

COMPOSITIONS FOR DELIVERING HIGHLY WATER SOLUBLE DRUGS

BACKGROUND OF THE INVENTION

Field of the Invention

This invention relates to an emulsion composition for delivering
5 highly water-soluble drugs such as vinca alkaloids.

Description of the Related Art

Despite years of research into the development of more effective
and safe therapeutic agents for cancer, cancer drugs remain extremely toxic to
patients. The common systemic toxicities associated with cancer drugs, such
10 as chemotherapeutic agents, include bone marrow suppression, asthenia,
peripheral neuropathy, dyspnea, alopecia, *etc.* To make things worse, many
intravenously injected cancer drugs can cause local reactions at the injection
site resulting in vein irritation, pain, tissue necrosis and/or thrombophlebitis.
The consequence of injection site reactions to a chemotherapy agent may
15 include extreme pain, early termination of drug treatment, wounds that are
difficult to heal, scars, permanent tissue damage, and, in the worst case,
amputation.

Alkaloids isolated from the periwinkle plant (*Vinca rosea*) and
derivatives thereof, collectively referred to as "vinca alkaloids," have proven
20 effective as first line therapy for many types of lymphomas, leukemias, and
other cancers. Vincristine and vinblastine consist of a catharanthine moiety
linked to vindoline, and the structures differ by a single substitution in the
vindoline group. Vindesine, a desacetyl carboxamide derivative of vinblastine,
was synthesized later. Subsequently, novel synthetic approaches were used to
25 generate compounds that differed from the natural compounds by the presence
of an eight- rather than a nine-member catharanthine ring, including vinorelbine,

which is commonly available as a tartrate salt, *i.e.*, vinorelbine bitartrate or vinorelbine tartrate.

Vinca alkaloids are highly cytotoxic drugs that disrupt microtubules, inhibit cell division and induce apoptosis. Without wishing to be
5 bound to a particular theory, it is believed that vinca alkaloids exert their cytotoxic effects by binding to tubulin, the protein subunit of microtubules.

Vincristine, vinblastine and vinorelbine are the best-known members of this drug family and are widely used clinically. Despite having similar structures and mechanisms of action, vinca alkaloids differ in their
10 antitumor activity and toxicities. For example, vincristine is used mostly to treat hematological cancer, and neurotoxicity is dose limiting. In contrast, vinorelbine is approved for the use as a single agent to treat breast and non-small cell lung cancer, and its injection site reaction is most severe amongst all vinca alkaloid drugs.

15 It is well known that all vinca alkaloid drugs are associated with adverse reactions at the injection site. For example, the current vinorelbine product approved in the U.S. (NAVELBINE[®]) has a "blackbox" warning due to its severe reaction at the injection site.

NAVELBINE[®] is reportedly associated with a high incidence
20 (51%-61%) of local reactions at the injection site, characterized by injection site pain and phlebitis. The injection site reaction or extravasation of NAVELBINE[®] can be severe, ranging from considerable pain, irritation, and tissue necrosis to thrombophlebitis (The NAVELBINE[®] Product Information by GlaxoSmithKline).

NAVELBINE[®] (vinorelbine tartrate) is a simple solution formulation
25 for intravenous administration. Each vial contains vinorelbine tartrate equivalent to 10 mg (1-mL vial) or 50 mg (5-mL vial) vinorelbine in water for injection at pH 3.5.

Rittenberg reported post-incident care and management of venous irritation or phlebitis due to NAVELBINE[®] (*Oncol. Nurs. Forum* 22: 707-
30 10, 1995). Mare reported methods for preventing venous toxicity of

NAVELBINE[®] by co-administering anti-inflammatory drugs as defibrotide or ketorolac, or changing infusion schedules from a bolus infusion to a slow infusion (*Support Care Cancer* 11: 593-6, 2003). However, the injection site toxicity problem of vinorelbine or other vinca alkaloids has not been properly
5 addressed from the drug formulation approach and vinca alkaloid products remain the most venous toxic drugs.

