2 h followed by a vacuum drying for 24 h to obtain dry ($A_w < 0.06$) granulated matrix particles. The granulated particles can be sieved to specific size range through a series of 50-500 micron sieves. The composition of the probiotic granulated matrix particles is provided in Table 2.

5

Table 2. Hydrogel composition

10	Trehalose	60 g
	Soy protein isolate	20 g
	Soy protein hydrolysate	2 g
	Sodium Alginate	4 g
	<i>L. acidophilus</i> paste	120 g
	Egg white	100 g

[0087] Recovery of *L. rhamnosus* after production of dry micromatrix particles.

15 [0088] *L. rhamnosus* micromatrix particles were produced as described above. CFU counts of concentrated *L. rhamnosus* after harvesting from a fermentor and centrifuging were 5×10^{10} /g paste and solid content was 24%. After mixing the probiotic with all protective components, forming hydrogel, chopping the hydrogel to small threads, drying in a freeze-drier for 24 h, grinding and sieving to particle size between 50-250 micron, the CFU counts were 9.4×10^9 /g dry micromatrix. This represents a loss of 0.28 log of probiotic activity during the manufacturing process (as illustrated in Figure 1).

[0089] Stability of *L. rhamnosus* in 40°C at 33% relative humidity (RH).

25 [0090] *L. rhamnosus* probiotic micromatrix particles were prepared and dried as described above. The dried probiotic particles were placed in temperature and humidity control incubators set at 40°C and 33% relative humidity for 4 weeks. Viabilities of bacteria dried in only in 10% trehalose solution or in the protective mixture described herein were measured on a weekly interval. Figure 2 shows that the protective mixture provided a significant protection to that of trehalose alone dried bacteria.

30

[0091] Stability of *L. rhamnosus* micromatrix particles in simulated gastric juices

[0092] Micromatrix particles (50-250 micron) containing either *L. rhamnosus* (LGG), *L. acidophilus* (LA5) or *L. paracase* (LPC) were prepared and dried as described above. The

micromatrix particles were then exposed for 2 hours to simulated gastric juice (full stomach -- 12% non fat skim milk, 2% glucose, 1% yeast extract and 0.05% cysteine; pH 2; or empty stomach -- 0.32% pepsin, 0.2% sodium chloride, pH 1.2). Bacterial viabilities were recorded before and after the exposure to the simulated gastric juices. Figures 3 demonstrate a significant protection of the probiotic bacteria in the micromatrix substance in simulated gastric conditions.

[0093] Coating of the probiotic substances

[0094] Probiotic substances were coated with a fat-based moisture barrier (a mixture of 20% jojoba oil, 70% cocoa butter and 10% beeswax) in a drum tumbler at a temperature of 40 degrees C. The moisture barrier was sprayed on using a spraying nozzle while the dry material is agitated in the drum tumbler to ensure homogeneous coating. The total amount of coating was about 20% (of the uncoated probiotic substances) for probiotic flakes and treats, and 40-50% for probiotic micromatrix particles.

[0095] Example 2

[0096] Preparation of probiotic pet food

[0097] Pet food for dogs that is commercially available was first dried in a convection oven to a water activity of 0.1, and then coated with probiotic micromatrix particles in a drum tumbler. The pellets were first sprayed with about 5% of fat-based moisture barrier (a mixture of 40% chicken fat, 40% cocoa butter and 20% beeswax), then mixed with the micro matrix particles (usually 0.1-0.5% of total pet food that provides a dosage of 10^8 CFU/g) and finally sprayed with additional coat of the fat-based moisture barrier. The total amount of coating was about 15% (of the pet food). Coating time was about 30 minutes.

