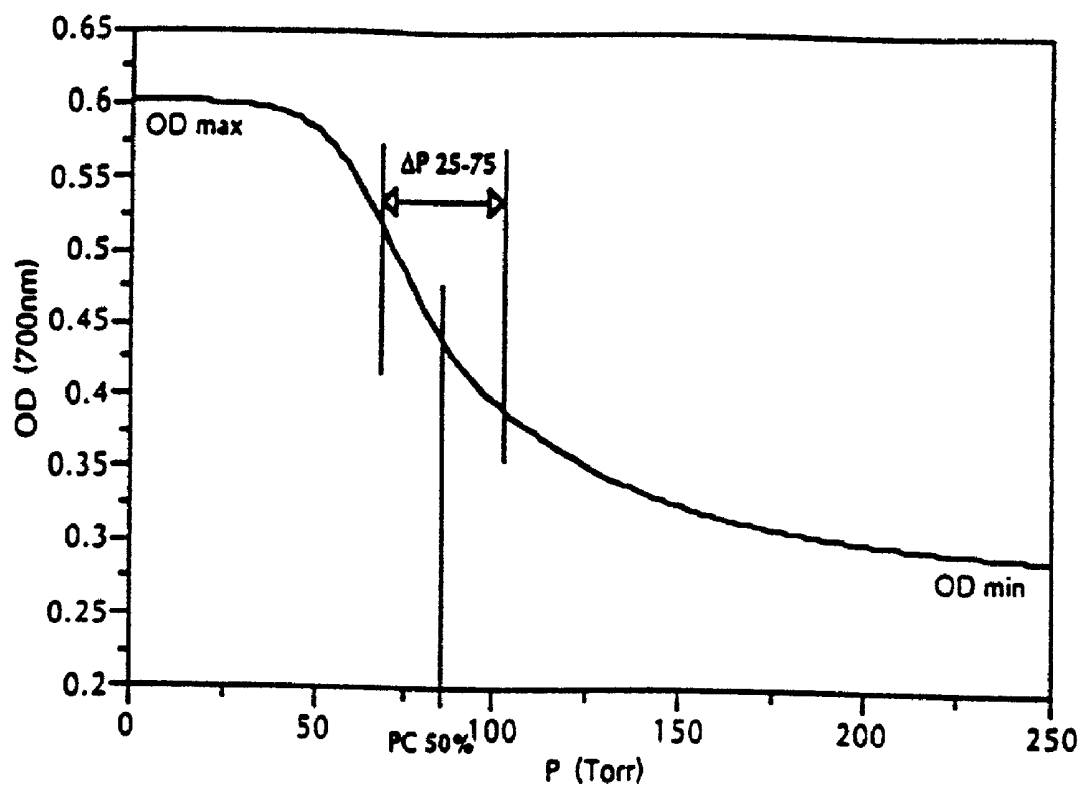




US 20010024640A1

(19) **United States**(12) **Patent Application Publication**
Schneider et al.(10) **Pub. No.: US 2001/0024640 A1**(43) **Pub. Date: Sep. 27, 2001**(54) **ULTRASOUND CONTRAST AGENTS AND
METHODS OF MAKING AND USING THEM**(76) Inventors: **Michel Schneider**, Troinex (CH);
Daniel Bichon, Montpellier (FR);
Philippe Bussat, Feigères (FR);
Jerome Puginier, Le Chable-Beaumont
(FR); **Eva Hybl**, Heidelberg (DE); **Feng
Yan**, Carouge (CH); **Pascal Grenier**,
Ambilly (FR); **Nadine Garcel**, legal
representative, Ambilly (FR);
Marie-Bernadette Barrau, Geneve
(CH)Correspondence Address:
NIXON & VANDERHYE P.C.
8th Floor
1100 North Glebe Road
Arlington, VA 22201 (US)(21) Appl. No.: **09/748,120**(22) Filed: **Dec. 27, 2000****Related U.S. Application Data**(60) Continuation-in-part of application No. 09/630,537,
filed on Aug. 1, 2000, which is a division of appli-
cation No. 09/021,150, filed on Feb. 10, 1998, now
Pat. No. 6,136,293, which is a division of application
No. 08/853,936, filed on May 9, 1997, now Pat. No.
6,110,443, which is a division of application No.
08/456,385, filed on Jun. 1, 1995, now Pat. No.
5,658,551, which is a division of application No.
08/315,347, filed on Sep. 30, 1994, now Pat. No.
5,531,980, which is a division of application No.
08/128,540, filed on Sep. 29, 1993, now Pat. No.
5,380,519, which is a division of application No.
07/775,989, filed on Nov. 20, 1991, now Pat. No.
5,271,928, which is a 371 of international applicationNo. PCT/EP91/00620, filed on Apr. 2, 1991. Con-
tinuation-in-part of application No. 08/740,653, filed
on Oct. 31, 1996, which is a division of application
No. 08/380,588, filed on Jan. 30, 1995, now Pat. No.
5,578,292, which is a division of application No.
07/991,237, filed on Dec. 16, 1992, now Pat. No.
5,413,774. Continuation-in-part of application No.
08/910,152, filed on Aug. 13, 1997, now Pat. No.
6,200,548, which is a division of application No.
08/288,550, filed on Aug. 10, 1994, now Pat. No.
5,711,933, which is a division of application No.
08/033,435, filed on Mar. 18, 1993, now abandoned,
which is a division of application No. 07/695,343,
filed on May 3, 1991, now abandoned.(30) **Foreign Application Priority Data**Apr. 2, 1990 (EP) 90810262.7
Jan. 24, 1992 (EP) 92810046.0
May 18, 1990 (EP) 90810367.4
Apr. 2, 1991 (EP) PCT/EP91/00620**Publication Classification**(51) **Int. Cl.⁷** **A61K 49/22**
(52) **U.S. Cl.** **424/9.52**(57) **ABSTRACT**Gas filled microbubble and microballoon suspensions in
aqueous phases usable as imaging contrast agents in ultra-
sonic echography. One can impart outstanding resistance
against collapse under pressure to these gas-filled
microbubbles and microballoons used as contrast agents in
ultrasonic echography by using as fillers gases whose solu-
bility in water, expressed in liter of gas by liter of water
under standard conditions, divided by the square root of the
molecular weight does not exceed 0.003. Contrast agents
with particular mixtures of gases are disclosed that have
advantageous properties. Dry formulations useful in making
the contrast agents are also disclosed.

Fig 1



ULTRASOUND CONTRAST AGENTS AND METHODS OF MAKING AND USING THEM

[0001] This application is a continuation-in-part of Ser. No. 09/630,537, filed Aug. 1, 2000, which is a divisional of Ser. No. 09/021,150, filed Feb. 10, 1998, U.S. Pat. No. 6,136,293, which is a divisional of Ser. No. 08/853,936, filed May 9, 1997, U.S. Pat. No. 6,110,443, which is a divisional of Ser. No. 08/456,385, filed Jun. 1, 1995, U.S. Pat. No. 5,658,551, which is a divisional of Ser. No. 08/315,347, filed Sep. 30, 1994, U.S. Pat. No. 5,531,980, which is a divisional of Ser. No. 08/128,540, filed Sep. 29, 1993, U.S. Pat. No. 5,380,519, which is a divisional of Ser. No. 07/775,989, filed Nov. 20, 1991, U.S. Pat. No. 5,271,928, which was the National Stage of International Application No. PCT/EP91/00620, filed Apr. 2, 1991, which originated from EP 90810262.7, filed Apr. 2, 1990; this application is also a continuation-in-part of Ser. No. 08/740,653, filed Oct. 31, 1996, which is a divisional of Ser. No. 08/380,588, filed Jan. 30, 1995, U.S. Pat. No. 5,578,292, which is a divisional of Ser. No. 07/991,237, filed Dec. 16, 1992, U.S. Pat. No. 5,413,774, which originated from EP 92810046.0, filed Jan. 24, 1992; this application is also a continuation-in-part of Ser. No. 08/910,152, filed Aug. 13, 1997, which is a divisional of Ser. No. 08/288,550, filed Aug. 10, 1994, U.S. Pat. No. 5,711,933, which is a divisional of Ser. No. 08/033,435, filed Mar. 18, 1993, which is a divisional of Ser. No. 07/695,343, filed May 3, 1991, which originated from EP 90810367.4, filed May 18, 1990. All of the above applications are hereby incorporated by reference herein in their entirety.

SUMMARY

[0002] The present invention concerns media adapted for injection into living bodies, e.g., for the purpose of ultrasonic echography and, more particularly, injectable liquid compositions comprising microbubbles of air or physiologically acceptable gases as stable dispersions or suspensions in an aqueous liquid carrier. These compositions are mostly usable as contrast agents in ultrasonic echography to image the inside of blood-stream vessels and other cavities of living beings, e.g., human patients and animals. Other uses however are also contemplated as disclosed hereafter.

[0003] The invention also comprises dry compositions which, upon admixing with an aqueous carrier liquid, will generate the foregoing sterile suspension of microbubbles thereafter usable as contrast agents for ultrasonic echography and other purposes. The present invention also concerns stable dispersions or compositions of gas filled microvesicles in aqueous carrier liquids. These dispersions are generally usable for most kinds of applications requiring gases homogeneously dispersed in liquids. One notable application for such dispersions is to be injected into living beings, for instance for ultrasonic echography and other medical applications. The invention also concerns the methods for making the foregoing compositions including some materials involved in the preparations, for instance pressure-resistant gas-filled microbubbles, microcapsules and microballoons.

BACKGROUND

[0004] It is well known that microbodies of air or a gas (defined here as microvesicles), e.g., microbubbles or

microballoons, suspended in a liquid are exceptionally efficient ultrasound reflectors for echography. In this disclosure, the term "microbubble" specifically designates air or gas globules in suspension in a liquid which generally results from the introduction therein of air or a gas in divided form, the liquid preferably also containing surfactants or tensides to control the surface properties thereof and the stability of the bubbles. More specifically, one may consider that the internal volume of the microbubbles is limited by the gas/liquid interface, or in other words, the microbubbles are only bounded by a rather evanescent envelope involving the molecules of the liquid and surfactant loosely bound at the gas to liquid junction boundary.

[0005] The term "microcapsule" or "microballoon" designates preferably air or gas bodies with a material boundary or envelope formed of molecules other than that of the liquid of suspension, e.g., a polymer membrane wall. Both microbubbles and microballoons are useful as ultrasonic contrast agents. For instance, injecting into the blood-stream of living bodies suspensions of gas microbubbles or microballoons (in the range of 0.5 to 10 μm) in a carrier liquid will strongly reinforce ultrasonic echography imaging, thus aiding in the visualization of internal organs. Imaging of vessels and internal organs can strongly help in medical diagnosis, for instance for the detection of cardiovascular and other diseases.

[0006] The formation of suspensions of microbubbles in an injectable liquid carrier suitable for echography can follow various routes, such as by the release of a gas dissolved under pressure in this liquid, or by a chemical reaction generating gaseous products, or by admixing with the liquid soluble or insoluble solids containing air or gas trapped or adsorbed therein. For instance in DE-A-3529195 (Max-Planck Gesell.), there is disclosed a technique for generating 0.5-50 μm bubbles in which an aqueous emulsified mixture containing a water soluble polymer, an oil and mineral salts is forced back and forth, together with a small amount of air, from one syringe into another through a small opening. Here, mechanical forces are responsible for the formation of bubbles in the liquid.

[0007] M. W. Keller et al. (J. Ultrasound Med. 5 (1986), 439-8) have reported subjecting to ultrasonic cavitation under atmospheric pressure solutions containing high concentrations of solutes such as dextrose, Renografin-76, Iopamidol (an X-ray contrast agent), and the like. There the air is driven into the solution by the energy of cavitation.

[0008] Other techniques rely on the shaking of a carrier liquid in which air containing microparticles have been incorporated, said carrier liquid usually containing, as stabilizers, viscosity enhancing agents, e.g., water soluble polypeptides or carbohydrates and/or surfactants. The stability of the microbubbles against decay or escape to the atmosphere is controlled by the viscosity and surface properties of the carrier liquid. The air or gas in the microparticles can consist of inter-particle or intra-crystalline entrapped gas, as well as surface adsorbed gas, or gas produced by reactions with the carrier liquid, usually aqueous. All this is fully described for instance in EP-A-0052575 (Ultra Med. Inc.) in which there are used aggregates of 1-50 μm particles of carbohydrates (e.g., galactose, maltose, sorbitol, gluconic acid, sucrose, glucose and the like) in aqueous solutions of glycols or polyglycols, or other water soluble polymers.

[0009] Also, in EP-A-0123235 and EP-A-0122624 (Schering, see also EP-A-0320433) use is made of air trapped in solids. For instance, EP-A-0122624 claims a liquid carrier contrast composition for ultrasonic echography containing microparticles of a solid surfactant, the latter being optionally combined with microparticles of a non-surfactant. As explained in this latter document, the formation of air bubbles in the solution results from the release of the air adsorbed on the surface of the particles, or trapped within the particle lattice, or caught between individual particles, this being so when the particles are agitated with the liquid carrier.

[0010] EP-A-0131540 (Schering) also discloses the preparation of microbubbles suspensions in which a stabilized injectable carrier liquid, e.g., a physiological aqueous solution of salt, or a solution of a sugar like maltose, dextrose, lactose or galactose, without viscosity enhancer, is mixed with microparticles (in the 0.1 to 1 μm range) of the same sugars containing entrapped air. In order that the suspension of bubbles can develop within the liquid carrier, the foregoing documents recommend that both liquid and solid components be violently agitated together under sterile conditions; the agitation of both components together is performed for a few seconds and, once made, the suspension must then be used immediately, i.e., it should be injected within 5-10 minutes for echographic measurements; this indicates that the bubbles in the suspensions are not longlived and one practical problem with the use of microbubbles suspensions for injection is lack of stability with time. The present invention fully remedies this drawback.

[0011] In an attempt to cure the evanescence problem, microballoons, i.e., microvesicles with a material wall, have been developed. As said before, while the microbubbles only have an immaterial or evanescent envelope, i.e., they are only surrounded by a wall of liquid whose surface tension is being modified by the presence of a surfactant, the microballoons or microcapsules have a tangible envelope made of substantive material, e.g., a polymeric membrane with definite mechanical strength. In other terms, they are microvesicles of material in which the air or gas is more or less tightly encapsulated.

[0012] In U.S. Pat. No. 4,466,442 (Schering), there is disclosed a series of different techniques for producing suspensions of gas microbubbles in a liquid carrier liquid carrier using (a) a solution of a tenside (surfactant) in a carrier liquid (aqueous) and (b) a solution of a viscosity enhancer as stabilizer. For generating the bubbles, the techniques used there include forcing at high velocity a mixture of (a), (b) and air through a small aperture; or injecting (a) into (b) shortly before use together with a physiologically acceptable gas; or adding an acid to (a) and a carbonate to (b), both components being mixed together just before use and the acid reacting with the carbonate to generate CO_2 bubbles; or adding an over-pressurized gas to a mixture of (a) and (b) under storage, said gas being released into microbubbles at the time when the mixture is used for injection.

[0013] The tensides used in component (a) of U.S. Pat. No. 4,466,442 comprise lecithins; esters and ethers of fatty acids and fatty alcohols with polyoxyethylene and polyoxyethylated polyols like sorbitol, glycols and glycerol, chole-

terol; and polyoxy-ethylene-polyoxypropylene polymers. The viscosity raising and stabilizing compounds include for instance mono- and polysaccharides (glucose, lactose, sucrose, dextran, sorbitol); polyols, e.g., glycerol, polyglycols; and polypeptides like proteins, gelatin, oxypolygelatin, plasma protein and the like.

[0014] In a typical preferred example of this latter document, equivalent volumes of (a) a 0.5% by weight aqueous solution of Pluronic® F-68 (a polyoxypropylene-polyoxyethylene polymer) and (b) a 10% lactose solution are vigorously shaken together under sterile conditions (closed vials) to provide a suspension of microbubbles ready for use as an ultrasonic contrast agent and lasting for at least 2 minutes. About 50% of the bubbles had a size below 50 μm .

[0015] Although the achievements of the prior art have merit, they suffer from several drawbacks which strongly limit their practical use by doctors and hospitals, namely their relatively short life-span (which makes test reproducibility difficult), relative low initial bubble concentration (the number of bubbles rarely exceeds 10^4 - 10^5 bubbles/ml and the count decreases rapidly with time) and poor reproducibility of the initial bubble count from test to test (which also makes comparisons difficult). Also it is admitted that for efficiently imaging certain organs, e.g., the left heart, bubbles smaller than 50 μm , preferably in the range of 0.5-10 μm , are required; with larger bubbles, there are risks of clots and consecutive emboly. Furthermore, the compulsory presence of solid microparticles or high concentrations of electrolytes and other relatively inert solutes in the carrier liquid may be undesirable physiologically in some cases. Finally, the suspensions are totally unstable under storage and cannot be marketed as such; hence great skill is required to prepare the microbubbles at the right moment just before use.