Oil-in-water emulsion formulations may provide advantages over a traditional solution formulation such as the one used by NAVELBINE[®] in control of venous toxicity at injection site for irritating drugs. For example, the
10 intramuscular or intravenous injection of erythromycin or clarithromycin in a solution formulation causes severe pain at the injection site, and erythromycin or clarithromycin fat emulsion (oil-in-water) is locally non-irritating (WO 90/14094). However, oil-in-water emulsion formulations are typically applied to only lipophilic drugs such as propofol, diazepam, erythromycin or
15 clarithromycin, *etc.* Desai (U.S. Pat. No. 4,816,247) disclosed emulsion compositions for administration of sparingly water soluble ionizable hydrophobic drugs.

Without wishing to be bound to a particular theory, it is believed that in an oil-in-water emulsion, a lipophilic drug is preferentially dissolved in the
20 oil phase and therefore is coated and/or encapsulated in the oil droplets, thus preventing direct contact of drug molecules at a high concentration with the venous endothelium tissue, thus reducing the venous toxicity of the drug.

However, to date, the utility of an emulsion in preventing venous toxicity of irritating drugs is limited to only lipophilic (or hydrophobic) drugs since
25 highly water-soluble drugs, such as vinca alkaloid drugs, do not partition well in the conventional emulsion oil droplets. For example, vinorelbine in the bitartrate salt form is highly soluble in water with an aqueous solubility is >1000 mg/mL in distilled water.

Thus, there remains a need in the art for developing emulsion compositions for delivering highly water soluble drugs. The present invention fulfills such a need and provides other related advantages.

BRIEF SUMMARY OF THE INVENTION

5 The present invention provides compositions for delivering highly water soluble drugs and methods for using such compositions.

 In one aspect, the present invention provides emulsion compositions for delivering highly water soluble drugs that comprises a triglyceride oil, an emulsifier, a stabilizer, and water, wherein the composition is
10 an emulsion having an oil and an aqueous phase, and the drug is substantially in the oil phase.

 In certain embodiments, the highly water soluble drug is venous toxic and/or weakly basic absent the emulsion. In certain embodiments, the drug is selected from the group consisting of dopamine, ciprofloxacin,
15 vancomycin, norvancomycin, doxorubicin, daunorubicin, vinca alkaloids (e.g., vinorelbine), and pharmaceutically acceptable salts thereof.

 In certain embodiments, the triglyceride oil is a triglyceride having long chain fatty acids, a triglyceride having medium chain fatty acids, or a mixture thereof.

20 In certain embodiments, the emulsifier is egg lecithin, soy lecithin, a synthetic phospholipid, or a mixture thereof.

 In certain embodiments, the stabilizer is a fatty acid (e.g., oleic acid), riboflavin-5-phosphate, vitamin-E succinate, cholesterol sulfate, or a mixture thereof.

25 In certain embodiments, the drug has an aqueous solubility of at least or over 30, 50, 75, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, or 1000 mg/ml.

 In certain embodiments, the charge ratio of the drug to the stabilizer is in a range of 1:1 to 1:10. In certain embodiments, the charge ratio

of the drug to the stabilizer is within the range of 1:1 to 1:5, 1:1 to 1:2, or 1:1 to 1:1.1.

In certain embodiments, no less than 80%, 85%, 90%, 92%, 94%, 95%, 96%, 97%, 98%, or 99% of the drug is present in the oil phase of the
5 emulsion.

In certain embodiments, the drug in the emulsion composition is in a concentration range of about 1 to about 50 mg/ml. In certain embodiments, the concentration of the drug in the emulsion is about 2, 3, 4, 5, 6, 7, 8, 9, 10, 15, 20, 25, 30, 35, 40, 45 or 50 mg/ml.

10 In another aspect, the present invention provides a composition for delivering a highly water soluble, venous toxic and weakly basic drug that comprises an oil-in-solid dispersion prepared by freeze-drying the emulsion described herein.

In another aspect, the present invention provides a lyophilized
15 formulation of a highly water soluble, venous toxic and weakly basic drug, wherein the formulation, when hydrated, produces the emulsion described herein.

In certain embodiments, when the lyophilized formulation is reconstituted in a liquid medium to provide particles, the particles increase in
20 size by less than one-fold as compared to particles before lyophilization.

In another aspect, the present invention provides a method for treating cancer comprising administering to a patient in need thereof the compositions described herein wherein the drug in the composition is a vinca alkaloid (e.g., vinorelbine bitartrate). In certain embodiments, the composition
25 is administered intravenously. In certain other embodiments, the composition is administered intramuscularly or intraarterially.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows the effects of various stabilizers on vinorelbine incorporation into the oil phase of an emulsion.