[0098] Example 3

[0099] Preparation of fish feed with several probiotic micro-organisms

[0100] Pelleted feed for fish was prepared with a mixture of several probiotics. Probiotic micromatrix particles containing a mixture of *L. rhamnosus*, *L. acidophilus* and *Bifidobacterium lactis* (DSM 20215) were prepared as described in Example 1. Fish feed that is commercially available was first dried in a convection oven to a water activity of 0.1, and

then coated with probiotics micromatrix particles in a drum tumbler. The pellets were first sprayed with about 5% of fat-based moisture barrier (a mixture of 40% fish oil, 40% cocoa butter and 20% beeswax), then mixed with the micro matrix particles (usually 0.1-0.5% of total fish feed that provides a dosage of 10^7 cfu/g) and finally sprayed with additional coat of the fat-based moisture barrier. The total amount of coating was about 10% (of the fish feed).

[0101] Example 4

[0102] Preparation of probiotic "chips" (as shown in Figure 4)

[0103] Probiotic chips were prepared with a mixture of *L. rhamnosus* and *L. acidophilus* as

described in Example 1. The basic formula was added with about 1-2% of probiotic pastes instead of the standard 10-20% load. Five gram of calcium phosphate dibasic was mixed in the basic formula followed by 5 g of gluconolactone and the slurry was pored into long plastic tubes of 1-inch diameter and allowed to harden (solid hydrogel) at room temperature over the next 4 hours. The firm "hot-dog" shaped gel was sliced to thin discs, loaded on a tray (13 x 10 inch) and placed in a freeze drier and a drying procedure applied as described in Example#1.

The dried probiotic chips product was taken out of the freeze drier and packed under nitrogen in small aluminum foiled packs that provide tasty and crispy treat containing an effective dose of 10^8 CFU/per serving. Figure 4 shows an example of probiotic chips containing 10^8 CFU/serving of *rhamnosus* produced as described herein.

[0104] Example 5

[0105] Preparation of probiotic cereals

[0106] Breakfast cereal that is commercially available was first dried in a convection oven to a water activity of 0.1. Probiotic chips containing a standard 10-20% load of probiotics were produced and further coated with a moisture barrier as described in Examples 1 and 4. The cereals were then mixed with 0.1-0.5% probiotic chips (to provides a dosage of 10^8 cfu/g).

[0107] Example 6

[0108] Alternative delivery forms of probiotic matrix substances in the lid of a container or a bottle of beverage drink

[0109] In general, all forms of the probiotic substances produced as in Example 1 or

5 Example 4 work similarly well, confirming the high versatility of usage and application of the materials. The probiotic substances produced as described herein can be provided in small bags or paper "stick" packaging or in any other manner along with an edible food product. The dry micromatrix particles or chips may also be provided in the lid of a yogurt container or a bottle of beverage drink to be mixed with the liquid food product for consumption.

10

[0110] References

[0111] The following references are referred to herein.

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5

[0124] The disclosure of every patent, patent application, and publication cited herein is hereby incorporated herein by reference in its entirety.

[0125] While this subject matter has been disclosed with reference to specific embodiments, it is apparent that other embodiments and variations can be devised by others skilled in the art without departing from the true spirit and scope of the subject matter described herein. The appended claims include all such embodiments and equivalent variations.

10

CLAIMS

1. A probiotic substance comprising viable probiotic bacteria preserved in a dried matrix of at least one sugar wherein the total amount of sugar compound in the matrix is from
5 about 10% to about 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix.
- 10 2. The probiotic substance according to claim 1, wherein the sugar in the dried matrix is a disaccharide.
3. The probiotic substance according to claim 2, wherein the disaccharide is selected from the group consisting of sucrose and trehalose.
- 15 4. The probiotic substance according to claim 1, wherein the protein in the dried matrix is an isolate or hydrolyzed protein selected from the group consisting of egg white, casein, soy protein isolate, soy hydrolysate and a combination thereof.
- 20 5. The probiotic substance according to claim 1, wherein the polysaccharide is a cross-linkable gel, selected from the group consisting of an alginate, pectin, and chitosan.
6. The probiotic substance according to claim 1, wherein the dried matrix is comprised of 20% w/v to 60% w/v trehalose, 2% w/v to 20% w/v egg white or soy protein isolate and
25 0.5% w/v to 10% w/v alginates.
7. The probiotic substance according to claim 1, wherein the probiotic microorganism comprises 10^5 to 10^{14} viable micro-organisms per gram of dry substance.
- 30 8. The probiotic substance according to claim 1, wherein the probiotic substance is coated with additional moisture barrier.