[0016] Of course there exists stable suspensions of microcapsules, i.e., microballoons with a solid, air-sealed rigid polymeric membrane which perfectly resist for long storage periods in suspension, which have been developed to remedy this shortcoming (see for instance K. J. Widder, EP-A-0324938); however the properties of microcapsules in which a gas is entrapped inside solid membrane vesicles essentially differ from that of the gas microbubbles of the present invention and belong to a different kind of art; for instance while the gas microbubbles discussed here will simply escape or dissolve in the blood-stream when the stabilizers in the carrier liquid are excreted or metabolized, the solid polymer material forming the walls of the aforementioned micro-balloons must eventually be disposed of by the organism being tested which may impose a serious afterburden upon it. Also capsules with solid, non-elastic membrane may break irreversibly under variations of pressure.

STABILIZED MICROBUBBLE COMPOSITIONS OF THE INVENTION

[0017] The compositions of the present invention fully remedy the aforementioned pitfalls.

[0018] The term "lamellar form" defining the condition of at least a portion of the surfactant or surfactants of the present composition indicates that the surfactants, in strong contrast with the microparticles of the prior art (for instance EP-A-0123235), are in the form of thin films involving one or more molecular layers (in laminate form). Converting

film forming surfactants into lamellar form can easily be done for instance by high pressure homogenization or by sonication under acoustical or ultrasonic frequencies. In this connection, it should be pointed out that the existence of liposomes is a well known and useful illustration of cases in which surfactants, more particularly lipids, are in lamellar form.

[0019] Liposome solutions are aqueous suspensions of microscopic vesicles, generally spherically shaped, which hold substances encapsulated therein. These vesicles are usually formed of one or more concentrically arranged layers (lamellae) of amphipathic compounds, i.e., compounds having a lipophobic hydrophilic moiety and a lipophilic hydrophobic moiety. See for instance "Liposome Methodology", Ed. L. D. Leserman et al, Insem 136, 2-8 (May 1982). Many surfactants or tensides, including lipids, particularly phospholipids, can be laminarized to correspond to this kind of structure. In this invention, one preferably uses the lipids commonly used for making liposomes, for instance the lecithins and other tensides disclosed in more detail hereafter, but this does in no way preclude the use of other surfactants provided they can be formed into layers or films.

[0020] It is important to note that no confusion should be made between the microbubbles of this invention and the disclosure of Ryan (U.S. Pat. No. 4,900,540) reporting the use of air or gas filled liposomes for echography. In this method Ryan encapsulates air or a gas within liposomic vesicles; in embodiments of the present invention microbubbles of air or a gas are formed in a suspension of liposomes (i.e., liquid filled liposomes) and the liposomes apparently stabilize the microbubbles. In Ryan, the air is inside the liposomes, which means that within the bounds of the presently used terminology, the air filled liposomes of Ryan belong to the class of microballoons and not to that of the microbubbles.

[0021] Practically, to achieve the suspensions of microbubbles according to the invention, one may start with liposomes suspensions or solutions prepared by any technique reported in the prior art, with the obvious difference that in the present case the liposomic vesicles are preferably "unloaded", i.e., they do not need to keep encapsulated therein any foreign material other than the liquid of suspension as is normally the object of classic liposomes. Hence, preferably, the liposomes of the present invention will contain an aqueous phase identical or similar to the aqueous phase of the solution itself. Then air or a gas is introduced into the liposome solution so that a suspension of microbubbles will form, said suspension being stabilized by the presence of the surfactants in lamellar form. Notwithstanding, the material making the liposome walls can be modified within the scope of the present invention, for instance by covalently grafting thereon foreign molecules designed for specific purposes as will be explained later.

[0022] The preparation of liposome solutions has been abundantly discussed in many publications, e.g., U.S. Pat. No. 4,224,179 and WO-A-88/09165 and all citations mentioned therein. This prior art is used here as reference for exemplifying the various methods suitable for converting film forming tensides into lamellar form. Another basic reference by M. C. Woodle and D. Papahadjopoulos is found in "Methods in Enzymology" 171 (1989), 193.

[0023] For instance, in a method disclosed in D. A. Tyrrell et al, *Biochimica & Biophysica Acta* 457 (1976), 259-302, a mixture of a lipid and an aqueous liquid carrier is subjected to violent agitation and thereafter sonicated at acoustic or ultrasonic frequencies at room or elevated temperature. In the present invention, it has been found that sonication without agitation is convenient. Also, an apparatus for making liposomes, a high pressure homogenizer such as the Microfluidizer®, which can be purchased from Microfluidics Corp., Newton, Mass. 02164 USA, can be used advantageously. Large volumes of liposome solutions can be prepared with this apparatus under pressures which can reach 600-1200 bar.

[0024] In another method, according to the teaching of GB-A-2,134,869 (Squibb), microparticles (10 μ m or less) of a hydrosoluble carrier solid (NaCl, sucrose, lactose and other carbohydrates) are coated with an amphipathic agent; the dissolution of the coated carrier in an aqueous phase will yield liposomic vesicles. In GB-A-2,135,647 insoluble particles, e.g., glass or resin microbeads are coated by moistening in a solution of a lipid in an organic solvent followed by removal of the solvent by evaporation. The lipid coated microbeads are thereafter contacted with an aqueous carrier phase, whereby liposomic vesicles will form in that carrier phase.

[0025] The introduction of air or gas into a liposome solution in order to form therein a suspension of microbubbles can be effected by usual means, inter alia by injection, that is, forcing said air or gas through tiny orifices into the liposome solution, or simply dissolving the gas in the solution by applying pressure and thereafter suddenly releasing the pressure. Another way is to agitate or sonicate the liposome solution in the presence of air or an entrappable gas. Also one can generate the formation of a gas within the solution of liposomes itself, for instance by a gas releasing chemical reaction, e.g., decomposing a dissolved carbonate or bicarbonate by acid. The same effect can be obtained by dissolving under pressure a low boiling liquid, for instance butane, in the aqueous phase and thereafter allowing said liquid to boil by suddenly releasing the pressure.

[0026] Notwithstanding, an advantageous method is to contact the dry surfactant in lamellar or thin film form with air or an adsorbable or entrappable gas before introducing said surfactant into the liquid carrier phase. In this regard, the method can be derived from the technique disclosed in GB-A-2,135,647, i.e., solid microparticles or beads are dipped in a solution of a film forming surfactant (or mixture of surfactants) in a volatile solvent, after which the solvent is evaporated and the beads are left in contact with air (or an adsorbable gas) for a time sufficient for that air to become superficially bound to the surfactant layer. Thereafter, the beads coated with air filled surfactant are put into a carrier liquid, usually water with or without additives, whereby air bubbles will develop within the liquid by gentle mixing, violent agitation being entirely unnecessary. Then the solid beads can be separated, for instance by filtration, from the microbubble suspension which is remarkably stable with time.

[0027] Needless to say that, instead of insoluble beads or spheres, one may use as supporting particles water soluble materials like that disclosed in GB-A-2,134,869 (carbohydrates or hydrophilic polymers), whereby said supporting

particles will eventually dissolve and final separation of a solid becomes unnecessary. Furthermore in this case, the material of the particles can be selected to eventually act as stabilizer or viscosity enhancer wherever desired.

[0028] In a variant of the method, one may also start with dehydrated liposomes, i.e., liposomes which have been prepared normally by means of conventional techniques in the form of aqueous solutions and thereafter dehydrated by usual means, e.g., such as disclosed in U.S. Pat. No. 4,229,360 also incorporated herein by reference. One of the methods for dehydrating liposomes recommended in this reference is freeze-drying (lyophilization), i.e., the liposome solution is frozen and dried by evaporation (sublimation) under reduced pressure. Prior to effecting freeze-drying, a hydrophilic stabilizer compound is dissolved in the solution, for instance a carbohydrate like lactose or sucrose or a hydrophilic polymer like dextran, starch, PVP, PVA and the like. This is useful in the present invention since such hydrophilic compounds also aid in homogenizing the microbubbles size distribution and enhance stability under storage. Actually making very dilute aqueous solutions (0.1-10% by weight) of freeze-dried liposomes stabilized with, for instance, a 5:1 to 10:1 weight ratio of lactose to lipid enables to produce aqueous microbubbles suspensions counting 10^8 - 10^9 microbubbles/ml (size distribution mainly 0.5-10 μ m) which are stable for at least a month (and probably much longer) without significant observable change. And this is obtained by simple dissolution of the air-stored dried liposomes without shaking or any violent agitation. Furthermore, the freeze-drying technique under reduced pressure is very useful because it permits, after drying, to restore the pressure above the dried liposomes with any entrappable gas, i.e., nitrogen, CO₂, argon, methane, freon, etc., whereby after dissolution of the liposomes processed under such conditions suspensions of microbubbles containing the above gases are obtained.

[0029] Microbubbles suspensions formed by applying gas pressure on a dilute solution of laminated lipids in water (0.1-10% by weight) and thereafter suddenly releasing the pressure have an even higher bubble concentration, e.g., in the order of 10_{10} - 10_{11} bubbles/ml. However, the average bubble size is somewhat above 10 μ m, e.g., in the 10-50 μ m range. In this case, bubble size distribution can be narrowed by centrifugation and layer decantation.

[0030] The tensides or surfactants which are convenient in this invention can be selected from all amphipathic compounds capable of forming stable films in the presence of water and gases. The preferred surfactants which can be laminarized include the lecithins (phosphatidylcholine) and other phospholipids, inter alia phosphatidic acid (PA), phosphatidylinositol, phosphatidylethanolamine (PE), phosphatidylserine (PS), phosphatidylglycerol (PG), cardiolipin (CL), sphingomyelins, the plasmogens, the cerebroside, etc. Examples of suitable lipids are the phospholipids in general, for example, natural lecithins, such as egg lecithin or soya bean lecithin, or synthetic lecithins such as saturated synthetic lecithins, for example, dimyristoylphosphatidylcholine, dipalmitoylphosphatidylcholine or distearoylphosphatidylcholine or unsaturated synthetic lecithins, such as dioleoylphosphatidylcholine or dilinoleoylphosphatidylcholine, with egg lecithin or soya bean lecithin being preferred. Additives like cholesterol and other substances (see below)

can be added to one or more of the foregoing lipids in proportions ranging from zero to 50% by weight.

[0031] Such additives may include other surfactants that can be used in admixture with the film forming surfactants and most of which are recited in the prior art discussed in the introduction of this specification. For instance, one may cite free fatty acids, esters of fatty acids with polyoxyalkylene compounds like polyoxypropylene glycol and polyoxyalkylene glycol; ethers of fatty alcohols with polyoxyalkylene glycols; esters of fatty acids with polyoxyalkylated sorbitan; soaps; glycerol-polyalkylene stearate; glycerol-polyoxyethylene ricinoleate; homo- and copolymers of polyalkylene glycols; polyethoxylated soya-oil and castor oil as well as hydrogenated derivatives; ethers and esters of sucrose or other carbohydrates with fatty acids, fatty alcohols, these being optionally polyoxyalkylated; mono- di and triglycerides of saturated or unsaturated fatty acids; glycerides of soya-oil and sucrose. The amount of the non-film forming tensides or surfactants can be up to 50% by weight of the total amount of surfactants in the composition but is preferably between zero and 30%.

[0032] The total amount of surfactants relative to the aqueous carrier liquid is best in the range of 0.01 to 25% by weight but quantities in the range 0.5-5% are advantageous because one always tries to keep the amount of active substances in an injectable solution as low as possible, this being to minimize the introduction of foreign materials into living beings even when they are harmless and physiologically compatible.

[0033] Further optional additives to the surfactants include:

[0034] a) substances which are known to provide a negative charge on liposomes, for example, phosphatidic acid, phosphatidyl-glycerol or dicetyl phosphate;

[0035] b) substances known to provide a positive charge, for example, stearyl amine, or stearyl amine acetate;

[0036] c) substances known to affect the physical properties of the lipid films in a more desirable way; for example, capro-lactam and/or sterols such as cholesterol, ergosterol, phytosterol, sitosterol, sitosterol pyroglutamate, 7-dehydro-cholesterol or lanosterol, may affect lipid films rigidity;

[0037] d) substances known to have antioxidant properties to improve the chemical stability of the components in the suspensions, such as tocopherol, propyl gallate, ascorbyl palmitate, or butylated hydroxy toluene.

[0038] The aqueous carrier in this invention is mostly water with possibly small quantities of physiologically compatible liquids such as isopropanol, glycerol, hexanol and the like (see for instance EP-A-052575). In general the amount of the organic hydrosoluble liquids will not exceed 5-10% by weight.

[0039] The present composition may also contain dissolved or suspended therein hydrophilic compounds and polymers defined generally under the name of viscosity enhancers or stabilizers. Although the presence of such compounds is not compulsory for ensuring stability to the air

or gas bubbles with time in the present dispersions, they are advantageous to give some kind of "body" to the solutions. When desired, the upper concentrations of such additives when totally innocuous can be very high, for instance up to 80-90% by weight of solution with lopamidol and other iodinated X-ray contrast agents. However, with the viscosity enhancers like for instance sugars, e.g., lactose, sucrose, maltose, galactose, glucose, etc. or hydrophilic polymers like starch, dextran, polyvinyl alcohol, polyvinyl-pyrrolidone, dextrin, xanthan or partly hydrolyzed cellulose oligomers, as well as proteins and polypeptides, the concentrations are best between about 1 and 40% by weight, a range of about 5-20% being preferred.

[0040] Like in the prior art, the injectable compositions of this invention can also contain physiologically acceptable electrolytes; an example is an isotonic solution of salt.

[0041] The present invention naturally also includes dry storable pulverulent blends which can generate the present microbubble containing dispersions upon simple admixing with water or an aqueous carrier phase. Preferably such dry blends or formulations will contain all solid ingredients necessary to provide the desired microbubbles suspensions upon the simple addition of water, i.e., principally the surfactants in lamellar form containing trapped or adsorbed therein the air or gas required for microbubble formation, and accessorially the other non-film forming surfactants, the viscosity enhancers and stabilizers and possibly other optional additives. As said before, the air or gas entrapment by the laminated surfactants occurs by simply exposing said surfactants to the air (or gas) at room or superatmospheric pressure for a time sufficient to cause said air or gas to become entrapped within the surfactant. This period of time can be very short, e.g., in the order of a few seconds to a few minutes although over-exposure, i.e., storage under air or under a gaseous atmosphere is in no way harmful. What is important is that air can well contact as much as possible of the available surface of the laminated surfactant, i.e., the dry material should preferably be in a "fluffy" light flowing condition. This is precisely this condition which results from the freeze-drying of an aqueous solution of liposomes and hydrophilic agent as disclosed in U.S. Pat. No. 4,229,360.

[0042] In general, the weight ratio of surfactants to hydrophilic viscosity enhancer in the dry formulations will be in the order of 0.1:10 to 10:1, the further optional ingredients, if any, being present in a ratio not exceeding 50% relative to the total of surfactants plus viscosity enhancers.

[0043] The dry blend formulations of this invention can be prepared by very simple methods. As seen before, one preferred method is to first prepare an aqueous solution in which the film forming lipids are laminarized, for instance by sonication, or using any conventional technique commonly used in the liposome field, this solution also containing the other desired additives, i.e., viscosity enhancers, non-film forming surfactants, electrolyte, etc., and thereafter freeze drying to a free flowable powder which is then stored in the presence of air or an entrappable gas.

[0044] The dry blend can be kept for any period of time in the dry state and sold as such. For putting it into use, i.e., for preparing a gas or air microbubble suspension for ultrasonic imaging, one simply dissolves a known weight of the dry pulverulent formulation in a sterile aqueous phase, e.g., water or a physiologically acceptable medium. The amount

of powder will depend on the desired concentration of bubbles in the injectable product, a count of about 10^8 - 10^9 bubbles/ml being generally that from making a 5-20% by weight solution of the powder in water. But naturally this figure is only indicative, the amount of bubbles being essentially dependent on the amount of air or gas trapped during manufacture of the dry powder. The manufacturing steps being under control, the dissolution of the dry formulations will provide microbubble suspensions with well reproducible counts.

[0045] The resulting microbubble suspensions (bubble in the 0.5-10 μ m range) are extraordinarily stable with time, the count originally measured at start staying unchanged or only little changed for weeks and even months; the only observable change is a kind of segregation, the larger bubbles (around 10 μ m) tending to rise faster than the small ones.

[0046] It has also been found that the microbubbles suspensions of this invention can be diluted with very little loss in the number of microbubbles to be expected from dilution, i.e., even in the case of high dilution ratios, e.g., $1/10^2$ to $1/10^4$, the microbubble count reduction accurately matches with the dilution ratio. This indicates that the stability of the bubbles depends on the surfactant in lamellar form rather than on the presence of stabilizers or viscosity enhancers like in the prior art. This property is advantageous in regard to imaging test reproducibility as the bubbles are not affected by dilution with blood upon injection into a patient.