Figure 2 shows the effects of concentrations of sodium oleate on the incorporation of vinorelbine into an emulsion.

Figure 3 shows average tumor growth curves of a vinorelbine emulsion according to Example 8 in comparison with a vinorelbine solution (MINNUOBIN[®] - Example 11) and dextrose solution (5%), in mice xenografted with human breast cancer (Bcap-37). The treatment details are described in Example 13.

Figure 4 shows the blood drug concentration profile of a vinorelbine emulsion prepared according to Example 8 after administering intravenously the emulsion to rats at three doses.

Figure 5 shows the blood drug concentration profile of a vinorelbine emulsion prepared according to Example 8 in comparison with a vinorelbine solution (MINNUOBIN[®] - Example 11), after intravenous injection to rats.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides compositions for delivering highly water soluble drugs (including those that are also venous toxic and/or weakly basic). The pharmaceutical compositions are oil-in-water emulsions containing sub-micron size oil droplets, and comprise triglyceride oil, stabilizers, emulsifiers, and water. The compositions may optionally comprise preservatives and/or other inactive ingredients.

The term of "highly water-soluble drugs," as used herein, refers to a drug (in its freebase or salt form) having solubility in water in excess of 30 mg/ml at room temperature (20-25°C). The solubility of a drug may be described in a variety of ways. The USP/NF generally expresses the solubility in terms of the volume of solvent required to dissolve 1 gram of the drug at a specified temperature (e.g., 1 g aspirin in 300 ml H₂O, 5 ml ethanol at 25°C). Other references may use more subjective terms to describe solubility, such as those given in the following table from Remington's Pharmaceutical Sciences, Mack Publishing Co., Easton, PA, latest edition.

Descriptive terms	Parts of solvent needed for 1 part solute
Very soluble	< 1
Freely soluble	1-10
Soluble	10-30
Sparingly soluble	30-100
Slightly soluble	100-1000
Very slightly soluble	1000-10,000
Practically insoluble or insoluble	> 10,000

Therefore, the "highly water-soluble drugs" of this invention include the drugs in the top 3 solubility categories, *i.e.*, "very soluble," "freely soluble," and "soluble."

The term of "venous toxic drugs," as used herein, refers to drugs which, when intravenously injected in a solution formulation, can cause local reactions at the injection site that result in vein irritation, pain, tissue necrosis and/or thrombophlebitis.

The term "weakly basic drugs," as used herein, refers to drugs having at least one weakly basic functional group.

A "highly water-soluble, venous toxic and weakly basic drug" is commonly provided in a salt form. Examples of some commercial drugs, which fall into this category include, but are not limited to, dopamine hydrochloride, ciprofloxacin lactate, vancomycin hydrochloride, norvancomycin hydrochloride, doxorubicin hydrochloride, daunorubicin hydrochloride, vincristine sulfate, vindestin sulfate, vinblastine sulfate, and vinorelbine bitartrate.

In certain embodiments, the highly water-soluble drugs for use in this invention are anti-neoplastic agents.

In other embodiments, the highly water-soluble drugs for use in this invention are vinca alkaloids, and the pharmaceutically acceptable salts

and derivatives thereof. Vinca alkaloids include, but are not limited to, vincristine, vinblastine, vindesine and vinorelbine.

In yet further embodiments, the highly water soluble drug for use in this invention is vinorelbine, and the pharmaceutically acceptable salts thereof.

Pharmaceutically acceptable salts," as used herein, refers to those salts which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of humans and lower animals without undue toxicity, irritation, allergic response, and the like, commensurate with a reasonable benefit/risk ratio, and effective for their intended use in the chemotherapy and prophylaxis of cancers. Among the more common pharmaceutically acceptable salts of vinca alkaloids are the tartrate, sulfate and hydrochloride forms. Other acid salts used in the pharmaceutical arts include adipate, acetate, bromide, mesylate, lactate, succinate, maleate, lactobionate, aspartate, benzoate, benzene sulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, gluconate, glycerophosphate, heptonate, hexanoate, hydrobromide, hydroiodide, 2-hydroxy ethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, pamoate, pantothenate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, thiocyanate, tosylate, and undecanoate.