9. The probiotic substance according to claim 8, wherein the moisture barrier comprises any food-grade substance having water repelling or impermeable properties.
- 5 10. The probiotic substance according to claim 8, wherein the moisture barrier comprises a mixture of oil based substances.
11. The probiotic substance according to claim 1, wherein the probiotic bacteria retain most of their initial biological activity after manufacturing, after extended exposure to
10 temperature and humidity conditions of gastric exposure.
12. A human food product comprising one or several probiotic substances comprising viable probiotic bacteria preserved in a dried matrix comprising at least one sugar, wherein the total amount of sugar compound in the matrix is from about 10% to about
15 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix, wherein the dried matrix has water activity below 0.2.
- 20 13. The human food product of claim 12, wherein the food product is in a moist, semi-moist, or semi-dry form.
14. The human food product of claim 12, wherein the food product is in a powdered,
25 particulate, pellet, tablet, capsule, colloidal suspension or liquid form.
15. The human food product of claim 12, wherein the food product is added to a bar, liquid formula, or cereal or another food product.
- 30 16. The human food product of claim 12, wherein the food product is in the form of a treat, a nutraceutical food additive or a pharmaceutical food additive.

17. An animal feed product comprising one or several probiotic substances comprising viable probiotic bacteria preserved in a dried matrix comprising at least one sugar wherein the total amount of sugar compound in the matrix is from about 10% to about 5 60% by weight of the matrix, at least one protein wherein the total amount of proteins in the matrix is from about 2% to about 20% by weight of the matrix, and at least one polysaccharide wherein the total amount of polysaccharides in the matrix is from about 0.5% to about 10% by weight of the matrix.
- 10 18. The animal feed product of claim 17, wherein the feed product is in a moist, semi-moist, or semi-dry form.
19. The animal feed product of claim 17, wherein the feed product is in powdered, particulate, pellet, tablet, capsule, colloidal suspension or liquid form.
- 15 20. The animal feed product of claim 17, wherein the feed product is added to a bar, liquid formula, or cereal or another food product.
21. The animal feed product of claim 17, wherein the feed product is in the form of a 20 treat, nutraceutical food additive or a pharmaceutical feed additive.
22. The animal feed product of claim 17, wherein the animal is chosen from terrestrial or aquatic animals.
- 25 23. A method of making probiotic substance or food product according to any of the above claims, the method comprising:
- a) mixing a preparation of probiotic micro-organisms with at least one sugar, at least one protein, and at least one polysaccharide to obtain a smooth and uniform slurry or gel in a desired shape and form; and
- 30 b) vacuum drying the preformed solid gel to a water activity below 0.1, wherein the vacuum drying comprises: setting the temperature of the above the freezing

temperature of the probiotic substance; and vacuum drying at reduced pressure of about 5000 mTOR at a shelf temperature of from 5-50°C and then at a pressure of about 100 mTOR while maintaining the shelf temperature.

- 5 24. The method of claim 23, wherein the slurry is not cross-linked before drying.