[0047] Another advantage of the bubbles of this invention versus the microbubbles of the prior art surrounded by a rigid but breakable membrane which may irreversibly fracture under stress is that when the present suspensions are subject to sudden pressure changes, the present bubbles will momentarily contract elastically and then resume their original shape when the pressure is released. This is important in clinical practice when the microbubbles are pumped through the heart and therefore are exposed to alternating pressure pulses.

[0048] The reasons why the microbubbles in this invention are so stable are not clearly understood. Since to prevent bubble escape the buoyancy forces should equilibrate with the retaining forces due to friction, i.e., to viscosity, it is theorized that the bubbles are probably surrounded by the laminated surfactant. Whether this laminar surfactant is in the form of a continuous or discontinuous membrane, or even as closed spheres attached to the microbubbles, is for the moment unknown but under investigation. However the lack of a detailed knowledge of the phenomena presently involved does not prelude full industrial operability of the present invention.

[0049] The bubble suspensions of the present invention are also useful in other medical/diagnostic applications where it is desirable to target the stabilized microbubbles to specific sites in the body following their injection, for instance to thrombi present in blood vessels, to atherosclerotic lesions (plaques) in arteries, to tumor cells, as well as for the diagnosis of altered surfaces of body cavities, e.g., ulceration sites in the stomach or tumors of the bladder. For this, one can bind monoclonal antibodies tailored by genetic engineering, antibody fragments or polypeptides designed to mimic antibodies, bioadhesive polymers, lectins and other site-recognizing molecules to the surfactant layer stabilizing

the microbubbles. Thus monoclonal antibodies can be bound to phospholipid bilayers by the method described by L. D. Leserman, P. Machy and J. Barbet ("Liposome Technology vol. III" p. 29 ed. by G. Gregoriadis, CRC Press 1984). In another approach a palmitoyl antibody is first synthesized and then incorporated in phospholipid bilayers following L. Huang, A. Huang and S. J. Kennel ("Liposome Technology vol. III" p. 51 ed. by G. Gregoriadis, CRC Press 1984). Alternatively, some of the phospholipids used in the present invention can be carefully selected in order to obtain preferential uptake in organs or tissues or increased half-life in blood. Thus GM1 gangliosides- or phosphatidylinositol-containing liposomes, preferably in addition to cholesterol, will lead to increased, half-lives in blood after intravenous administration in analogy with A. Gabizon, D. Papahadjopoulos, Proc. Natl Acad. Sci USA 85 (1988) 6949.

[0050] The gases in the microbubbles of the present invention can include, in addition to current innocuous physiologically acceptable gases like CO₂, nitrogen, N₂O, methane, butane, freon (organic compounds containing one or more carbon atoms and fluorine such as CF₄, C₂F₆, C₃F₈ or C₄F₈ and the like) and mixtures thereof, radioactive gases such as ¹³³Xe or ⁸¹Kr are of particular interest in nuclear medicine for blood circulation measurements, for lung scintigraphy, etc.

[0051] The invention described up until this point can be further elucidated by the description of the following representative (but not limiting) embodiments, numbered 1-27:

[0052] 1. A composition adapted for injection into the bloodstream and body cavities of living beings, e.g., for the purpose of ultrasonic echography consisting of a suspension of air or gas microbubbles in a physiologically acceptable aqueous carrier phase comprising from about 0.01 to about 20% by weight of one or more dissolved or dispersed surfactants, characterized in that at least one of the surfactants is a film forming surfactant present in the composition at least partially in lamellar or laminar form.

[0053] 2. The composition of embodiment 1, characterized in that the lamellar surfactant is in the form of mono- or pluri-molecular membrane layers.

[0054] 3. The composition of embodiment 1, characterized in that the lamellar surfactant is in the form of liposome vesicles.

[0055] 4. The composition of embodiment 1, characterized in that it essentially consists of a liposome solution containing air or gas microbubbles developed therein.

[0056] 5. The composition of embodiment 4, characterized in that the size of most of both liposomes and microbubbles is below 50 μ m, preferably below 10 μ m.

[0057] 6. The composition of embodiment 1, containing about 10⁸-10⁹ bubbles of 0.5-10 μ m size/ml, said concentration showing little or substantially no variability under storage for at least a month.

[0058] 7. The composition of embodiment 1, characterized in that the surfactants are selected from phospholipids including the lecithins such as phosphatidic acid, phosphatidylcholine, phosphatidylethanolamine, phosphatidylserine, phosphatidylglycerol, phosphatidylinositol, cardiolipin and sphingomyelin.

[0059] 8. The composition of embodiment 7, characterized in further containing substances affecting the properties of liposomes selected from phosphatidylglycerol, dicytlylphosphate, cholesterol, ergosterol, phytosterol, sitosterol, lanosterol, tocopherol, propyl gallate, ascorbyl palmitate and butylated hydroxytoluene.

[0060] 9. The composition of embodiment 1, further containing dissolved viscosity enhancers or stabilizers selected from linear and cross-linked poly- and oligo-saccharides, sugars, hydrophilic polymers and iodinated compounds such as lopamidol in a weight ratio to the surfactants comprised between about 1:5 to 100:1.

[0061] 10. The composition of embodiment 1, in which the surfactants comprise up to 50% by weight of non-laminar surfactants selected from fatty acids, esters, and ethers of fatty acids and alcohols with polyols such as polyalkylene glycols, polyalkylenated sugars and other carbohydrates, and polyalkylenated glycerol.

[0062] 11. A method for the preparation of the suspensions of embodiment 1, characterized by the following steps:

[0063] (a) selecting at least one film forming surfactant and converting it into lamellar form;

[0064] (b) contacting the surfactant in lamellar form with air or an adsorbable or entrappable gas for a time sufficient for that air or gas to become bound by said surfactant; and

[0065] (c) admixing the surfactant in lamellar form with an aqueous liquid carrier, whereby a stable dispersion of air or gas microbubbles in said liquid carrier will result.

[0066] 12. The method of embodiment 11, in which step (c) is brought about before step (b), the latter being effected by introducing pressurized air or gas into the liquid carrier and thereafter releasing the pressure.

[0067] 13. The method of embodiment 11, in which step (c) is brought about by gentle mixing of the components, no shaking being necessary, whereby the air or gas bound to the lamellar surfactant in step (b) will develop into a suspension of stable microbubbles.

[0068] 14. The method of embodiments 11 or 12, in which the liquid carrier contains dissolved therein stabilizer compounds selected from hydrosoluble proteins, polypeptides, sugars, poly- and oligo-saccharides and hydrophilic polymers.

[0069] 15. The method of embodiment 11, in which the conversion of step (a) is effected by coating the surfactant onto particles of soluble or insoluble materials; step (b) is effected by letting the coated particles stand for a while under air or a gas; and step (c) is effected by admixing the coated particles with an aqueous liquid carrier.

[0070] 16. The method of embodiment 11, in which the conversion of step (a) is effected by sonicating or homogenizing under high pressure an aqueous solution of film forming lipids, this operation leading, at least partly, to the formation of liposomes.

[0071] 17. The method of embodiment 16, in which step (b) is effected by freeze-drying the liposome containing solution, the latter optionally containing hydrophilic stabi-

lizers and contacting the resulting freeze-dried product with air or gas for a period of time.

[0072] 18. The method of embodiments 16 and 17, in which the water solution of film forming lipids also contains viscosity enhancers or stabilizers selected from hydrophilic polymers and carbohydrates in weight ratio relative to the lipids comprised between 1:5 and 100:1.

[0073] 19. A dry pulverulent formulation which, upon dissolution in water, will form an aqueous suspension of microbubbles for ultrasonic echography, characterized in containing one or more film forming surfactants in laminar form and hydrosoluble stabilizers.

[0074] 20. The dry formulation of embodiment 19, in which the surfactants in laminar form are in the form of fine layers deposited on the surface of soluble or insoluble solid particulate material.

[0075] 21. The dry formulation of embodiment 20, in which the insoluble solid particles are glass or polymer beads.

[0076] 22. The dry formulation of embodiment 20, in which the soluble particles are made of hydrosoluble carbohydrates, polysaccharides, synthetic polymers, albumin, gelatin or iopamidol.

[0077] 23. The dry formulation of embodiment 19, which comprises freeze-dried liposomes.

[0078] 24. The use of the injectable composition of embodiment 1 for ultrasonic echography.

[0079] 25. The use of the injectable composition of embodiments 1-10 for transporting in the blood-stream or body cavities bubbles of foreign gases active therapeutically or diagnostically.

[0080] 26. The composition of embodiment 4, in which the surfactant comprises, bound thereto, bioactive species designed for specific targeting purposes, e.g., for immobilizing the bubbles in specifically defined sites in the circulatory system, or in organs, or in tissues.

[0081] 27. The composition of embodiment 4, in which the surfactant comprises, bound thereto, bioactive species selected from monoclonal antibodies, antibody fragments or polypeptides designed to mimic antibodies, bioadhesive polymers, lectins and other receptor recognizing molecules.

[0082] The following Examples further illustrate the invention from a practical standpoint.

[0083] Echogenic Measurements

[0084] Echogenicity measurements were performed in a pulse—echo system made of a plexiglas specimen holder (diameter 30 mm) and a transducer holder immersed in a constant temperature water bath, a pulser-receiver (Accutron M3010S) with for the receiving part an external pre-amplifier with a fixed gain of 40 dB and an internal amplifier with adjustable gain from -40 to +40 dB. A 10 MHz low-pass filter was inserted in the receiving part to improve the signal to noise ratio. The A/D board in the IBM PC was a Sonotek STR 832. Measurements were carried out at 2.25, 3.5, 5 and 7.5 MHz.

EXAMPLE 1

[0085] A liposome solution (50 mg lipids per ml) was prepared in distilled water by the REV method (see F. Szoka

Jr. and D. Papahadjopoulos, Proc. Natl. Acad. Sci. USA 75 (1978) 4194) using hydrogenated soya lecithin (NC 95 H, Nattermann Chemie, Koln, W. Germany) and dicetylphosphate in a molar ratio 9/1. This liposome preparation was extruded at 65° C. (to calibrate the vesicle size) through a 1 μ m polycarbonate filter (Nucleopore). Two ml of this solution were admixed with 5 ml of a 75% iopamidol solution in water and 0.4 ml of air and the mixture was forced back and forth through a two syringe system as disclosed in DE-A-3529195, while maintaining continuously a slight overpressure. This resulted in the formation of a suspension of microbubbles of air in the liquid (10^5 - 10^6 bubbles per ml, bubble size 1-20 μ m as estimated by light microscopy) which was stable for several hours at room temperature. This suspension gave a strong echo signal when tested by ultrasonic echography at 7.5, 5, 3.5 and 2.25 MHz.

EXAMPLE 2

[0086] A distilled water solution (100 ml) containing by weight 2% of hydrogenated soya lecithin and dicetylphosphate in a 9/1 molar ratio was sonicated for 15 min at 60°-65° C. with a Branson probe sonifier (Type 250). After cooling, the solution was centrifuged for 15 min at 10,000 g and the supernatant was recovered and lactose added to make a 7.5% b.w. solution. The solution was placed in a tight container in which a pressure of 4 bar of nitrogen was established for a few minutes while shaking the container. Afterwards, the pressure was released suddenly whereby a highly concentrated bubble suspension was obtained (10^{10} - 10^{11} bubbles/ml). The size distribution of the bubbles was however wider than in Example 1, i.e., from about 1 to 50 μ m. The suspension was very stable but after a few days a segregation occurred in the standing phase, the larger bubbles tending to concentrate in the upper layers of the suspension.

EXAMPLE 3

[0087] Twenty g of glass beads (diameter about 1 mm) were immersed into a solution of 100 mg of dipalmitoylphosphatidylcholine (Fluka A. G. Buchs) in 10 ml of chloroform. The beads were rotated under reduced pressure in a rotating evaporator until all CHCl_3 had escaped. Then the beads were further rotated under atmospheric pressure for a few minutes and 10 ml of distilled water were added. The beads were removed and a suspension of air microbubbles was obtained which was shown to contain about 10^6 bubbles/ml after examination under the microscope. The average size of the bubbles was about 3-5 μ m. The suspension was stable for several days at least.

EXAMPLE 4

[0088] A hydrogenated soya lecithin/dicetylphosphate suspension in water was laminarized using the REV technique as described in Example 1. Two ml of the liposome preparation were added to 8 ml of 15% maltose solution in distilled water. The resulting solution was frozen at -30° C., then lyophilized under 0.1 Torr. Complete sublimation of the ice was obtained in a few hours. Thereafter, air pressure was restored in the evacuated container so that the lyophilized powder became saturated with air in a few minutes.

[0089] The dry powder was then dissolved in 10 ml of sterile water under gentle mixing, whereby a microbubble

suspension (10^8 - 10^9 microbubbles per ml, dynamic viscosity <20 mPa.s) was obtained. This suspension containing mostly bubbles in the 1 - $5\text{ }\mu\text{m}$ range was stable for a very long period, as numerous bubbles could still be detected after 2 months standing. This microbubble suspension gave a strong response in ultrasonic echography. If in this example the solution is frozen by spraying in air at -30° to -70° C. to obtain a frozen snow instead of a monolithic block and the snow is then evaporated under vacuum, excellent results are obtained.

EXAMPLE 5

[0090] Two ml samples of the liposome solution obtained as described in Example 4 were mixed with 10 ml of an 5% aqueous solution of gelatin (sample 5A), human albumin (sample SB), dextran (sample 5C) and iopamidol (sample 5D). All samples were lyophilized. After lyophilization and introduction of air, the various samples were gently mixed with 20 ml of sterile water. In all cases, the bubble concentration was above 10^8 bubbles per ml and almost all bubbles were below $10\text{ }\mu\text{m}$. The procedure of the foregoing Example was repeated with 9 ml of the liposome preparation (450 mg of lipids) and only one ml of a 5% human albumin solution. After lyophilization, exposure to air and addition of sterile water (20 ml), the resulting solution contained 2×10^8 bubbles per ml, most of them below $10\text{ }\mu\text{m}$.

EXAMPLE 6

[0091] Lactose (500 mg), finely milled to a particle size of 1 - $3\text{ }\mu\text{m}$, was moistened with a chloroform (5 ml) solution of 100 mg of dimyristoylphosphatidylcholine/cholesterol/dipalmitoylphosphatidic acid (from Fluka) in a molar ratio of 4:1:1 and thereafter evaporated under vacuum in a rotating evaporator. The resulting free flowing white powder was rotated a few minutes under nitrogen at normal pressure and thereafter dissolved in 20 ml of sterile water. A microbubble suspension was obtained with about 10^5 - 10^6 microbubbles per ml in the 1 - $10\text{ }\mu\text{m}$ size range as ascertained by observation under the microscope. In this Example, the weight ratio of coated surfactant to water-soluble carrier was 1:5. Excellent results (10^7 - 10^8 microbubbles/ml) are also obtained when reducing this ratio to lower values, i.e., down to 1:20, which will actually increase the surfactant efficiency for the intake of air, that is, this will decrease the weight of surfactant necessary for producing the same bubble count.

EXAMPLE 7

[0092] An aqueous solution containing 2% of hydrogenated soya lecithin and 0.4% of Pluronic® F68 (a non ionic polyoxyethylenepolyoxypropylene copolymer surfactant) was sonicated as described in Example 2. After cooling and centrifugation, 5 ml of this solution were added to 5 ml of a 15% maltose solution in water. The resulting solution was frozen at -30° C. and evaporated under 0.1 Torr. Then air pressure was restored in the vessel containing the dry powder. This was left to stand in air for a few seconds, after which it was used to make a 10% by weight aqueous solution which showed under the microscope to be a suspension of very tiny bubbles (below $10\text{ }\mu\text{m}$); the bubble concentration was in the range of 10^7 bubbles per ml. This preparation gave a very strong response in ultrasonic echography at 2.25, 3.5, 5 and 7.5 MHz.

EXAMPLE 8

[0093] Two-dimensional echocardiography was performed in an experimental dog following peripheral vein injection of 0.1-2 ml of the preparation obtained in Example 4. Opacification of the left heart with clear outlining of the endocardium was observed, thereby confirming that the microbubbles (or at least a significant part of them) were able to cross the pulmonary capillary circulation.

EXAMPLE 9

[0094] A phospholipid/maltose lyophilized powder was prepared as described in Example 4. However, at the end of the lyophilization step, a ^{133}Xe containing gas mixture was introduced in the evacuated container instead of air. A few minutes later, sterile water was introduced and after gentle mixing a microbubble suspension containing ^{133}Xe in the gas phase was produced. This microbubble suspension was injected into living bodies to undertake investigations requiring use of ^{133}Xe as tracer. Excellent results were obtained.