In certain embodiments, the vinorelbine salt is the bitartrate salt having a chemical name is 3',4'-didehydro-4'-deoxy- C'-norvincal leukoblastine [$R-(R^*, R^*)$ -2,3-dihydroxybutanedioate (1:2)(salt)].

As used herein, the term "oil-in-water emulsion" refers to a colloidal dispersion systems in which liquid oil is dispersed in small droplets (the discrete phase) in an aqueous medium (the continuous phase), wherein in excess of 80% of the drug is dissolved and remains in the oil droplets. In certain embodiments, greater than 85%, 90%, 92%, 94%, 95%, 96%, 97%, 98%, or 99% of the drug is present in the oil phase.

As used herein, the term "an acidic pH" is meant the pH measured for the emulsion to be in the range of pH of 2 to 6. In certain embodiment, the emulsion has a pH of 3 to 5, pH 3 to 4, or pH 3.5 to 4.0.

In certain embodiments, the pH of the emulsion of the present invention may be in the range of 2 to 7 (e.g., 3 to 7), including any values therebetween.

As used herein, the term "sub-micron size droplet" is meant oil droplets in an oil-in-water emulsion having an average diameter of less than 1 micron as measured by conventional sizing techniques such as laser light scattering spectrometry. In certain embodiments, the oil droplets of the compositions of the present invention have an average diameter of less than 500, 450, 400, 350, 300, or 250 nm. Oil droplets of sub-micron size are desired for the safe passage of these droplets in the capillary blood vessel in the circulation. Droplets of greater than 5 micron in diameter are believed to be unsafe for intravenous injection since they may block the capillary vessel resulting in pulmonary embolism. In certain embodiments, the oil droplets of the compositions of the present invention have an average diameter of less than 0.2-micron (200 nm) so the emulsion may be sterilized by filtering through a 0.2 micron sized filter membrane.

In certain embodiments, the oil droplets of the compositions of the present invention have an average diameter of less than about 150, 100, 75, 50, 25, 20, 15, or 10 nm.

"Triglyceride oil," as used herein, refers to a triglyceride composition which is liquid at room temperature (20-25°C), and which comprises primarily triglycerides of C6 to C18 fatty acids. Triglyceride oil is used in this invention to form the discrete phase of the emulsion or the oil droplets in which the drug is encapsulated. The triglyceride oil is thus desired to be non-toxic, biocompatible, and capable of forming stable droplets of the desired size and encapsulating the highly water-soluble drugs.

The triglyceride oil used in this invention can be glycerol esters of short chain (C4 to C6), medium chain (C8-C12), or long chain (C14 to C18) fatty acids or mixture thereof.

In certain embodiments, triglyceride oils of medium chain fatty acids may be used. Such oils comprise predominantly glycerol triesters of C8 to C12 fatty acids. These oils can be prepared synthetically by well-known techniques, or can be obtained from natural sources by known techniques of thermal or solvent fractionation of suitable natural oils, such as palm oil or coconut oil, to yield fractions rich in the desired low-melting triglycerides. An exemplary low-melting, low molecular weight triglyceride oil is a low molecular weight fraction of coconut or palm oil which is rich in mixed esters of caprylic (octanoic) and capric (decanoic) acids. Such oil is commercially available as Miglyol 812 from SASOL GmbH Germany, CRODAMOL GTCC-PN from Croda Inc. of Parsippany, New Jersey, or Neobees M-5 oil from PVO International, Inc., of Boonton, New Jersey. Other low-melting medium chain oils may also be used in the present invention.

In certain embodiments, triglyceride oils with a high percentage of glycerol triesters of unsaturated or polyunsaturated C14 to C18 fatty acids (long chain fatty acids) may be used. An example of such an oil is soybean oil, which typically has a fatty acid composition of about 80% oleic and linoleic acids. An injectable grade of soybean oil is commercially available as Super Refined USP grade oil from Croda Inc. of Parsippany, New Jersey. Another example of such an oil is safflower oil. Other low-melting vegetable oils or low-melting fractions of oils, including cottonseed, menhaden, olive, peanut, corn, sesame and flaxseed oil, which can be obtained by conventional thermal or solvent fractionation, may also be used in the present invention. While such unsaturated or polyunsaturated vegetable oils may offer a cost advantage in formulating compositions according to this invention, they also exhibit a greater tendency to oxidative deterioration, and may require the addition of oil soluble antioxidants, such as tocopherols.