FIG 1. Effect of each component in the composition of the present invention on viability Loss of LGG Probiotic after vacuum-drying

LGG paste	Trehalose	Egg White	Alginate	Log Loss
10%	0%	0%	0%	2.27
10%	30%	0%	2%	1.96
10%	0%	30%	2%	1.87
10%	30%	10%	2%	0.28

* Component amount is in percent of water content in the composition

FIG 2. Shelf Life of Probiotic LGG Under Accelerated Conditions at High Temperature (40°C) and Humidity (33% RH)

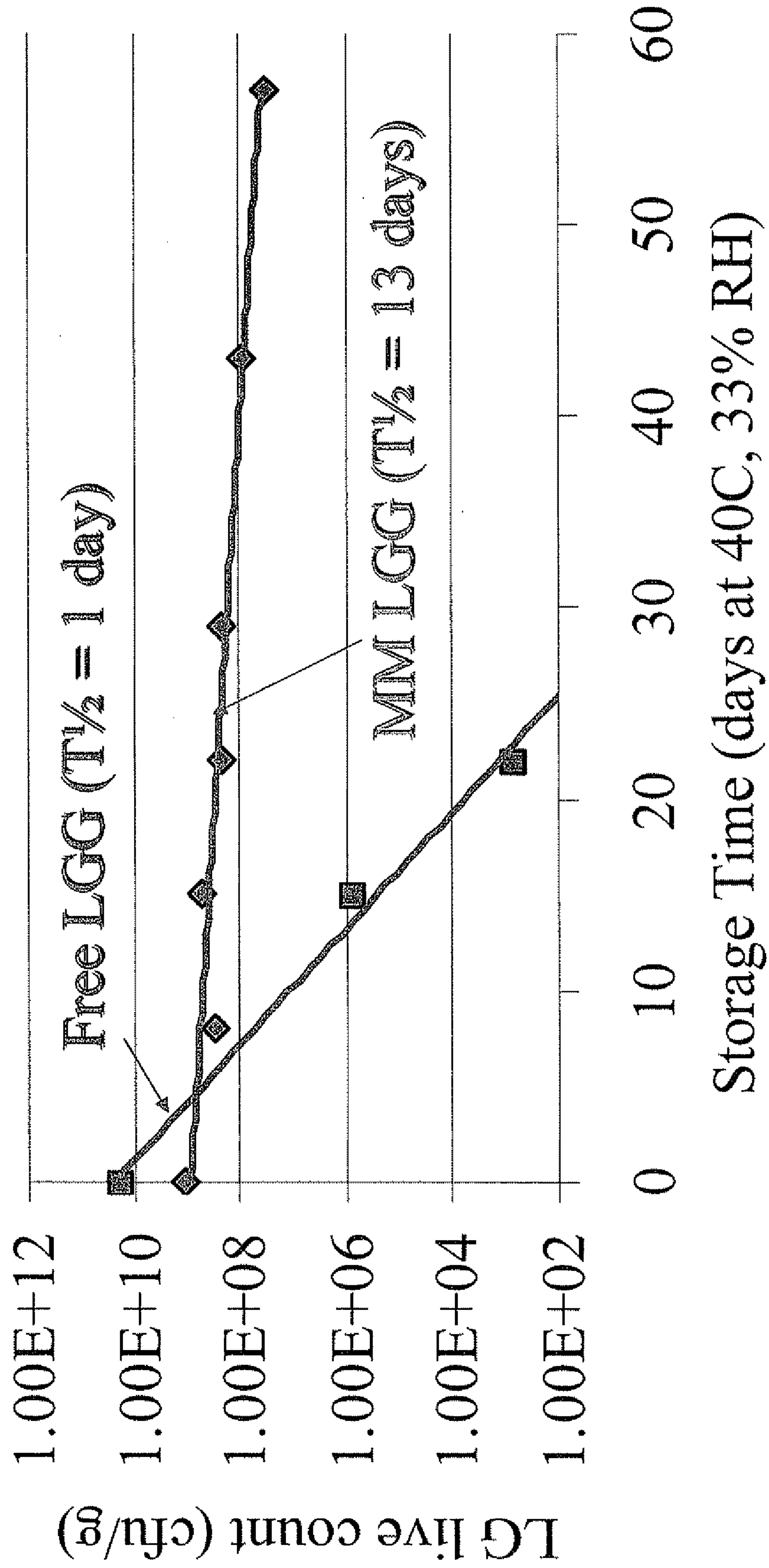
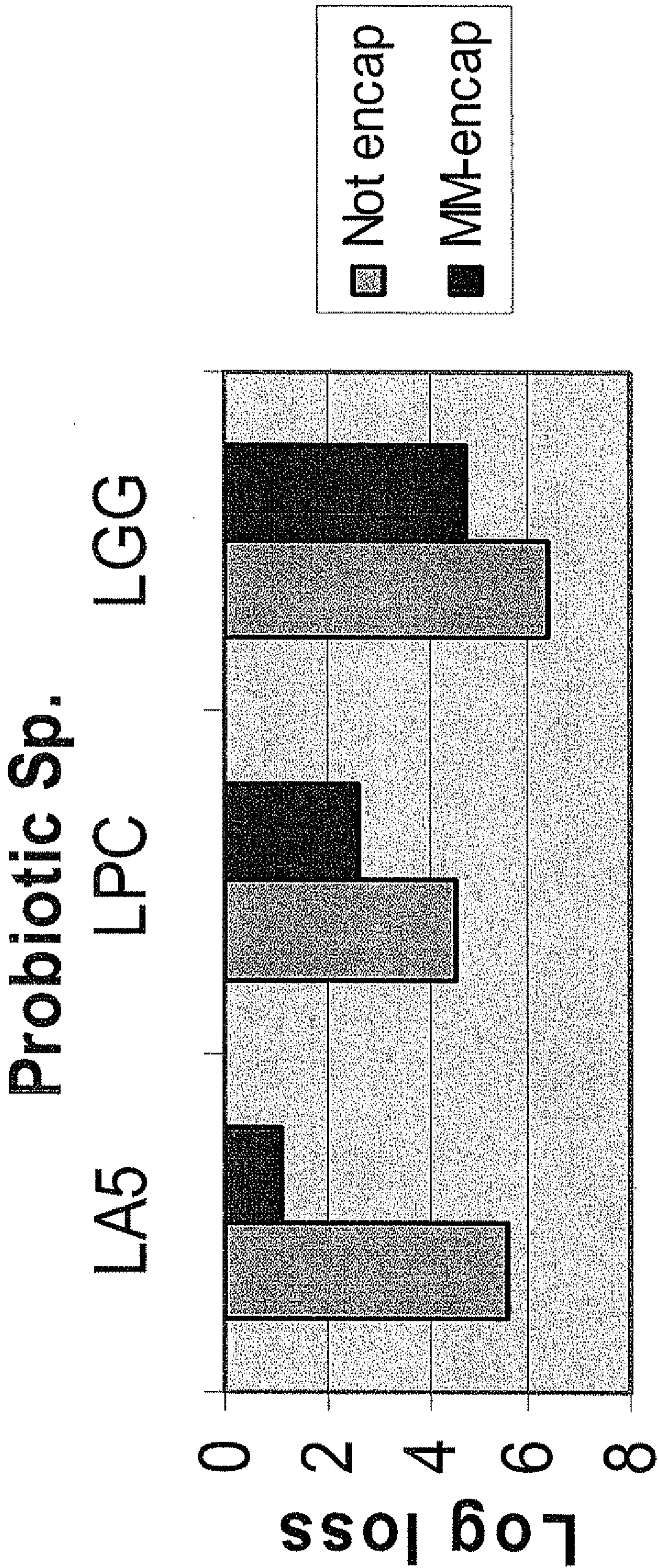