EXAMPLE 10 (COMPARATIVE)

[0095] In U.S. Pat. No. 4,900,540, Ryan et al. disclose gas filled liposomes for ultrasonic investigations. According to the citation, liposomes are formed by conventional means but with the addition of a gas or gas precursor in the aqueous composition forming the liposome core (col. 2, lines 15-27).

[0096] Using a gas precursor (bicarbonate) is detailed in Examples 1 and 2 of the reference. Using an aqueous carrier with an added gas for encapsulating the gas in the liposomes (not exemplified by Ryan et al.) will require that the gas be in the form of very small bubbles, i.e., of size similar or smaller than the size of the liposome vesicles.

[0097] Aqueous media in which air can be entrapped in the form of very small bubbles (2.5 - $5\text{ }\mu\text{m}$) are disclosed in M. W. Keller et al., J. Ultrasound Med. 5 (1986), 413-498.

[0098] A quantity of 126 mg of egg lecithin and 27 mg of cholesterol were dissolved in 9 ml of chloroform in a 200 ml round bottom flask. The solution of lipids was evaporated to dryness on a Rotavapor whereby a film of the lipids was formed on the walls of the flask. A 10 ml of a 50% by weight aqueous dextrose solution was sonicated for 5 min according to M. W. Keller et al. (ibid) to generate air microbubbles therein and the sonicated solution was added to the flask containing the film of lipid, whereby hand agitation of the vessel resulted into hydration of the phospholipids and formation of multilamellar liposomes within the bubbles containing carrier liquid.

[0099] After standing for a while, the resulting liposome suspension was subjected to centrifugation under 5000 g for 15 min to remove from the carrier the air not entrapped in the vesicles. It was also expected that during centrifugation, the air filled liposomes would segregate to the surface by buoyancy.

[0100] After centrifugation the tubes were examined and showed a bottom residue consisting of agglomerated dextrose filled liposomes and a clear supernatant liquid with substantially no bubbles left. The quantity of air filled liposomes having risen by buoyancy was negligibly small and could not be ascertained.

EXAMPLE 11 (COMPARATIVE)

[0101] An injectable contrast composition was prepared according to Ryan (U.S. Pat. No. 4,900,540, col. 3, Example 1). Egg lecithin (126 mg) and cholesterol (27 mg) were dissolved in 9 ml of diethylether. To the solution were added 3 ml of 0.2 molar aqueous bicarbonate and the resulting two phase systems was sonicated until becoming homogeneous. The mixture was evaporated in a Rotavapor apparatus and 3 ml of 0.2 molar aqueous bicarbonate were added.

[0102] A 1 ml portion of the liposome suspension was injected into the jugular vein of an experimental rabbit, the animal being under condition for heart ultrasonic imaging using an Acuson 128-XP5 ultrasonic imager (7.5 transducer probe for imaging the heart). The probe provided a cross-sectional image of the right and left ventricles (mid-papillary muscle). After injection, a light and transient (a few seconds) increase in the outline of the right ventricle was observed. The effect was however much inferior to the effect observed using the preparation of Example 4. No improvement of the imaging of the left ventricle was noted which probably indicates that the CO₂ loaded liposomes did not pass the pulmonary capillaries barrier.

Further Methods of the Invention and Gases Used Therein

[0103] Despite the many progresses achieved regarding the stability under storage of aqueous microbubble suspensions, this being either in the precursor or final preparation stage, there still remained until now the problem of vesicle durability when the suspensions are exposed to overpressure, e.g., pressure variations such as that occurring after injection in the blood stream of a patient and consecutive to heart pulses, particularly in the left ventricle. Actually, the present inventors have observed that, for instance in anaesthetized rabbits, the pressure variations are not sufficient to substantially alter the bubble count for a period of time after injection. In contrast, in dogs and human patients, typical microbubbles or microballoons filled with common gases such as air, methane or CO₂ will collapse completely in a matter of seconds after injection due to the blood pressure effect. It became hence important to solve the problem and to increase the useful life of suspensions of microbubbles and membrane bounded microballoons under pressure in order to ensure that echographic measurements can be performed in vivo safely and reproducibly.

[0104] It should be mentioned at this stage that another category of echogenic image enhancing agents has been proposed which resist overpressures as they consist of plain microspheres with a porous structure, such porosity containing air or a gas. Such microspheres are disclosed for instance in WO-A-91/12823 (Delta Biotechnology), EP-A-0327490 (Schering) and EP-A-0458079 (Hoechst). The drawback with the plain porous microspheres is that the encapsulated gas-filled free space is generally too small for good echogenic response and the spheres lack adequate elasticity. Hence the preference generally remains with the hollow microvesicles and a solution to the collapsing problem was searched.

[0105] This problem has now been solved by using gases or gas mixtures in conformity with the criteria outlined in the embodiments shown below. Briefly, it has been found that when the echogenic microvesicles are made in the presence

of a gas, respectively are filled at least in part with a gas, having physical properties in conformity with the equation below, then the microvesicles remarkably resist pressure >60 Torr after injection for a time sufficient to obtain reproducible echographic measurements:

$$\frac{s_{\text{gas}}}{s_{\text{air}}} \times \frac{\sqrt{Mw_{\text{air}}}}{\sqrt{Mw_{\text{gas}}}} \leq 1$$

[0106] In the foregoing equation, "s" designates the solubilities in water expressed as the "Bunsen" coefficients, i.e., as volume of gas dissolved by unit volume of water under standard conditions (1 bar, 25° C.), and under partial pressure of the given gas of 1 atm (see the Gas Encyclopaedia, Elsevier 1976). Since, under such conditions and definitions, the solubility of air is 0.0167, and the square root of its average molecular weight (Mw) is 5.39, the above relation simplifies to:

$$s_{\text{gas}}/\sqrt{Mw_{\text{gas}}} \leq 0.0031$$

[0107] In the Examples to be found hereafter there is disclosed the testing of echogenic microbubbles and microballoons (see the Tables) filled with a number of different gases and mixtures thereof, and the corresponding resistance thereof to pressure increases, both in vivo and in vitro. In the Tables, the water solubility factors have also been taken from the aforesaid Gas Encyclopaedia from "L'Air Liquide", Elsevier Publisher (1976).

[0108] The microvesicles in aqueous suspension containing gases according to the invention include most microbubbles and microballoons disclosed until now for use as contrast agents for echography. The preferred microballoons are those disclosed in EP-A-0324938, PCT/EP91/01706 and EP-A-0458745; the preferred microbubbles are those of the compositions disclosed herein (e.g., supra) and in PCT/EP91/00620; these microbubbles are advantageously formed from an aqueous liquid and a dry powder (microvesicle precursors) containing lamellarized freeze-dried phospholipids and stabilizers; the microbubbles are developed by agitation of this powder in admixture with the aqueous liquid carrier. The microballoons of EP-A-0458745 have a resilient interfacially precipitated polymer membrane of controlled porosity. They are generally obtained from emulsions into microdroplets of polymer solutions in aqueous liquids, the polymer being subsequently caused to precipitate from its solution to form a fibrogenic membrane at the droplet/liquid interface, which process leads to the initial formation of liquid-filled microvesicles, the liquid core thereof being eventually substituted by a gas.

[0109] In order to carry out the method of the present invention, i.e., to form or fill the microvesicles, whose suspensions in aqueous carriers constitute the desired echogenic additives, with the gases according to the foregoing relation, one can either use, as a first embodiment, a two step route consisting of (1) making the microvesicles from appropriate starting materials by any suitable conventional technique in the presence of any suitable gas, and (2) replacing this gas originally used (first gas) for preparing the microvesicles with a new gas (second gas) according to the invention (gas exchange technique).

[0110] Otherwise, according to a second embodiment, one can directly prepare the desired suspensions by suitable usual methods under an atmosphere of the new gas according to the invention.

[0111] If one uses the two-step route, the initial gas can be first removed from the vesicles (for instance by evacuation under suction) and thereafter replaced by bringing the second gas into contact with the evacuated product, or alternatively, the vesicles still containing the first gas can be contacted with the second gas under conditions where the second gas will displace the first gas from the vesicles (gas substitution). For instance, the vesicle suspensions, or preferably precursors thereof (precursors here may mean the materials the microvesicle envelopes are made of, or the materials which, upon agitation with an aqueous carrier liquid, will generate or develop the formation of microbubbles in this liquid), can be exposed to reduced pressure to evacuate the gas to be removed and then the ambient pressure is restored with the desired gas for substitution. This step can be repeated once or more times to ensure complete replacement of the original gas by the new one. This embodiment applies particularly well to precursor preparations stored dry, e.g., dry powders which will regenerate or develop the bubbles of the echogenic additive upon admixing with an amount of carrier liquid. Hence, in one preferred case where microbubbles are to be formed from an aqueous phase and dry laminarized phospholipids, e.g., powders of dehydrated lyophilized liposomes plus stabilizers, which powders are to be subsequently dispersed under agitation in a liquid aqueous carrier phase, it is advantageous to store this dry powder under an atmosphere of a gas selected according to the invention. A preparation of such kind will keep indefinitely in this state and can be used at any time for diagnosis, provided it is dispersed into sterile water before injection.

[0112] Otherwise, and this is particularly so when the gas exchange is applied to a suspension of microvesicles in a liquid carrier phase, the latter is flushed with the second gas until the replacement (partial or complete) is sufficient for the desired purpose. Flushing can be effected by bubbling from a gas pipe or, in some cases, by simply sweeping the surface of the liquid containing the vesicles under gentle agitation with a stream (continuous or discontinuous) of the new gas. In this case, the replacement gas can be added only once in the flask containing the suspension and allowed to stand as such for a while, or it can be renewed one or more times in order to assure that the degree of renewal (gas exchange) is more or less complete.

[0113] Alternatively, in a second embodiment as said before, one will effect the full preparation of the suspension of the echogenic additives starting with the usual precursors thereof (starting materials), as recited in the prior art and operating according to usual means of said prior art, but in the presence of the desired gases or mixture of gases according to the invention instead of that of the prior art which usually recites gases such as air, nitrogen, CO₂ and the like.

[0114] It should be noted that in general the preparation mode involving one first type of gas for preparing the microvesicles and, thereafter, substituting the original gas by a second kind of gas, the latter being intended to confer different echogenic properties to said microvesicles, has the

following advantage: As will be best seen from the results in the Examples hereinafter, the nature of the gas used for making the microvesicles, particularly the microballoons with a polymer envelope, has a definitive influence on the overall size (i.e., the average mean diameter) of said microvesicles; for instance, the size of microballoons prepared under air with precisely set conditions can be accurately controlled to fall within a desired range, e.g., the 1 to 10 μ m range suitable for echographing the left and right heart ventricles. This not so easy with other gases, particularly the gases in conformity with the requirements of the present invention; hence, when one wishes to obtain microvesicles in a given size range but filled with gases the nature of which would render the direct preparation impossible or very hard, one will much advantageously rely on the two-steps preparation route, i.e., one will first prepare the microvesicles with a gas allowing more accurate diameter and count control, and thereafter replace the first gas by a second gas by gas exchange.

[0115] In the description of the Experimental part that follows (Examples), gas-filled microvesicles suspended in water or other aqueous solutions have been subjected to pressures over that of ambient. It was noted that when the overpressure reached a certain value (which is generally typical for a set of microsphere parameters and working conditions like temperature, compression rate, nature of carrier liquid and its content of dissolved gas (the relative importance of this parameter will be detailed hereinafter), nature of gas filler, type of echogenic material, etc.), the microvesicles started to collapse, the bubble count progressively decreasing with further increasing the pressure until a complete disappearance of the sound reflector effect occurred. This phenomenon was better followed optically, (nephelometric measurements) since it is paralleled by a corresponding change in optical density, i.e., the transparency of the medium increases as the bubble progressively collapse. For this, the aqueous suspension of microvesicles (or an appropriate dilution thereof was placed in a spectrophotometric cell maintained at 25° C. (standard conditions) and the absorbance was measured continuously at 600 or 700 nm, while a positive hydrostatic overpressure was applied and gradually increased. The pressure was generated by means of a peristaltic pump (Gilson's Mini-puls) feeding a variable height liquid column connected to the spectrophotometric cell, the latter being sealed leak-proof. The pressure was measured with a mercury manometer calibrated in Torr. The compression rate with time was found to be linearly correlated with the pump's speed (rpm's). The absorbance in the foregoing range was found to be proportional to the microvesicle concentration in the carrier liquid.

[0116] The invention will now be further described with reference to **FIG. 1** which is a graph which relates the bubble concentration (bubble count), expressed in terms of optical density in the aforementioned range, and the pressure applied over the bubble suspension. The data for preparing the graph are taken from the experiments reported in Example 15.

[0117] **FIG. 1** shows graphically that the change of absorbance versus pressure is represented by a sigmoid-shaped curve. Up to a certain pressure value, the curve is nearly flat which indicates that the bubbles are stable. Then, a relatively fast absorbance drop occurs, which indicates the existence of a relatively narrow critical region within which any pressure

increase has a rather dramatic effect on the bubble count. When all the microvesicles have disappeared, the curve levels off again. A critical point on this curve was selected in the middle between the higher and lower optical readings, i.e., intermediate between the "full"-bubble (OD max) and the "no"-bubble (OD min) measurements, this actually corresponding where about 50% of the bubbles initially present have disappeared, i.e., where the optical density reading is about half the initial reading, this being set, in the graph, relative to the height at which the transparency of the pressurized suspension is maximal (base line). This point which is also in the vicinity where the slope of the curve is maximal is defined as the critical pressure PC. It was found that for a given gas, PC does not only depend on the aforementioned parameters but also, and particularly so, on the actual concentration of gas (or gases) already dissolved in the carrier liquid: the higher the gas concentration, the higher the critical pressure. In this connection, one can therefore increase the resistance to collapse under pressure of the microvesicles by making the carrier phase saturated with a soluble gas, the latter being the same, or not, (i.e., a different gas) as the one that fills the vesicles. As an example, air-filled microvesicles could be made very resistant to overpressures (>120 Torr) by using, as a carrier liquid, a saturated solution of CO₂. Unfortunately, this finding is of limited value in the diagnostic field since once the contrast agent is injected to the bloodstream of patients (the gas content of which is of course outside control), it becomes diluted therein to such an extent that the effect of the gas originally dissolved in the injected sample becomes negligible.

[0118] Another readily accessible parameter to reproducibly compare the performance of various gases as microsphere fillers is the width of the pressure interval (AP) limited by the pressure values under which the bubble counts (as expressed by the optical densities) is equal to the 75% and 25% of the original bubble count. Now, it has been surprisingly found that for gases where the pressure difference $\Delta P = P_{25} - P_{75}$ exceeds a value of about 25-30 Torr, the killing effect of the blood pressure on the gas-filled microvesicles is minimized, i.e., the actual decrease in the bubble count is sufficiently slow not to impair the significance, accuracy and reproducibility of echographic measurements.

[0119] It was found, in addition, that the values of PC and AP also depend on the rate of rising the pressure in the test experiments illustrated by FIG. 1, i.e., in a certain interval of pressure increase rates (e.g., in the range of several tens to several hundreds of Torr/min), the higher the rate, the larger the values for PC and AP. For this reason, the comparisons effected under standard temperature conditions were also carried out at the constant increase rate of 100 Torr/min. It should however be noted that this effect of the pressure increase rate on the measure of the PC and AP values levels off for very high rates; for instance the values measured under rates of several hundreds of Torr/min are not significantly different from those measured under conditions ruled by heart beats.

[0120] Although the very reasons why certain gases obey the aforementioned properties, while others do not, have not been entirely clarified, it would appear that some relation possibly exists in which, in addition to molecular weight and water solubility, dissolution kinetics, and perhaps other

parameters, are involved. However these parameters need not be known to practice the present invention since gas eligibility can be easily determined according to the aforediscussed criteria.

[0121] The gaseous species which particularly suit the invention are, for instance, halogenated hydrocarbons like the freons (organic compounds containing one or more carbon atoms and fluorine such as CF₄, C₂F₆, C₃F₈ or C₄F₈ and the like) and stable fluorinated chalcogenides like SF₆, SeF₆ and the like.

[0122] It has been mentioned above that the degree of gas saturation of the liquid used as carrier for the microvesicles according to the invention has an importance on the vesicle stability under pressure variations. Indeed, when the carrier liquid in which the microvesicles are dispersed for making the echogenic suspensions of the invention is saturated at equilibrium with a gas, preferably the same gas with which the microvesicles are filled, the resistance of the microvesicles to collapse under variations of pressure is markedly increased. Thus, when the product to be used as a contrast agent is sold dry to be mixed just before use with the carrier liquid (see for instance the products disclosed in PCT/EP91/00620 mentioned hereinbefore), it is quite advantageous to use, for the dispersion, a gas saturated aqueous carrier. Alternatively, when marketing ready-to-use microvesicle suspensions as contrast agents for echography, one will advantageously use as the carrier liquid for the preparation a gas saturated aqueous solution; in this case the storage life of the suspension will be considerably increased and the product may be kept substantially unchanged (no substantial bubble count variation) for extended periods, for instance several weeks to several months, and even over a year in special cases. Saturation of the liquid with a gas may be effected most easily by simply bubbling the gas into the liquid for a period of time at room temperature.