The triglyceride oil is generally present in a range of from about 2 to about 40% in the final emulsion formulation. In certain embodiments, triglyceride oil is present at about 5%, 10 %, 15%, 20%, 25%, 30%, or 35% by weight in the final emulsion formulation.

5 In certain embodiments, the triglyceride oil comprises a 1:1 weight ratio mixture of a medium chain triglyceride and a long chain triglyceride.

As used herein, the term "stabilizers" refers to those ingredients that retain the highly water-soluble drug in the oil droplets of an oil-in-water emulsion.

10 In other embodiments, the "stabilizers" comprise compounds selected from groups of fatty acids, cholesterol sulfate, riboflavin-5-phosphate, and vitamin E succinate or a mixture thereof.

Exemplary fatty acids include saturated fatty acids, monoenoic acids and polyenoic acids. The saturated fatty acids include, but are not limited to, those listed in the table below:

Systematic name	Common name	Shorthand designation	Molecular wt.
butanoic	Butyric	4:0	88.1
pentanoic	Valeric	5:0	102.1
hexanoic	Caproic	6:0	116.1
octanoic	Caprylic	8:0	144.2
nonanoic	Pelargonic	9:0	158.2
decanoic	Capric	10:0	172.3
dodecanoic	Lauric	12:0	200.3
tetradecanoic	Myristic	14:0	228.4
hexadecanoic	Palmitic	16:0	256.4
heptadecanoic	margaric (daturic)	17:0	270.4
octadecanoic	Stearic	18:0	284.4
eicosanoic	Arachidic	20:0	412.5
docosanoic	Behenic	22:0	340.5

Systematic name	Common name	Shorthand designation	Molecular wt.
tetracosanoic	Lignoceric	24:0	368.6
hexacosanoic	Cerotic	26:0	396.7
heptacosanoic	Carboceric	27:0	410.7
octacosanoic	Montanic	28:0	424.8
triacontanoic	Melissic	30:0	452.9
dotriacontanoic	Lacceroic	32:0	481
tritriacontanoic	ceromelissic (psyllic)	33:0	495
tetratriacontanoic	Geddlic	34:0	509.1
pentatriacontanoic	Ceroplactic	35:0	523.1

Exemplary monoenoic fatty acids include those listed in the table

below:

Systematic name	Common name	Shorthand designation	Molecular wt.
cis-4-decenoic	Obtusilic	10:1(n-6)	170.3
cis-9-decenoic	Caproleic	10:1(n-1)	170.3
cis-5-lauroleic	Lauroleic	12:1(n-7)	198.4
cis-4-dodecenoic	Linderic	12:1(n-8)	198.4
cis-9-tetradecenoic	myristoleic	14:1(n-5)	226.4
cis-5-tetradecenoic	Physeteric	14:1(n-9)	226.4
cis-4-tetradecenoic	Tsuzuic	14:1(n-10)	226.4
cis-9-hexadecenoic	palmitoleic	16:1(n-7)	254.4
cis-6-octadecenoic	petroselinic	18:1(n-12)	282.4
cis-9-octadecenoic	oleic	18:1(n-9)	282.4
cis-11-octadecenoic	vaccenic (asclepic)	18:1(n-7)	282.4
cis-9-eicosenoic	Gadoleic	20:1(n-11)	310.5

Systematic name	Common name	Shorthand designation	Molecular wt.
cis-11-eicosenoic	Gondoic	20:1(n-9)	310.5
cis-11-docosenoic	Cetoleic	22:1(n-11)	338.6
cis-13-docosenoic	Erucic	22:1(n-9)	338.6
cis-15-tetracosenoic	Nervonic	24:1(n-9)	366.6

Exemplary polyenoic fatty acids include those listed in the table

below:

Systematic name	Common name	Shorthand designation	Molecular wt.
9,12-octadecadienoic	linoleic	18:2(n-6)	280.4
6,9,12-octadecatrienoic	linolenic	18:3(n-6)	278.4
8,11,14-eicosatrienoic	dihomolinolenic	20:3(n-6)	306.5
5,8,11,14-eicosatetraenoic	arachidonic	20:4(n-6)	304.5
7,10,13,16-docosatetraenoic	-	22:4(n-6)	332.6
4,7,10,13,16-docosapentaenoic	-	22:5(n-6)	330.6
9,12,15-octadecatrienoic	-	18:3(n-3)	278.4
6,9,12,15-octadecatetraenoic	stearidonic	18:4(n-3)	276.4
8,11,14,17-eicosatetraenoic	-	20:4(n-3)	304.5
5,8,11,14,17-eicosapentaenoic	EPA	20:5(n-3)	302.5
7,10,13,16,19-docosapentaenoic	DPA	22:5(n-3)	330.6
4,7,10,13,16,19-docosahexaenoic	DHA	22:6(n-3)	328.6
5,8,11-eicosatrienoic	Mead acid	20:3(n-9)	306.5

5 In certain embodiments, the fatty acid used is oleic acid.

The amount of stabilizers in the emulsions of this invention may be defined by a drug-to-stabilizer charge ratio, which is calculated as:

Charge ratio = Net positive charge of the drug/Net negative charge of the stabilizer

The amount of stabilizers in the emulsions is within a drug-to-stabilizer charge ratio range of 1:1 to 1:10. In certain embodiments, the charge ratio of the drug to the stabilizer is within the range of 1:1 to 1:5, 1:1 to 1:2, or 1:1 to 1:1.1.

As used herein, the term "emulsifiers" refers to compounds that allow the formation of a stable oil-in-water emulsion wherein the droplets are of sub-micron size and contain the highly water-soluble drug. Exemplary emulsifiers include compounds selected from phospholipids, bile salts, polyoxyethylene sorbitan fatty acid esters (e.g., TWEENS), polyoxyethylene castor oil derivatives (e.g., CREMOPHOR), albumin and poloxamer (e.g., PLURONIC), and mixtures thereof.

A "stable oil-in-water emulsion" refers to an oil-in-water emulsion wherein more than 50% of the oil droplets in the emulsion do not increase their size more than one-fold under appropriate storage conditions for at least 3 months.

In certain embodiments, the emulsion of the present invention is stable for at least 4, 5, 6, 7, 8, 9, 10, 11, 12, 14, 16, 18, 20, 22, or 24 months.

In certain embodiments, the average size of the oil droplets in the emulsion of the present invention does not increase by about 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 100%, 120%, 140%, 160%, 180%, 200%, or 250%.

In further embodiments, phospholipids may be used as emulsifiers. Phospholipids are available from naturally occurring sources or by organic synthesis. Lecithin is a naturally occurring mixture of the diglycerides of stearic, palmitic, and oleic acids, linked to the choline ester of phosphoric acid, commonly called phosphatidylcholine. Hydrogenated lecithin is the product of controlled hydrogenation of lecithin.

According to the United State Pharmacopoeia (USP), lecithin is a non-proprietary name describing a complex mixture of acetone-insoluble phospholipids, which consists primarily of phosphotidylcholine, phosphotidylethanolamine, phosphotidylserine and phosphotidylinositol, combined with various amounts of other substances such as triglycerides, fatty acids, and carbohydrates.

Pharmaceutically, lecithins are mainly used as dispersing, emulsifying, and stabilizing agents and are included in intramuscular and intravenous injections, parenteral nutritional formulations and topical products.

- 10 Lecithin is also listed in the FDA Inactive Ingredients Guide for use in inhalations, intramuscular and intravenous injections, oral capsules, suspensions and tablets, rectal, topical, and vaginal preparations.

Phospholipids can also be synthesized and the common synthetic phospholipids are listed below:

15 Diacylglycerols

- 1,2-Dilauroyl-sn-glycerol (DLG)
- 1,2-Dimyristoyl-sn-glycerol (DMG)
- 1,2-Dipalmitoyl-sn-glycerol (DPG)
- 1,2-Distearoyl-sn-glycerol (DSG)

20 Phosphatidic Acids

- 1,2-Dimyristoyl- sn-glycero-3-phosphatidic acid, sodium salt (DMPA,Na)
- 1,2-Dipalmitoyl- sn-glycero-3-phosphatidic acid, sodium salt (DPPA,Na)
- 1,2-Distearoyl- sn-glycero-3-phosphatidic acid, sodium salt (DSPA,Na)