FIG 3. Gastric Loss of Probiotics, Produced as Described Herein (pH 1.2)

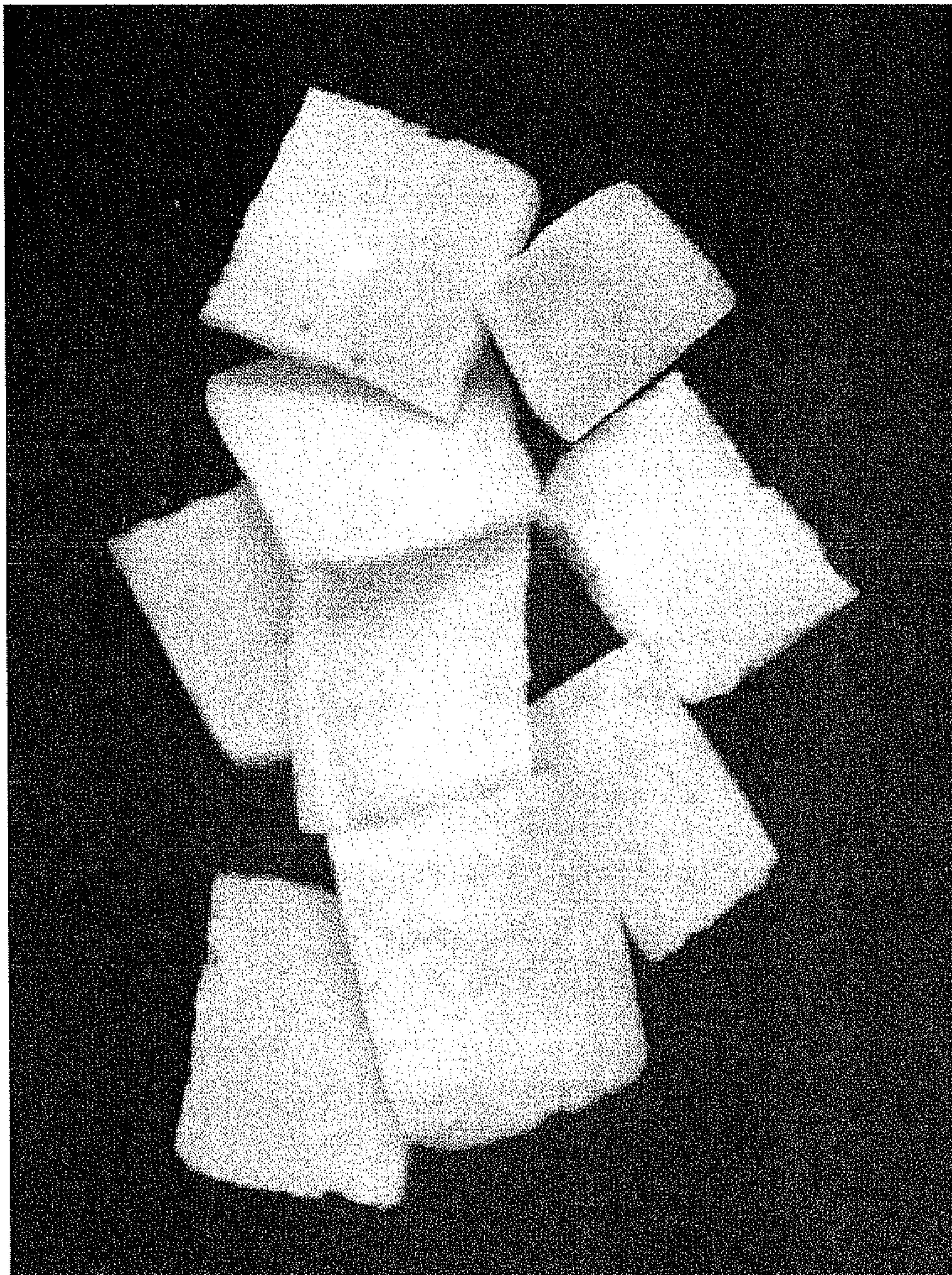
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**FIG 4. Examples of Probiotic Chips Produced
as Described Herein**



Examples of Probiotic Chips Produced as Described Herein