[0123] The invention described herein can be further elucidated by the description of the following representative (but not limiting) embodiments, numbered 1-18:

[0124] 1. A method for imparting resistance against collapsing to contrast agents for ultrasonic echography which consist of gas-filled microvesicles in suspension in aqueous liquid carrier phases, i.e., either microbubbles bounded by an evanescent gas/liquid interfacial closed surface, or microballoons bounded by a material envelope, said collapsing resulting, at least in part, from pressure increases effective, e.g., when the said suspensions are injected into the blood stream of patients, said method comprising forming said microvesicles in the presence of a gas, or if the microvesicles are already made filling them with this gas, which is a physiologically acceptable gas, or gas mixture, at least a fraction of which has a solubility in water expressed in liters of gas by liter of water under standard conditions divided by the square root of the molecular weight in daltons which does not exceed 0.003.

[0125] 2. The method of embodiment 1, which is carried out in two steps, in the first step the microvesicles or dry precursors thereof are initially prepared under an atmosphere of a first gas, then in the second step at least a fraction of the first gas is substantially substituted by a second gas, the latter being said physiologically acceptable gas.

[0126] 3. The method of embodiment 1, in which the physiologically acceptable gas used is selected from SF₆ or

freon such as CF_4 , CBrF_3 , C_4F_8 , CClF_3 , CCl_2F_2 , C_2F_6 , C_2ClF_5 , CBrClF_2 , $\text{C}_2\text{Cl}_2\text{F}_4$, CBr_2F_2 and C_4F_{10} .

[0127] 4. The method of embodiment 2, in which the gas used in the first step is a kind that allows effective control of the average size and concentration of the microvesicles in the carrier liquid, and the physiologically acceptable gas added in the second step ensures prolonged useful echogenic life to the suspension for in vivo ultrasonic imaging.

[0128] 5. The method of embodiment 1, in which the aqueous phase carrying the microbubbles contains dissolved film-forming surfactants in lamellar or laminar form, said surfactants stabilizing the microbubbles boundary at the gas to liquid interface.

[0129] 6. The method of embodiment 5, in which said surfactants comprise one or more phospholipids.

[0130] 7. The method of embodiment 6, in which at least part of the phospholipids are in the form of liposomes.

[0131] 8. The method of embodiment 6, in which at least one of the phospholipids is a diacylphosphatidyl compound wherein the acyl group is a C_{16} fatty acid residue or a higher homologue thereof.

[0132] 9. The method of embodiments 1 and 2, in which the microballoon material envelope is made of an organic polymeric membrane.

[0133] 10. The method of embodiment 9, in which the polymers of the membrane are selected from polylactic or polyglycolic acid and their copolymers, reticulated serum albumin, reticulated haemoglobin, polystyrene, and esters of polyglutamic and polyaspartic acids.

[0134] 11. The method of embodiment 1, in which the forming of the microvesicles with said physiologically acceptable gas is effected by alternately subjecting dry precursors thereof to reduced pressure and restoring the pressure with said gas, and finally dispersing the precursors in a liquid carrier.

[0135] 12. The method of embodiment 1, in which the filling of the microballoons with said physiologically acceptable gas is effected by simply flushing the suspension with said gas under ambient pressure.

[0136] 13. The method of embodiment 1, which comprises making the microvesicles by any standard method known in the art but operating under an atmosphere composed at least in part of said gas.

[0137] 14. Suspensions of gas filled microvesicles distributed in an aqueous carrier liquid to be used as contrast agents in ultrasonic echography, characterized in that the gas is physiologically acceptable and such that at least a portion thereof has a solubility in water, expressed in liter of gas by liter of water under standard conditions, divided by the square root of the molecular weight which does not exceed 0.003.

[0138] 15. The aqueous suspensions of embodiment 14, characterized in that the gas is such that the pressure difference ΔP between those pressures which, when applied under standard conditions and at a rate of about 100 Torr/min to the suspension cause the collapsing of about 75%, respectively 25%, of the microvesicles initially present, is at least 25 Torr.

[0139] 16. Aqueous suspensions according to embodiment 14, in which the microvesicles are microbubbles filled with said physiologically acceptable gas suspended in an aqueous carrier liquid containing phospholipids whose fatty acid residues contain 16 carbons or more.

[0140] 17. Contrast agents for echography in precursor form consisting of a dry powder comprising lyophilized liposomes and stabilizers, this powder being dispersible in aqueous liquid carriers to form echogenic suspensions of gas-filled microbubbles, characterized in that it is stored under an atmosphere comprising a physiologically acceptable gas whose solubility in water, expressed in liter of gas by liter of water under standard conditions, divided by the square root of the molecular weight does not exceed 0.003.

[0141] 18. The contrast agent precursors of embodiment 17, in which the liposomes comprise phospholipids whose fatty acid residues have 16 or more carbon atoms.

[0142] The following Examples further illustrate various aspects of the invention.

EXAMPLE 12

[0143] Albumin microvesicles filled with air or various gases were prepared as described in EP-A-0324938 using a 10 ml calibrated syringe filled with a 5% human serum albumin (HSA) obtained from the Blood Transfusion Service, Red-Cross Organization, Bern, Switzerland. A sonicator probe (Sonifier Model 250 from Branson Ultrasonic Corp, USA) was lowered into the solution down to the 4 ml mark of the syringe and sonication was effected for 25 sec (energy setting=8). Then the sonicator probe was raised above the solution level up to the 6 ml mark and sonication was resumed under the pulse mode (cycle=0.3) for 40 sec. After standing overnight at 4° C., a top layer containing most of the microvesicles had formed by buoyancy and the bottom layer containing unused albumin debris of denatured protein and other insolubles was discarded. After resuspending the microvesicles in fresh albumin solution the mixture was allowed to settle again at room temperature and the upper layer was finally collected. When the foregoing sequences were carried out under the ambient atmosphere, air filled microballoons were obtained. For obtaining microballoons filled with other gases, the albumin solution was first purged with a new gas, then the foregoing operational sequences were effected under a stream of this gas flowing on the surface of the solution; then at the end of the operations, the suspension was placed in a glass bottle which was extensively purged with the desired gas before sealing.

[0144] The various suspensions of microballoons filled with different gases were diluted to 1:10 with distilled water saturated at equilibrium with air, then they were placed in an optical cell as described above and the absorbance was recorded while increasing steadily the pressure over the suspension. During the measurements, the suspensions temperature was kept at 25° C.

[0145] The results are shown in the Table 1 below and are expressed in terms of the critical pressure PC values registered for a series of gases defined by names or formulae, the characteristic parameters of such gases, i.e., Mw and water solubility being given, as well as the original bubble count and bubble average size (mean diameter in volume).

TABLE 1

Sample	Gas	Mw	Solubility	Bubble Count (10 ⁹ /ml)	Bubble size (μm)	PC (Torr)	S gas/√Mw
AFre1	CF ₄	88	.0038	0.8	5.1	120	.0004
AFre2	CBrF ₃	149	.0045	0.1	11.1	104	.0004
ASF1	SF ₆	146	.005	13.9	6.2	150	.0004
ASF2	SF ₆	146	.005	2.0	7.9	140	.0004
AN1	N ₂	28	.0144	0.4	7.8	62	.0027
A14	Air	29	.0167	3.1	11.9	53	.0031
A18	Air	29	.0167	3.8	9.2	52	—
A19	Air	29	.0167	1.9	9.5	51	—
AMel	CH ₄	16	.032	0.25	8.2	34	.008
AKr1	Kr	84	.059	0.02	9.2	86	.006
AX1	Xe	131	.108	0.06	17.2	65	.009
AX2	Xe	131	.108	0.03	16.5	89	.009

[0146] From the results of Table 1, it is seen that the critical pressure PC increases for gases of lower solubility and higher molecular weight. It can therefore be expected that microvesicles filled with such gases will provide more durable echogenic signals in vivo. It can also be seen that average bubble size generally increases with gas solubility.

EXAMPLE 13

[0147] Aliquots (1 ml) of some of the microballoon suspensions prepared in Example 12 were injected in the Jugular vein of experimental rabbits in order to test echogenicity in vivo. Imaging of the left and right heart ventricles was carried out in the grey scale mode using an Acuson 128-XP5 echography apparatus and a 7.5 MHz transducer. The duration of contrast enhancement in the left ventricle was determined by recording the signal for a period of time. The results are gathered in Table 2 below which also shows the PC of the gases used.

TABLE 2

Sample (Gas)	Duration of contrast (sec)	PC (Torr)
AMe1 (CH ₄)	zero	34
A14 (air)	10	53
A18 (air)	11	52
AX1 (Xe)	20	65
AX2 (Xe)	30	89
ASF2 (SF ₆)	>60	140

[0148] From the above results, one can see the existence of a definite correlation between the critical pressure of the gases tried and the persistence in time of the echogenic signal.

EXAMPLE 14

[0149] A suspension of echogenic air-filled galactose microparticles (Echovistg from Schering AG) was obtained by shaking for 5 sec 3 g of the solid microparticles in 8.5 ml of a 20% galactose solution. In other preparations, the air above a portion of Echovist® particles was evacuated (0.2 Torr) and replaced by an SF₆ atmosphere, whereby, after addition of the 20% galactose solution, a suspension of microparticles containing associated sulfur hexafluoride was obtained. Aliquots (1 ml) of the suspensions were administered to experimental rabbits (by injection in the jugular

vein) and imaging of the heart was effected as described in the previous example. In this case the echogenic microparticles do not transit through the lung capillaries, hence imaging is restricted to the right ventricle and the overall signal persistence has no particular significance. The results of Table 3 below show the value of signal peak intensity a few seconds after injection.

TABLE 3

Sample No	Gas	Signal peak (arbitrary units)
Gal1	air	114
Gal2	air	108
Gal3	SF ₆	131
Gal4	SF ₆	140

[0150] It can be seen that sulfur hexafluoride, an inert gas with low water solubility, provides echogenic suspensions which generate echogenic signals stronger than comparable suspensions filled with air. These results are particularly interesting in view of the teachings of EP-A-0441468 and EP-A-0357163 (Schering) which disclose the use for echography purposes of microparticles, respectively, cavitate and clathrate compounds filled with various gases including SF₆; these documents do not however report particular advantages of SF₆ over other more common gases with regard to the echogenic response.

EXAMPLE 15

[0151] A series of echogenic suspensions of gas-filled microbubbles were prepared by the general method set forth below:

[0152] One gram of a mixture of hydrogenated soya lecithin (from Nattermann Phospholipids GmbH, Germany) and dicetyl-phosphate (DCP), in 9/1 molar ratio, was dissolved in 50 ml of chloroform, and the solution was placed in a 100 ml round flask and evaporated to dryness on a Rotavapor apparatus. Then, 20 ml of distilled water were added and the mixture was slowly agitated at 75° C. for an hour. This resulted in the formation of a suspension of multilamellar liposomes (MLV) which was thereafter extruded at 75° C. through, successively, 3 μm and 0.8 μm polycarbonate membranes (Nuclepore(D)). After cooling, 1 ml aliquots of the extruded suspension were diluted with 9 ml of a concentrated lactose solution (83 g/l), and the diluted suspensions were frozen at -45° C. The frozen samples were thereafter freeze-dried under high vacuum to a free-flowing powder in a vessel which was ultimately filled with air or a gas taken from a selection of gases as indicated in Table 4 below. The powdery samples were then resuspended in 10 ml of water as the carrier liquid, this being effected under a stream of the same gas used to fill the said vessels. Suspension was effected by vigorously shaking for 1 min on a vortex mixer.

[0153] The various suspensions were diluted 1:20 with distilled water equilibrated beforehand with air at 25° C. and the dilutions were then pressure tested at 25° C. as disclosed in Example 12 by measuring the optical density in a spectrophotometric cell which was subjected to a progressively increasing hydrostatic pressure until all bubbles had collapsed. The results are collected in Table 4 below which, in

addition to the critical pressure PC, gives also the AP values (see FIG. 1).

TABLE 4

Sample No	Gas	Mw	Solubility in H ₂ O	Bubble Count (10 ⁸ /ml)	PC (Torr)	Δ P (Torr)
LFre1	CF ₄	88	.0038	1.2	97	35
LFre2	CBrF ₃	149	.0045	0.9	116	64
LSF1	SF ₆	146	.005	1.2	92	58
LFre3	C ₄ F ₈	200	.016	1.5	136	145
L1	air	29	.0167	15.5	68	17
L2	air	29	.0167	11.2	63	17
LAr1	Ar	40	.031	14.5	71	18
LKr1	Kr	84	.059	12.2	86	18
LXe1	Xe	131	.108	10.1	92	23
LFre4	CHClF ₂	86	.78	—	83	25

[0154] The foregoing results clearly indicate that the highest resistance to pressure increases is provided by the most water-insoluble gases. The behavior of the microbubbles is therefore similar to that of the microballoons in this regard. Also, the less water-soluble gases with the higher molecular weights provide the flattest bubble-collapse/pressure curves (i.e., ΔP is the widest) which is also an important factor of echogenic response durability in vivo, as indicated hereinbefore.

EXAMPLE 16

[0155] Some of the microbubble suspensions of Example 15 were injected to the jugular vein of experimental rabbits as indicated in Example 13 and imaging of the left heart ventricle was effected as indicated previously. The duration of the period for which a useful echogenic signal was detected was recorded and the results are shown in Table 5 below in which C₄F₈ indicates octafluorocyclobutane.

TABLE 5

Sample No	Type of gas	Contrast duration (sec)
L1	Air	38
L2	Air	29
LMe1	CH ₄	47
LKr1	Krypton	37
LFre1	CF ₄	>120
LFre2	CBrF ₃	92
LSF1	SF ₆	>112
LFre3	C ₄ F ₈	>120

[0156] These results indicate that, again in the case of microbubbles, the gases according to the criteria of the present invention will provide ultrasonic echo signal for a much longer period than most gases used until now.

EXAMPLE 17

[0157] Suspensions of microbubbles were prepared using different gases exactly as described in Example 15, but replacing the lecithin phospholipid ingredient by a mole equivalent of diarachidoylphosphatidylcholine (C₂₀ fatty acid residue) available from Avanti Polar Lipids, Birmingham, Ala. USA. The phospholipid to DCP molar ratio was

still 9/1. Then the suspensions were pressure tested as in Example 15; the results, collected in Table 6A below, are to be compared with those of Table 4.

TABLE 6A

Sample No	Type of Gas	Mw of Gas	Solubility in water	Bubble Count (10 ⁸ /ml)	PC (Torr)	Δ P (Torr)
LFre1	CF ₄	88	.0038	3.4	251	124
LFre2	CBrF ₃	149	.0045	0.7	121	74
LSF1	SF ₆	146	.005	3.1	347	>150
LFre3	C ₄ F ₈	200	.016	1.7	>350	>200
L1	Air	29	.0167	3.8	60	22
LBu1	Butane	58	.027	0.4	64	26
LAr1	Argon	40	.031	3.3	84	47
LMe1	CH ₄	16	.032	3.0	51	19
LEt1	C ₂ H ₆	44	.034	1.4	61	26
LKr1	Kr	84	.059	2.7	63	18
LXe1	Xe	131	.108	1.4	60	28
LFre4	CHClF ₂	86	.78	0.4	58	28

[0158] The above results, compared to that of Table 4, show that, at least with low solubility gases, by lengthening the chain of the phospholipid fatty acid residues, one can dramatically increase the stability of the echogenic suspension toward pressure increases. This was further confirmed by repeating the foregoing experiments but replacing the phospholipid component by its higher homolog, i.e., di-behenoylphosphatidylcholine (C₂₂ fatty acid residue). In this case, the resistance to collapse with pressure of the microbubbles suspensions was still further increased.

[0159] Some of the microbubbles suspensions of this Example were tested in dogs as described previously for rabbits (imaging of the heart ventricles after injection of 5 ml samples in the anterior cephalic vein). A significant enhancement of the useful in vivo echogenic response was noted, in comparison with the behavior of the preparations disclosed in Example 15, i.e., the increase in chain length of the fatty-acid residue in the phospholipid component increases the useful life of the echogenic agent in vivo.

[0160] In the next Table below, there is shown the relative stability in the left ventricle of the rabbit of microbubbles (SF₆) prepared from suspensions of a series of phospholipids whose fatty acid residues have different chain lengths (<injected dose: 1 ml/rabbit).