Phosphocholines

- 1,2-Dilauroyl-sn-glycero-3-phosphocholine (DLPC)
- 1,2-Dimyristoyl-sn-glycero-3-phosphocholine (DMPC)
- 1,2-Dipalmitoyl-sn-glycero-3-phosphocholine (DPPC)
- 5 1,2-Dipalmitoyl-sn-glycero-3-phosphocholine (DPPC)
- 1,2-Distearoyl-sn-glycero-3-phosphocholine (DSPC)
- 1,2-Distearoyl-sn-glycero-3-phosphocholine (DSPC)

Phosphoethanolamines

- 1,2-Dilauroyl-sn-glycero-3-phosphoethanolamine (DLPE)
- 10 1,2-Dimyristoyl-sn-glycero-3-phosphoethanolamine (DMPE)
- 1,2-Dipalmitoyl-sn-glycero-3-phosphoethanolamine (DPPE)
- 1,2-Distearoyl-sn-glycero-3-phosphoethanolamine (DSPE)

Phosphoglycerols

- 1,2-Dilauroyl-sn-glycero-3-phosphoglycerol, sodium salt (DLPG)
- 15 1,2-Dimyristoyl-sn-glycero-3-phosphoglycerol, sodium salt (DMPG)
- 1,2-Dimyristoyl-sn-glycero-3-phospho- sn-1-glycerol, ammonium salt
(DMP-sn-1-G,NH₄)
- 1,2-Dipalmitoyl- sn-glycero-3-phosphoglycerol, sodium salt (DPPG,Na)
- 1,2-Distearoyl- sn-glycero-3-phosphoglycerol, sodium salt (DSPG,Na)
- 20 1,2-Distearoyl- sn-glycero-3-phospho- sn-1-glycerol, sodium salt
(DSP-sn-1G,Na)

Phosphoserines

- 1,2-Dipalmitoyl- sn-glycero-3-phospho-L-serine, sodium salt (DPPS,Na)

Mixed Chain Phospholipids

- 25 1-Palmitoyl-2-oleoyl- sn-glycero-3-phosphocholine (POPC)
- 1-Palmitoyl-2-oleoyl- sn-glycero-3-phosphoglycerol, sodium salt (POPG,Na)
- 1-Palmitoyl-2-oleoyl- sn-glycero-3-phosphoglycerol, ammonium salt
(POPG,NH₄)

Lysophospholipids

1-Palmitoyl-2-lyso- sn-glycero-3-phosphocholine (P-lyso-PC)

1-Stearoyl-2-lyso- sn-glycero-3-phosphocholine (S-lyso-PC)

Pegylated Phospholipids

5 N-(Carbonyl-methoxypolyethyleneglycol 2000)- MPEG-2000-DPPE

1,2-dipalmitoyl- sn-glycero-3-phosphoethanolamine, sodium salt

N-(Carbonyl-methoxypolyethyleneglycol 5000)- MPEG-5000-DSPE

1,2-distearoyl- sn-glycero-3-phosphoethanolamine, sodium salt

N-(Carbonyl-methoxypolyethyleneglycol 5000)- MPEG-5000-DPPE

10 1,2-dipalmitoyl- sn-glycero-3-phosphoethanolamine, sodium salt

N-(Carbonyl-methoxypolyethyleneglycol 750)- MPEG-750-DSPE

1,2-distearoyl- sn-glycero-3-phosphoethanolamine, sodium salt

N-(Carbonyl-methoxypolyethyleneglycol 2000)- MPEG-2000-DSPE

1,2-distearoyl- sn-glycero-3-phosphoethanolamine, sodium salt

15

In certain embodiments, the phospholipids in the formulations of the present invention may be egg or soy lecithin.

The amount of phospholipids, by weight, in the emulsions of the present invention may be within a range of about 2% to about 15%, such as at
20 about 5% to about 10%.

As used herein, the term "bile salts" refers to salts of bile acid, *i.e.*, steroids having 1-3 hydroxyl groups and a five-carbon atom side chain terminating in a carboxyl group, which can be conjugated to glycine or taurine. Bile salts include, but are not limited to, cholate, deoxycholate,
25 chenodeoxycholate, or ursodeoxycholate, and their glycine or taurine conjugates, *e.g.*, glycodeoxycholate (GDC), glycocholate (GC), or taurodeoxycholate (TDC).