TABLE 6B

Phospholipid	Chain length (C _n)	PC (Torr)	Δ P (Torr)	Duration of contrast (sec)
DMPC	14	57	37	31
DPPC	16	100	76	105
DSPC	18	115	95	120
DAPC	20	266	190	>300

[0161] It has been mentioned hereinabove that for the measurement of resistance to pressure described in these Examples, a constant rate of pressure rise of 100 Torr/min was maintained. This is justified by the results given below which show the variations of the PC values for different gases in function to the rate of pressure increase. In these samples DMPC was the phospholipid used.

Gas	PC (Torr)		
	Rate of pressure increase (Torr/min)		
Sample	40	100	200
SF ₆	51	57	82
Air	39	50	62
CH ₄	47	61	69
Xe	38	43	51
Freon 22	37	54	67

EXAMPLE 18

[0162] A series of albumin microballoons as suspensions in water were prepared under air in a controlled sphere size fashion using the directions given in Example 12. Then the air in some of the samples was replaced by other gases by the gas-exchange sweep method at ambient pressure. Then, after diluting to 1:10 with distilled water as usual, the samples were subjected to pressure testing as in Example 12. From the results gathered in Table 7 below, it can be seen that the two-steps preparation mode gives, in some cases, echo-generating agents with better resistance to pressure than the one-step preparation mode of Example 12.

TABLE 7

Sample No	Type of gas	Mw of the gas	Solubility in water	Initial Bubble Count (10 ⁸ /ml)	PC (Torr)
A14	Air	29	.0167	3.1	53
A18	Air	29	.0167	3.8	52
A18/SF ₆	SF ₆	146	.005	0.8	115
A18/C ₂ H ₆	C ₂ H ₆	30	.042	3.4	72
A19	Air	29	.0167	1.9	51
A19/SF ₆	SF ₆	146	.005	0.6	140
A19/Xe	Xe	131	.108	1.3	67
A22/CF ₄	CF ₄	88	.0038	1.0	167
A22/Kr	Kr	84	.059	0.6	85

EXAMPLE 19

[0163] The method of the present invention was applied to an experiment as disclosed in the prior art, for instance Example 1 WO-92/11873. Three grams of Pluronic® F68 (a copolymer of polyoxyethylene-polyoxypropylene with a molecular weight of 8400), 1 g of dipalmitoylphosphatidylglycerol (Na salt, Avanti Polar Lipids) and 3.6 g of glycerol were added to 80 ml of distilled water. After heating at about 80° C., a clear homogenous solution was obtained. The tenside solution was cooled to room temperature and the volume was adjusted to 100 ml. In some experiments (see Table 8) dipalmitoylphosphatidylglycerol was replaced by a mixture of diarachidoylphosphatidylcholine (920 mg) and 80 mg of dipalmitoylphosphatidic acid (Na salt, Avanti Polar lipids).

[0164] The bubble suspensions were obtained by using two syringes connected via a three-way valve. One of the syringes was filled with 5 ml of the tenside solution while the other was filled with 0.5 ml of air or gas. The three-way valve was filled with the tenside solution before it was connected to the gas-containing syringe. By alternatively operating the two pistons, the tenside solutions were transferred back and forth between the two syringes (5 times in

each direction), milky suspensions were formed. After dilution (1:10 to 1:50) with distilled water saturated at equilibrium with air, the resistance to pressure of the preparations was determined according to Example 12, the pressure increase rate was 240 Torr/min. The following results were obtained:

TABLE 8

Phospholipid	Gas	Pc (mm Hg)	DP (mm Hg)
DPPG	air	28	17
DPPG	SF ₆	138	134
DAPC/DPPA 9/1	air	46	30
DAPC/DPPA 9/1	SF ₆	269	253

[0165] It follows that by using the method of the invention and replacing air with other gases, e.g., SF₆, even with known preparations a considerable improvements, i.e., increase in the resistance to pressure, may be achieved. This is true both in the case of negatively charged phospholipids (e.g., DPPG) and in the case of mixtures of neutral and negatively charged phospholipids (DAPC/DPPA).

[0166] The above experiment further demonstrates that the recognized problem sensitivity of microbubbles and microballoons to collapse when exposed to pressure, i.e., when suspensions are injected into living beings, has advantageously been solved by the method of the invention. Suspensions with microbubbles or microballoons with greater resistance against collapse and greater stability can advantageously be produced providing suspensions with better reproducibility and improved safety of echographic measurements performed in vivo on a human or animal body.

Stable Microballoon Compositions of the Invention

[0167] Desirable features have also now been achieved with the microballoons of the present invention which are of micronic or submicronic size bounded by a polymer membrane filled with air or a gas suitable, when in the form of suspensions in a liquid carrier, to be administered to human or animal patients for therapeutic or diagnostic applications, e.g., for the purpose of ultrasonic echography imaging. The polymer of the membrane is a deformable and resilient interfacially deposited polymer. The invention also includes air or gas filled microballoons bounded by an elastic interfacial polymeric membrane adapted to form with suitable physiologically acceptable aqueous carrier liquids suspensions to be taken orally, rectally and urethrally, or injectable into living organisms for therapeutic or diagnostic purposes. These microballoons are characterized as being non-coalescent, dry and instantly dispersible by admixing with a liquid carrier. Moreover, although the present microspheres can generally be made relatively short-lived, i.e., susceptible to biodegradation to cope with the foregoing metabolism problems by using selected types of polymers, this feature (which is actually controlled by the fabrication parameters) is not a commercial drawback because either the microballoons can be stored and shipped dry, a condition in which they are stable indefinitely, or the membrane can be made substantially impervious to the carrier liquid, degradation starting to occur only after injection. In the first case, the microballoons supplied in dry powder form are simply admixed with a proportion of an aqueous phase carrier

before use, this proportion being selected depending on the needs. Note that this is an additional advantage over the prior art products because the concentration can be chosen at will and initial values far exceeding the aforementioned $10^8/\text{ml}$, i.e., in the range 10^5 to 10^{10} , are readily accessible. It should be noted that the method of the invention (to be disclosed hereafter) enables to control porosity to a wide extent; hence microballoons with a substantially impervious membrane can be made easily which are stable in the form of suspensions in aqueous liquids and which can be marketed as such also.

[0168] Microspheres with membranes of interfacially deposited polymers, although in the state where they are filled with liquid, are well known in the art. They may normally result from the emulsification into droplets (the size of which is controllable in function to the emulsification parameters) of a first aqueous phase in an organic solution of polymer followed by dispersion of this emulsion into a second water phase and subsequent evaporation of the organic solvent. During evaporation of the volatile solvent, the polymer deposits interfacially at the droplets boundary and forms a microporous membrane which efficiently bounds the encapsulated first aqueous phase from the surrounding second aqueous phase. This technique, although possible, is not preferred in the present invention.

[0169] Alternatively, one may emulsify with an emulsifier a hydrophobic phase in an aqueous phase (usually containing viscosity increasing agents as emulsion stabilizers) thus obtaining an oil-in-water type emulsion of droplets of the hydrophobic phase and thereafter adding thereto a membrane forming polymer dissolved in a volatile organic solvent not miscible with the aqueous phase.

[0170] If the polymer is insoluble in the hydrophobic phase, it will deposit interfacially at the boundary between the droplets and the aqueous phase. Otherwise, evaporation of the volatile solvent will lead to the formation of said interfacially deposited membrane around the droplets of the emulsified hydrophobic phase. Subsequent evaporation of the encapsulated volatile hydrophobic phase provides water filled microspheres surrounded by interfacially deposited polymer membranes. This technique which is advantageously used in the present invention is disclosed by K. Uno et al. in *J. Microencapsulation* 1 (1984), 3-8 and K. Makino et al., *Chem. Pharm. Bull.* 33 (1984), 1195-1201. As said before, the size of the droplets can be controlled by changing the emulsification parameters, i.e., nature of emulsifier (more effective the surfactant, i.e., the larger the hydrophilic to lipophilic balance, the smaller the droplets) and the stirring conditions (faster and more energetic the agitation, the smaller the droplets).

[0171] In another variant, the interfacial wall forming polymer is dissolved in the starting hydrophobic phase itself; the latter is emulsified into droplets in the aqueous phase and the membrane around the droplets will form upon subsequent evaporation of this encapsulated hydrophobic phase. An example of this is reported by J. R. Famand et al., *Powder Technology* 22 (1978), 11-16 who emulsify a solution of polymer (e.d., polyethylene) in naphthalene in boiling water, then after cooling they recover the naphthalene in the form of a suspension of polymer bounded microbeads in cold water and, finally, they remove the naphthalene by subjecting the microbeads to sublimation, whereby 25

micron microballoons are produced. Other examples exist, in which a polymer is dissolved in a mixed hydrophobic phase comprising a volatile hydrophobic organic solvent and a water-soluble organic solvent, then this polymer solution is emulsified in a water phase containing an emulsifier, whereby the water-soluble solvent disperses into the water phase, thus aiding in the formation of the emulsion of microdroplets of the hydrophobic phase and causing the polymer to precipitate at the interface; this is disclosed in EP-A-274,961 (H. Fessi).

[0172] The aforementioned techniques can be adapted to the preparation of air or gas filled microballoons suited for ultrasonic imaging provided that appropriate conditions are found to control sphere size in the desired ranges, cell-wall permeability or imperviousness and replacement of the encapsulated liquid phase by air or a selected gas. Control of overall sphere size is obviously important to adapt the microballoons to use purposes, i.e., injection or oral intake. The size conditions for injection (about 0.5-10 μm average size) have been discussed previously. For oral application, the range can be much wider, being considered that echogenicity increases with size; hence microballoons in several size ranges between say 1 and 1000 microns can be used depending on the needs and provided the membrane is elastic enough not to break during transit in the stomach and intestine. Control of cell-wall permeability is important to ensure that infiltration by the injectable aqueous carrier phase is absent or slow enough not to impair the echographic measurements but, in cases, still substantial to ensure relatively fast after-test biodegradability, i.e., ready metabolization of the suspension by the organism. Also the microporous structure of the microballoons envelope (pores of a few nm to a few hundreds of nm or more for microballoons envelopes of thickness ranging from 50-500 nm) is a factor of resiliency, i.e., the microspheres can readily accept pressure variations without breaking. The preferred range of pore sizes is about 50-2000 nm.

[0173] The conditions for achieving these results are met by using the method including the steps of (1) emulsifying a hydrophobic organic phase into a water phase so as to obtain droplets of the hydrophobic phase as an oil-in-water emulsion in the water phase; (2) adding to said emulsion a solution of at least one polymer in a volatile solvent insoluble in the water phase, so that a layer of said polymer will form around said droplets; then (3) evaporating the volatile solvent so that the polymer will deposit by interfacial precipitation around the droplets which then form beads with a core of the hydrophobic phase encapsulated by a membrane of the polymer, the beads being in suspension in the water phase; and finally (4) subjecting the suspension to reduced pressure under conditions such that the encapsulated hydrophobic phase can be removed by evaporation.

[0174] The hydrophobic phase selected in step (4) so it evaporates substantially simultaneously with the water phase and is replaced by air or gas, whereby dry, free flowing, readily dispersible microballoons are obtained.

[0175] One factor which enables to control the permeability of the microballoons membrane is the rate of evaporation of the hydrophobic phase relative to that of water in step (4) of the method, e.g., under conditions of freeze drying. For instance if the evaporation is carried out between about -40° and 0°C , and hexane is used as the hydrophobic phase,

polystyrene being the interfacially deposited polymer, beads with relatively large pores are obtained; this is so because the vapour pressure of the hydrocarbon in the chosen temperature range is significantly greater than that of water, which means that the pressure difference between the inside and outside of the spheres will tend to increase the size of the pores in the spheres membrane through which the inside material will be evaporated. In contrast, using cyclooctane as the hydrophobic phase (at -17°C the vapour pressure is the same as that of water) will provide beads with very tiny pores because the difference of pressures between the inside and outside of the spheres during evaporation is minimized.

[0176] Depending on degree of porosity the microballoons of this invention can be made stable in an aqueous carrier from several hours to several months and give reproducible echographic signals for a long period of time. Actually, depending on the polymer selected, the membrane of the microballoons can be made substantially impervious when suspended in carrier liquids of appropriate osmotic properties, i.e., containing solutes in appropriate concentrations. It should be noted that the existence of micropores in the envelope of the microballoons of the present invention appears to be also related with the echographic response, i.e., all other factors being constant, microporous vesicles provide more efficient echographic signal than corresponding non-porous vesicles. The reason is not known but it can be postulated that when a gas is in resonance in a closed structure, the damping properties of the latter may be different if it is porous or non-porous.

[0177] Other non water soluble organic solvents which have a vapour pressure of the same order of magnitude between about -40°C and 0°C are convenient as hydrophobic solvents in this invention. These include hydrocarbons such as for instance n-octane, cyclooctane, the dimethylcyclohexanes, ethyl-cyclohexane, 2-, 3- and 4-methylheptane, 3-ethyl-hexane, toluene, xylene, 2-methyl-2-heptane, 2,2,3,3-tetramethylbutane and the like. Esters such as propyl and isopropyl butyrate and isobutyrate, butylformate and the like, are also convenient in this range. Another advantage of freeze drying is to operate under reduced pressure of a gas instead of air, whereby gas filled microballoons will result. Physiologically acceptable gases such as CO_2 , N_2O , methane, Freon (organic compounds containing one or more carbon atoms and fluorine such as CF_4 , C_2F_6 , C_3F_8 or C_4F_8 and the like), helium and other rare gases are possible. Gases with radioactive tracer activity can be contemplated.

[0178] As the volatile solvent insoluble in water to be used for dissolving the polymer to be precipitated interfacially, one can cite halo-compounds such as CCl_4 , CH_3Br , CH_2Cl_2 , chloroform, Freon (organic compounds containing one or more carbon atoms and fluorine such as CF_4 , C_2F_6 , C_3F_8 or C_4F_8 and the like), low boiling esters such as methyl, ethyl and propyl acetate as well as lower ethers and ketones of low water solubility. When solvents not totally insoluble in water are used, e.g., diethyl-ether, it is advantageous to use, as the aqueous phase, a water solution saturated with said solvent beforehand.

[0179] The aqueous phase in which the hydrophobic phase is emulsified as an oil-in-water emulsion preferably contains 1-20% by weight of water-soluble hydrophilic compounds like sugars and polymers as stabilizers, e.g., polyvinyl

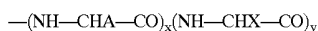
alcohol (PVA), polyvinyl pyrrolidone (PVP), polyethylene glycol (PEG), gelatin, polyglutamic acid, albumin, and polysaccharides such as starch, dextran, agar, xanthan and the like. Similar aqueous phases can be used as the carrier liquid in which the microballoons are suspended before use.

[0180] Part of this water-soluble polymer can remain in the envelope of the microballoons or it can be removed by washing the beads before subjecting them to final evaporation of the encapsulated hydrophobic core phase.

[0181] The emulsifiers to be used (0.1-5% by weight) to provide the oil-in-water emulsion of the hydrophobic phase in the aqueous phase include most physiologically acceptable emulsifiers, for instance egg lecithin or soya bean lecithin, or synthetic lecithins such as saturated synthetic lecithins, for example, dimyristoyl phosphatidyl choline, dipalmitoyl phosphatidyl choline or distearoyl phosphatidyl choline or unsaturated synthetic lecithins, such as dioleoyl phosphatidyl choline or dilinoleoyl phosphatidyl choline. Emulsifiers also include surfactants such as free fatty acids, esters of fatty acids with polyoxyalkylene compounds like polyoxypropylene glycol and polyoxyethylene glycol; ethers of fatty alcohols with polyoxyalkylene glycols; esters of fatty acids with polyoxyalkylated sorbitan; soaps: glycerol-polyalkylene stearate; glycerol-polyoxyethylene ricinoleate; homo- and copolymers of polyalkylene glycols; polyethoxylated soya-oil and castor oil as well as hydrogenated derivatives; ethers and esters of sucrose or other carbohydrates with fatty acids, fatty alcohols, these being optionally polyoxyalkylated; mono-, di- and triglycerides of saturated or unsaturated fatty acids; glycerides or soya-oil and sucrose.

[0182] The polymer which constitutes the envelope or bounding membrane of the injectable microballoons can be selected from most hydrophilic, biodegradable physiologically compatible polymers. Among such polymers one can cite polysaccharides of low water solubility, polyactides and polyglycolides and their copolymers, copolymers of lactides and lactones such as δ -caprolactone, 5-valerolactone, polypeptides, and proteins such as gelatin, collagen, globulins and albumins. The great versatility in the selection of synthetic polymers is another advantage of the present invention since, as with allergic patients, one may wish to avoid using microballoons made of natural proteins (albumin, gelatin) like in U.S. Pat. No. 4,276,885 or EP-A-324,938. Other suitable polymers include poly-(ortho)esters (see for instance U.S. Pat. No. 4,093,709; U.S. Pat. No. 4,131,648; U.S. Pat. No. 4,138,344; U.S. Pat. No. 4,180,646); polylactic and polyglycolic acid and their copolymers, for instance DEXON (see J. Heller, *Biomaterials* 1 (1980), 51; poly(DL-lactide-co- δ -caprolactone), poly(DL-lactide-co- δ -valerolactone), poly(DL-lactide-co- δ -butyrolactone), polyalkylcyanoacrylates; polyamides, polyhydroxybutyrate; polydioxanone; poly- β -aminoketones (Polymer 23 (1982), 1693); polyphosphazenes (Science 193 (1976), 1214); and polyanhydrides. References on biodegradable polymers can be found in R. Langer et al., *Macromol. Chem. Phys.* C23 (1983), 61-126. Polyamino-acids such as polyglutamic and polyaspartic acids can also be used as well as their derivatives, i.e., partial esters with lower alcohols or glycols. One useful example of such polymers is poly-(t-butyl-glutamate). Copolymers with other amino-acids such as methionine, leucine, valine, proline, glycine, alanine, etc. are also possible. Recently some novel derivatives of polyglutamic and

polyaspartic acid with controlled biodegradability have been reported (see W087/03891; U.S. Pat. No. 4,888,398 and EP-130.935 incorporated here by reference). These polymers (and copolymers with other amino-acids) have formulae of the following type:

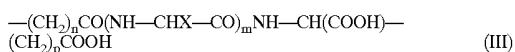


[0183] where X designates the side chain of an amino-acid residue and A is a group of formula $-(\text{CH}_2)_n\text{COOR}^1\text{R}^2-\text{OCOR}$ (II), with R^1 and R^2 being H or lower alkyls, and R being alkyl or aryl; or R and R^1 are connected together by a substituted or unsubstituted linking member to provide 5- or 6-membered rings.

[0184] A can also represent groups of formulae:



[0185] and



[0186] and corresponding anhydrides. In all these formulae n, m and p are lower integers (not exceeding 5) and x and y are also integers selected for having molecular weights not below 5000.

[0187] The aforementioned polymers are suitable for making the microballoons according to the invention and, depending on the nature of substituents R, R^1 , R^2 and X, the properties of the membrane can be controlled, for instance, strength, elasticity and biodegradability. For instance X can be methyl (alanine), isopropyl (valine), isobutyl (leucine and isoleucine), benzyl (phenylalanine).

[0188] Additives can be incorporated into the polymer wall of the microballoons to modify the physical properties such as dispersibility, elasticity and water permeability. For incorporation in the polymer, the additives can be dissolved in the polymer carrying phase, e.g., the hydrophobic phase to be emulsified in the water phase, whereby they will co-precipitate with the polymer during inter-facial membrane formation.

[0189] Among the useful additives, one may cite compounds which can "hydrophobize" the microballoons membrane in order to decrease water permeability, such as fats, waxes and high molecular-weight hydrocarbons. Additives which improve dispersibility of the microballoons in the injectable liquid-carrier are amphipatic compounds like the phospholipids; they also increase water permeability and rate of biodegradability.

[0190] Non-biodegradable polymers for making microballoons to be used in the digestive tract can be selected from most water-insoluble, physiologically acceptable, bioresistant polymers including polyolefins (polystyrene), acrylic resins (polyacrylates, polyacrylonitrile), polyesters (polycarbonate), polyurethanes, polyurea and their copolymers. ABS (acryl-butadienestyrene) is a preferred copolymer.

[0191] Additives which increase membrane elasticity are the plasticisers like isopropyl myristate and the like. Also, very useful additives are constituted by polymers akin to that of the membrane itself but with relatively low molecular weight. For instance when using copolymers of poly(lactic/polyglycolic) type as the membrane forming material, the properties of the membrane can be modified advantageously (enhanced softness and biodegradability) by incorporating,

as additives, low molecular weight (1,000 to 15,000 Dalton) polyglycolides or polylactides. Also polyethylene glycol of moderate to low M_w (e.g., PEG 2000) is a useful softening additive.

[0192] Preferably the plasticizers include isopropyl myristate, glyceryl monostearate and the like to control flexibility, the amphipatic substances include surfactants and phospholipids like the lecithins to control permeability by increasing porosity while the hydrophobic compounds include high molecular weight hydrocarbon like the paraffin-waxes to reduce porosity.

[0193] The quantity of additives to be incorporated in the polymer forming the inter-facially deposited membrane of the present microballoons is extremely variable and depends on the needs. In some cases no additive is used at all; in other cases amounts of additives which may reach about 20% by weight of the polymer are possible.

[0194] The injectable microballoons of the present invention can be stored dry in the presence or in the absence of additives to improve conservation and prevent coalescence. As additives, one may select from 0.1 to 25% by weight of water-soluble physiologically acceptable compounds such as mannitol, galactose, lactose or sucrose or hydrophilic polymers like dextran, xanthan, agar, starch, PVP, polyglutamic acid, polyvinylalcohol (PVA), albumin and gelatin. The useful life-time of the microballoons in the injectable liquid carrier phase, i.e., the period during which useful echographic signals are observed, can be controlled to last from a few minutes to several months depending on the needs; this can be done by controlling the porosity of the membrane from substantial imperviousness toward carrier liquids to porosities having pores of a few nanometers to several hundreds of nanometers. This degree of porosity can be controlled, in addition to properly selecting the membrane forming polymer and polymer additives, by adjusting the evaporation rate and temperature in step (4) of the method and properly selecting the nature of the compound (or mixture of compounds) constituting the hydrophobic phase, i.e., the greater the differences in its partial pressure of evaporation with that of the water phase, the coarser the pores in the microballoons membrane will be. Of course, this control by selection of the hydrophobic phase can be further refined by the choice of stabilizers and by adjusting the concentration thereof in order to control the rate of water evaporation during the forming of the microballoons. All these changes can easily be made by skilled persons without exercising inventiveness and need not be further discussed.

[0195] It should be remarked that although the microballoons of this invention can be marketed in the dry state, more particularly when they are designed with a limited life time after injection, it may be desirable to also sell ready preparations, i.e., suspensions of microballoons in an aqueous liquid carrier ready for injection or oral administration. This requires that the membrane of the microballoons be substantially impervious (at least for several months or more) to the carrier liquid. It has been shown in this description that such conditions can be easily achieved with the present method by properly selecting the nature of the polymer and the interfacial deposition parameters. Actually parameters have been found (for instance using the polyglutamic polymer (where A is the group of formula II) and cyclooctane as the hydrophobic phase) such that the porosity of the mem-

brane after evaporation of the hydrophobic phase is so tenuous that the microballoons are substantially impervious to the aqueous carrier liquid in which they are suspended.

[0196] A preferred administrable preparation for diagnostic purposes comprises a suspension in buffered or unbuffered saline (0.9% aqueous NaCl; buffer 10 mM tris-HCl) containing 10^8 - 10^{10} vesicles/ml. This can be prepared mainly according to the directions of the Examples below, preferably Examples 22 and 23, using poly-(DL-lactide) polymers from the Company Boehringer, Ingelheim, Germany.

[0197] The invention described up until this point can be further elucidated by the description of the following representative (but not limiting) embodiments, numbered 1-31:

[0198] 1. Microcapsules or microballoons of micronic or submicronic size bounded by a polymer membrane filled with air or a gas suitable, when in the form of suspensions in a liquid carrier, to be administered to human or animal patients for therapeutic or diagnostic applications, e.g., for the purpose of ultrasonic echography imaging, characterized in that the polymer of the membrane is a deformable and resilient interfacially deposited polymer.

[0199] 2. Air or gas filled microballoons bounded by an elastic interfacial polymeric membrane * adapted to form with suitable physiologically acceptable aqueous carrier liquids suspensions to be taken orally, rectally and urethrally, or injectable into living organisms for therapeutic or diagnostic purposes, characterized in being non-coalescent dry and instantly dispersible by admixing with said liquid carrier.

[0200] 3. The microballoons of embodiments 1 or 2 having size mostly in the 0.5-10 μ m range suitable for injection into the bloodstream of living beings, characterized in that the membrane polymer is biodegradable and the membrane is either impervious or contains pores permeable to bioactive liquids for increasing the rate of biodegradation.

[0201] 4. The microballoons of embodiment 3, in which the polymer membrane has a porosity ranging from a few nanometers to several hundreds or thousands of nanometers, preferable 50-2000 nm.

[0202] 5. The microballoons of embodiment 3, in which the membrane is elastic, has a thickness of 50-500 nm, and resists pressure variations produced by heart beat pulsations in the bloodstream.

[0203] 6. The microballoons of embodiment 3, in which the polymer of the membrane is a biodegradable polymer selected from polysaccharides, polyamino-acids, polylactides and polyglycolides and their copolymers, copolymers of lactides and lactones, polypeptides, poly-(ortho)esters, polydioxanone, poly- β -aminoketones, polyphosphazenes, polyanhydrides and poly(alkyl-cyano-acrylates).

[0204] 7. The microballoons of embodiment 3, in which the membrane polymer is selected from polyglutamic or polyaspartic acid derivatives and their copolymers with other amino-acids.

[0205] 8. The microballoons of embodiment 7, in which the polyglutamic and polyaspartic acid derivatives are

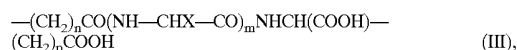
selected from esters and amides involving the carboxylated side function thereof, said side functions having formulae



[0206] or



[0207] or



[0208] in which R is an alkyl or aryl substituent; R^1 and R^2 are H or lower alkyls, or R and R^1 are connected together by a substituted or unsubstituted linking member to form a 5- or 6-membered ring; n is 1 or 2; p is 1, 2 or 3; m is an integer from 1 to 5 and X is a side chain of an aminoacid residue.

[0209] 9. The microballoons of embodiment 3, in which the membrane polymer contains additives to control the degree of elasticity, and the size and density of the pores for permeability control.

[0210] 10. The microballoons of embodiment 9, in which said additives include plasticizers, amphipatic substances and hydrophobic compounds.

[0211] 11. The microballoons of embodiment 10, in which the plasticizers include isopropyl myristate, glyceryl monostearate and the like to control flexibility, the amphipatic substances include surfactants and phospholipids like the lecithins to control permeability by increasing porosity and the hydrophobic compounds include high molecular weight hydrocarbon like the paraffin-waxes to reduce porosity.

[0212] 12. The microballoons of embodiment 10, in which the additives include polymers of low molecular weight, e.g., in the range of 1,000 to 15,000, to control softness and resiliency of the microballoon membrane.

[0213] 13. The microballoons of embodiment 12, in which the low molecular weight polymer additives are selected from polylactides, polyglycolides, polyalkylene glycols like polyethylene glycol and polypropylene glycol, and polyols like polyglycerol.

[0214] 14. The microballoons of embodiments 1 or 2, having size up to about 1000 μ m suitable for oral, rectal and urethral applications, characterized in that the membrane polymer is not biodegradable in the digestive tract and impervious to biological liquids.

[0215] 15. The microballoons of embodiment 14, in which the polymer is selected from polyolefins, polyacrylates, polyacrylonitrile, non-hydrolyzable polymesters, polyurethanes and polyureas.

[0216] 16. Aqueous suspension of the microballoons according to embodiments 1 or 2 for administration to patients, characterized in containing a concentration of about 10^6 to 10^{10} microballoons/ml, this being stable for period exceeding a month.

[0217] 17. A method for making air or gas filled microballoons usable as suspensions in a carrier liquid for oral, rectal and urethral applications, or for injections into living organisms, this method comprising the steps of:

[0218] (a) emulsifying a hydrophobic organic phase into a water phase so as to obtain droplets of said hydrophobic phase as an oil-in-water emulsion in said water phase;

- [0219] (b) adding to said emulsion a solution of at least one polymer in a volatile solvent insoluble in the water phase, so that a later of said polymer will form around said droplets;
- [0220] (c) evaporating said volatile solvent so that the polymer will deposit by interfacial precipitation around the droplets which then form beads with a core of said hydrophobic phase encapsulated by a membrane of said polymer, said beads being in suspension in said water phase;
- [0221] (d) subjecting said suspension to reduced pressure under conditions such that said encapsulated hydrophobic phase be removed by evaporation;
- [0222] characterized in that said hydrophobic phase is selected so that in step (4) it evaporates substantially simultaneously with the water phase and is replaced by air or gas, whereby dry, free flowing, readily dispersible microballoons are obtained.
- [0223] 18. The method of embodiment 17, in which said polymer is dissolved in said hydrophobic phase, so that steps (2) and (3) can be omitted and the polymer membrane will form by interfacial precipitation during step (4).
- [0224] 19. The method of embodiment 17, characterized in that evaporation of said hydrophobic phase in step (4) is performed at a temperature where the partial vapour pressure of said hydrophobic phase is of the same order as that of water vapour.
- [0225] 20. The method of embodiment 17, in which said evaporation of step (4) is carried out under freeze-drying conditions.
- [0226] 21. The method of embodiment 20, in which freeze-drying is effected at temperatures of from -40°C . to 0°C .
- [0227] 22. The method of embodiments 17 or 19, in which the hydrophobic phase is selected from organic compounds having a vapour pressure of about 1 Torr at a temperature comprised in the interval of about -40°C . to 0°C .
- [0228] 23. The method of embodiments 17 or 18, in which the aqueous phase comprises, dissolved, from about 1 to 20% by weight of stabilizers comprising hydrophilic compound selected from sugars, PVA, PVP, gelatin, starch, dextran, polydextrose, albumin and the like.
- [0229] 24. The method of embodiment 18, in which additives to control the degree of permeability of the microballoons membrane are added to the hydrophobic phase, the rate of biodegradability of the polymer after injecting the microballoons into living organisms being a function of said degree of permeability.
- [0230] 25. The method of embodiment 24, in which the said additives include hydrophobic solids like fats, waxes and high molecular weight hydrocarbons, the presence of which in the membrane polymer of the microballoons will reduce permeability toward aqueous liquids.
- [0231] 26. The method of embodiment 24, in which the said additives include amphipatic compounds like the phospholids, or low molecular weight polymers, the presence of which in the membrane polymer will increase permeability of the microballoons to aqueous liquids.
- [0232] 27. The method of embodiment 18, in which the hydrophobic phase subjected to emulsification in said water phase also contains a water-soluble solvent which, upon being diluted into said water phase during emulsification, will aid in reducing the size of droplets and induce interfacial precipitation of the polymer before step (4) is carried out.
- [0233] 28. A method for making air or gas filled microballoons usable as suspensions in a carrier liquid for oral, rectal and urethral applications, or for injections into living organisms, this method comprising the steps of;
- [0234] (a) emulsifying a hydrophobic organic phase into a water phase so as to obtain droplets of said hydrophobic organic phase as an oil-in-water emulsion in said water phase, said organic phase containing, dissolved therein, one or more water-insoluble polymers;
- [0235] (b) subjecting said emulsion to reduced pressure under conditions such that said hydrophobic phase be removed by evaporation, whereby the polymer dissolved in the droplets will deposit interfacially and form a polymer bounding membrane, the droplets being simultaneously converted to microballoons, characterized in that said hydrophobic phase is selected so that in step (2) it evaporates substantially simultaneously with the water phase and, upon evaporation, is replaced by air or gas, whereby the microballoons obtained are in dry, free flowing and readily dispersible form.
- [0236] 29. The method of embodiment 28, in which the hydrophobic polymer solution phase subjected to emulsification in said water phase also contains a water-soluble solvent which, upon being diluted into said water phase during emulsification, will aid in reducing the size of droplets and induce interfacial precipitations of the polymer before step (2) is carried out.
- [0237] 30. The method of embodiment 28, in which said organic hydrophobic phase emulsified in step (1) contains no polymer dissolved therein, and before carrying through step (2), the following additional steps are performed:
- [0238] (a) adding to said emulsion a solution of at least one polymer in a volatile solvent insoluble in the water phase, so that a layer of said polymer will form around said droplets;
- [0239] (b) evaporating said volatile solvent so that the polymer will deposit by interfacial precipitation around the droplets, thus forming microballoons or beads with a core of said hydrophobic phase encapsulated by a membrane of said polymer, said beads being in suspension in said water phase, whereby in step (2) evaporation of said hydrophobic phase takes place through said membrane and provides it with substantial microporosity.
- [0240] 31. An injectable aqueous suspension of microballoons containing 10^8 - 10^{10} vesicles/ml bounded by a membrane of interfacially precipitated DL-lactide polymer defined by the commercial name of Resomer.
- [0241] The following Examples illustrate the invention practically:

EXAMPLE 20

[0242] One gram of polystyrene was dissolved in 19 g of liquid naphthalene at 100° C. This naphthalene solution was emulsified at 90°-95° C. into 200 ml of a water solution of polyvinyl alcohol (PVA) (4% by weight) containing 0.1% of Tween-40 emulsifier. The emulsifying head was a Polytron PT-3000 at about 10,000 rpm. Then the emulsion was diluted under agitation with 500 ml of the same aqueous phase at 15° C. whereby the naphthalene droplets solidified into beads of less than 50 microns as ascertained by passing through a 50 micron mesh screen. The suspension was centrifugated under 1000 g and the beads washed with water and recentrifugated. This step was repeated twice.

[0243] The beads were resuspended in 100 ml of water with 0.8 g of dissolved lactose and the suspension was frozen into a block at -30° C. The block was thereafter evaporated under about 0.5-2 Torr between about -20° and -10° C. Air filled microballoons of average size 5-10 microns and controlled porosity were thus obtained which gave an echographic signal at 2.25 and 7.5 MHz after being dispersed in water (3% dispersion by weight). The stability of the microballoons in the dry state was effective for an indefinite period of time; once suspended in an aqueous carrier liquid the useful life-time for echography was about 30 min or more. Polystyrene being non-biodegradable, this material was not favored for injection echography but was useful for digestive tract investigations. This Example clearly establishes the feasibility of the method of the invention.

EXAMPLE 21

[0244] A 50:50 copolymer mixture (0.3 g) of DL-lactide and glycolide (Du Pont Medisorb) and 16 mg of egg-lecithin were dissolved in 7.5 ml of CHCl_3 to give solution (1).

[0245] A solution (2) containing 20 mg of paraffin-wax (M.P. 54°-56° C) in 10 ml of cyclooctane (M.P. 10-13°) was prepared and emulsified in 150 ml of a water solution (0.13% by weight) of 0.13% by weight of Pluronic F-108 (a block copolymer of ethylene oxide and propylene oxide) containing also 1.2 g of CHCl_3 . Emulsification was carried out at room temperature for 1 min with a Polytron head at 7000 rpm. Then solution (1) was added under agitation (7000 rpm) and, after about 30-60 sec, the emulsifier head was replaced by a helical agitator (500 rpm) and stirring was continued for about 3 hrs at room temperature (22° C.). The suspension was passed through a 50 micron screen and frozen to a block which was subsequently evaporated between -20° and 0° C under high-vacuum (catching trap -60° to -80° C). There were thus obtained 0.264 g (88%) of air-filled microballoons stable in the dry state.

[0246] Suspensions of said microballoons in water (no stabilizers) gave a strong echographic signal for at least one hour. After injection in the organism, they biodegraded in a few days.

EXAMPLE 22

[0247] A solution was made using 200 ml of tetrahydrofuran (THF), 0.8 g of a 50:50 DL-lactide/glycolide copolymer (Boehringer AG), 80 mg of egg-lecithin, 64 mg of paraffin-wax and 4 ml of octane. This solution was emulsified by adding slowly into 400 ml of a 0.1% aqueous

solution of Pluronic F-108 under helical agitation (500 r.p.m.). After stirring for 15 min, the milky dispersion was evaporated under 10-12 Torr 25° C. in a rotavapor until its volume was reduced to about 400 ml. The dispersion was sieved on a 50 micron grating, then it was frozen to -40° C. and freeze-dried under about 1 Torr. The residue, 1.32 g of very fine powder, was taken with 40 ml of distilled water which provided, after 3 min of manual agitation, a very homogeneous dispersion of microballoons of average size 4.5 microns as measured using a particle analyzer (Mastersizer from Malvern). The concentration of microballoons (Coulter Counter) was about 2×10^9 /ml. This suspension gave strong echographic signals which persisted for about 1 hr.

[0248] If in the present example, the additives to the membrane polymer are omitted, i.e., there is used only 800 mg of the lactide/glycolide copolymer in the THF/octane solution, a dramatic decrease in cell-wall permeability is observed, the echographic signal of the dispersion in the aqueous carrier not being significantly attenuated after 3 days.

[0249] Using intermediate quantities of additives provided beads with controlled intermediate porosity and life-time.

EXAMPLE 23

[0250] There was used in this Example a polymer of formula defined in embodiment 1 in which the side group has formula (II) where R^1 and R^2 are hydrogen and R is tert.butyl. The preparation of this polymer (defined as poly-POMEG) is described in U.S. Pat. No. 4,888,398.

[0251] The procedure was like in Example 14, using 0.1 g poly-POMEG, 70 ml of THF, 1 ml of cyclooctane and 100 ml of a 0.1% aqueous solution of Pluronic F-108. No lecithin or high-molecular weight hydrocarbon was added. The milky emulsion was evaporated at 27° C./10 Torr until the residue was about 100 ml, then it was screened on a 50 micron mesh and frozen. Evaporation of the frozen block was carried out (0.5-1 Torr) until dry. The yield was 0.18 g because of the presence of the surfactant. This was dispersed in 10 ml of distilled water and counted with a Coulter Counter. The measured concentration was found to be 1.43×10^9 microcapsules/ml, average size 5.21 microns as determined with a particle analyzer (Mastersizer from Malvern). The dispersion was diluted 100x, i.e., to give about 1.5×10^7 microspheres/ml and measured for echogenicity. The amplitude of the echo signal was 5 times greater at 7.5 MHz than at 2.25 MHz. These signals were reproducible for a long period of time.

[0252] Echogenicity measurements were performed with a pulse-echo system consisting of a plexiglas specimen holder (diameter 30 mm) with a 20 micron thick Mylar acoustic window, a transducer holder immersed in a constant temperature water bath, a pulser-receiver (Accutron M3010JS) with an external pre-amplifier with a fixed gain of 40 dB and an internal amplifier with gain adjustable from -40 to +40 dB and interchangeable 13 mm unfocused transducers. A 10 MHz low-pass filter was inserted in the receiving part to improve the signal to noise ratio. The A/D board in the IBM PC was a Sonotek STH 832. Measurements were carried out at 2.25, 3.5, 5 and 7.5 MHz.

[0253] If in the present Example, the polymer used is replaced by lactic-lactone copolymers, the lactones being

δ -butyrolactone, δ -valerolactone or ϵ -caprolactone (see Fukuzaki et al., J. Biomedical Mater. Res. 25 (1991), 315-328), similar favorable results were obtained. Also in a similar context, polyalkylcyano-acrylates and particularly a 90:10 copolymer poly(DL-lactide-co-glycolide) gave satisfactory results. Finally, a preferred polymer is a poly(DL-lactide) from the Company Boehringer-Ingelheim sold under the name "Resomer R-206" or Resomer R-207.

EXAMPLE 24

[0254] Two-dimensional echocardiography was performed using an Acuson-128 apparatus with the preparation of Example 15 (1.43×10^9 /ml) in an experimental dog following peripheral vein injection of 0.1-2 ml of the dispersion. After normally expected contrast enhancement imaging of the right heart, intense and persistent signal enhancement of the left heart with clear outlining of the endocardium was observed, thereby confirming that the microballoons made with poly-POMEG (or at least a significant part of them) were able to cross the pulmonary capillary circulation and to remain in the blood-stream for a time sufficient to perform efficient echographic analysis.

[0255] In another series of experiments, persistent enhancement of the Doppler signal from systemic arteries and the portal vein was observed in the rabbit and in the rat following peripheral vein injection of 0.5-2 ml of a preparation of microballoons prepared as disclosed in Example 15 but using poly(DL-lactic acid) as the polymer phase. The composition used contained 1.9×10^8 vesicles/ml.

[0256] Another composition prepared also according to the directions of Example 15 was achieved using poly(tert-butylglutamate). This composition (0.5 ml) at dilution of 3.4×10^8 microballoons/ml was injected in the portal vein of rats and gave persistent contrast enhancement of the liver parenchyma.

EXAMPLE 25

[0257] A microballoon suspension (1.1×10^9 vesicles/ml) was prepared as disclosed in Example 12 (resin=polystyrene). One ml of this suspension was diluted with 100 ml of 300 mM mannitol solution and 7 ml of the resulting dilution was administered intragastrically to a laboratory rat. The animal was examined with an Acuson-128 apparatus for 2-dimensional echography imaging of the digestive tract which clearly showed the single loops of the small intestine and of the colon.

We claim:

1. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a freon.

2. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is an organic compound containing one or more carbon atoms and fluorine.

3. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a halogenated hydrocarbon.

4. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection the ultrasound contrast agent of claim 1; and,

(2) echographically imaging said body.

5. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection the ultrasound contrast agent of claim 2; and,

(2) echographically imaging said body.

6. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection the ultrasound contrast agent of claim 3; and,

(2) echographically imaging said body.

7. A method of making the ultrasound contrast agent of claim 1 comprising the step of: utilizing a gas that is a freon.

8. A method of making the ultrasound contrast agent of claim 2 comprising the step of: utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

9. A method of making the ultrasound contrast agent of claim 3 comprising the step of: utilizing a gas that is a halogenated hydrocarbon.

10. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a freon.

11. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is an organic compound containing one or more carbon atoms and fluorine.

12. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a halogenated hydrocarbon.

13. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 10; and,
- (2) echographically imaging said body.

14. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 11; and,
- (2) echographically imaging said body.

15. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 12; and,
- (2) echographically imaging said body.

16. A method of making the ultrasound contrast agent of claim 10 comprising the step of:

utilizing a gas that is a freon.

17. A method of making the ultrasound contrast agent of claim 11 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

18. A method of making the ultrasound contrast agent of claim 12 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

19. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is a freon.

20. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is an organic compound containing one or more carbon atoms and fluorine.

21. An ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: air, CO₂ and nitrogen; and a second compound that is a halogenated hydrocarbon.

22. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 19; and,
- (2) echographically imaging said body.

23. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 20; and,
- (2) echographically imaging said body.

24. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 21; and,
- (2) echographically imaging said body.

25. A method of making the ultrasound contrast agent of claim 19 comprising the step of:

utilizing a gas that is a freon.

26. A method of making the ultrasound contrast agent of claim 20 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

27. A method of making the ultrasound contrast agent of claim 21 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

28. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a freon.

29. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is an organic compound containing one or more carbon atoms and fluorine.

30. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a halogenated hydrocarbon.

31. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 28; and,
- (2) echographically imaging said body.

32. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast agent made from the dry formulation of claim 29; and,
- (2) echographically imaging said body.

33. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast agent made from the dry formulation of claim 30; and,
- (2) echographically imaging said body.

34. A method of making the dry formulation of claim 28 comprising the step of:

utilizing a gas that is a freon.

35. A method of making the dry formulation of claim 29 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

36. A method of making the dry formulation of claim 30 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

37. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is a freon.

38. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is an organic compound containing one or more carbon atoms and fluorine.

39. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: air, CO₂ and nitrogen; and a second compound that is a halogenated hydrocarbon.

40. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 37; and,
- (2) echographically imaging said body.

41. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 38; and,
- (2) echographically imaging said body.

42. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 39; and,
- (2) echographically imaging said body.

43. A method of making the dry formulation of claim 37 comprising the step of:

utilizing a gas that is a freon.

44. A method of making the dry formulation of claim 38 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

45. A method of making the dry formulation of claim 39 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

46. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a freon.

47. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is an organic compound containing one or more carbon atoms and fluorine.

48. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a halogenated hydrocarbon.

49. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection a contrast agent made from the dry formulation of claim 46; and,

(2) echographically imaging said body.

50. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection a contrast agent made from the dry formulation of claim 47; and,

(2) echographically imaging said body.

51. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection a contrast agent made from the dry formulation of claim 48; and,

(2) echographically imaging said body.

52. A method of making the dry formulation of claim 46 comprising the step of:

utilizing a gas that is a freon.

53. A method of making the dry formulation of claim 47 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

54. A method of making the dry formulation of claim 48 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

55. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is a freon.

56. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: CO₂, nitrogen, N₂O, methane and butane; and a second compound that is an organic compound containing one or more carbon atoms and fluorine.

57. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microbubbles in a physiologically compatible aqueous carrier,

wherein said suspension comprises a film-forming surfactant that is a phospholipid, and

wherein the gas in the microbubbles is a mixture of a first compound selected from the group consisting of: air, CO₂ and nitrogen; and a second compound that is a halogenated hydrocarbon.

58. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 55; and,

(2) echographically imaging said body.

59. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 56; and,

(2) echographically imaging said body.

60. A method of imaging an organ in a living body, said method comprising the steps of:

(1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 57; and,

(2) echographically imaging said body.

61. A method of making the dry formulation of claim 55 comprising the step of:

utilizing a gas that is a freon.

62. A method of making the dry formulation of claim 56 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

63. A method of making the dry formulation of claim 57 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

64. An ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a freon.

65. An ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is an organic compound containing one or more carbon atoms and fluorine.

66. An ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a halogenated hydrocarbon.

67. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 64; and,
- (2) echographically imaging said body.

68. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 65; and,
- (2) echographically imaging said body.

69. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 66; and,
- (2) echographically imaging said body.

70. A method of making the ultrasound contrast agent of claim 64 comprising the step of: utilizing a gas that is a freon.

71. A method of making the ultrasound contrast agent of claim 65 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

72. A method of making the ultrasound contrast agent of claim 66 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

73. An ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a mixture of a first compound selected from the group consisting of: air, CO₂ and nitrogen; and a second compound that is a halogenated hydrocarbon.

74. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection the ultrasound contrast agent of claim 73; and,
- (2) echographically imaging said body.

75. A method of making the ultrasound contrast agent of claim 73 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

76. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a freon.

77. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier, wherein the gas in the microballoons is an organic compound containing one or more carbon atoms and fluorine.

78. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a halogenated hydrocarbon.

79. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 76; and,
- (2) echographically imaging said body.

80. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 77; and,
- (2) echographically imaging said body.

81. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 78; and,
- (2) echographically imaging said body.

82. A method of making the dry formulation of claim 76 comprising the step of:

utilizing a gas that is a freon.

83. A method of making the dry formulation of claim 77 comprising the step of:

utilizing a gas that is an organic compound containing one or more carbon atoms and fluorine.

84. A method of making the dry formulation of claim 78 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

85. A dry formulation which, upon mixing with an aqueous carrier phase, will generate an ultrasound contrast agent comprising:

a suspension of gas filled microballoons in a physiologically compatible aqueous carrier,

wherein the gas in the microballoons is a mixture of a first compound selected from the group consisting of: air, CO₂ and nitrogen; and a second compound that is a halogenated hydrocarbon.

86. A method of imaging an organ in a living body, said method comprising the steps of:

- (1) administering to said body by injection an ultrasound contrast made from the dry formulation of claim 85; and,
- (2) echographically imaging said body.

87. A method of making the dry formulation of claim 85 comprising the step of:

utilizing a gas that is a halogenated hydrocarbon.

* * * * *

专利名称(译)	超声造影剂及其制备和使用方法		
公开(公告)号	US20010024640A1	公开(公告)日	2001-09-27
申请号	US09/748120	申请日	2000-12-27
[标]申请(专利权)人(译)	SCHNEIDER MICHEL DANIEL BICHON BUSSAT PHILIPPE 皮吉涅JEROME HYBL EVA 闫峰 GRENIER PASCAL GARCEL NADINE BARRAU MARIE贝尔纳黛特		
申请(专利权)人(译)	SCHNEIDER MICHEL DANIEL BICHON BUSSAT PHILIPPE 皮吉涅JEROME HYBL EVA 严峰 GRENIER PASCAL GARCEL NADINE BARRAU MARIE-贝尔纳黛特		
当前申请(专利权)人(译)	SCHNEIDER MICHEL DANIEL BICHON BUSSAT PHILIPPE 皮吉涅JEROME HYBL EVA 严峰 GRENIER PASCAL GARCEL NADINE BARRAU MARIE-贝尔纳黛特		
[标]发明人	SCHNEIDER MICHEL BICHON DANIEL BUSSAT PHILIPPE PUGINIER JEROME HYBL EVA YAN FENG GRENIER PASCAL GARCEL NADINE BARRAU MARIE BERNADETTE		
发明人	SCHNEIDER, MICHEL BICHON, DANIEL BUSSAT, PHILIPPE PUGINIER, JEROME HYBL, EVA YAN, FENG GRENIER, PASCAL GARCEL, NADINE BARRAU, MARIE-BERNADETTE		

IPC分类号	A61B6/00 A61B8/00 A61K49/22
CPC分类号	A61B8/481 A61K49/223
优先权	1990810262 1990-04-02 EP 1992810046 1992-01-24 EP 1990810367 1990-05-18 EP PCT/EP1991/000620 1991-04-02 WO 07/775989 1991-11-20 US
外部链接	Espacenet USPTO

摘要(译)

气相填充的微泡和微球悬浮液在水相中可用作超声波回波描记术中的成像造影剂。通过使用在水中溶解度的填充气体，在标准条件下以升升的气体升的气体表示的气体填充气体，可以在压力作用下对这些气体填充的微气泡和用作超声波回波描记术中的造影剂的微球提供出色的抗塌陷性能除以平方根的分子量不超过0.003。公开了具有特定气体混合物的造影剂，其具有有利的性质。还公开了可用于制造造影剂的干燥制剂。

Fig 1