As used herein, the term "preservatives" refers to agents that can prevent microbial growth in the emulsion formulation of this invention. The oil-
30 in-water emulsions of this invention contain nutrients for microbes and may thus

be conducive to microbial growth or contamination. Therefore, a preservative may be desirable in the formulation, especially for a vial product that is intended to provide multiple doses where multiple punctures of the vial stopper by syringe needles are needed. The preservatives useful for this invention
5 include, but are not limited to, sodium edetate (EDTA), sodium metabisulfite, sodium benzoate, benzyl alcohol, bronopol, parabens, cresol, phenol, phenoxyethanol, phenylmercuric acetate, phenylmercuric borate, phenylmercuric nitrate, sorbate, benzoate, sorbic acid thimerosal, cetrimide, chlorhexidine, chlorobutanol, chlorocresol, benzalkonium chloride and
10 benzethonium chloride or a mixture thereof.

The other inactive ingredients used in the emulsion compositions of this invention may include virtually any component, such as an acid or base for pH adjustment such as hydrochloric acid and sodium hydroxide, agents to adjust the tonicity of the emulsion including sodium chloride, mannitol, sucrose,
15 dextrose, lactose, polyethylene glycols (PEG) and glycerin or a mixture thereof.

The emulsion formulation of the present invention can be prepared so that it is ready-to-use or can be prepared with a cryoprotectant(s) as a lyophilized solid, *i.e.*, "an oil-in-solid dispersion system" that can be reconstituted at a later date and diluted with water to reform the oil-in-water
20 emulsion before injection.

As used herein, the term "an oil-in-solid dispersion system" refers to a solid matrix prepared by freeze-drying (lyophilizing) an oil-in-water emulsion of this invention and the solid matrix can reform an oil-in-water emulsion of similar droplet size upon mixing with water (reconstitution). In
25 certain embodiments, the average droplet size of the reformed emulsion is no more than about 500%, 300%, or 150% of the average droplet size of the emulsion before the freeze-drying. An oil-in-solid dispersion system of this invention may be optionally prepared by spray drying.

"Cryoprotectants" used in the emulsion compositions of the
30 present invention refers to those ingredients which are added to maintain the

discrete and submicron droplets of the emulsion during the freeze-drying process and, upon the removal of aqueous phase of the emulsion, to provide a solid matrix for the droplets to form the an oil-in-solid dispersion system.

Cryoprotectants that may be used in the emulsion compositions of
5 this invention include, but are not limited to, polyols, monosaccharides, disaccharides, polysaccharides, amino acids, peptides, proteins, and hydrophilic polymers, or mixtures thereof.

Polyols that may be used in the present invention include, but are not limited to, glycerin, mannitol, erythritol, maltitol, xylitol, sorbitol, polyglycitol
10 or mixtures thereof.

Monosaccharides that may be used in this invention include, but are not limited to, glucose, mannose, fructose, lactulose, allose, altrose, gulose, idose, galactose, talose, ribose, arabinose, xylose, lyxose or mixtures thereof.

Disaccharides that may be used in this invention include, but are
15 not limited to, sucrose, lactose, maltose, isomaltose, trehalose, cellubiose or mixtures thereof.

Polysaccharides that may be used in this invention include, but are not limited to, cellulose, amylose, inulin, chitin, chitosan, amylopectin, glycogen, pectin, hyaruronic acid or mixtures thereof.

20 Amino acids that may be used in this invention include, but are not limited to, alanine, arginine, asparagine, aspartic acid, cysteine, glutamine, glutamic acid, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine, valine or mixtures thereof.

25 Peptides that may be used in this invention include, but are not limited to, diglycine and triglycine.

Proteins that may be used in this invention include, but are not limited to, albumin, collagen, casein, and gelatin.

Hydrophilic polymers that may be used in this invention include,
30 but are not limited to, polyethylene glycols povidones, poloxamers, polyvinyl

alcohols or mixtures thereof. The most preferred hydrophilic polymers are polyethylene glycols and povidones.

The concentration of the cryoprotectants used in the liquid emulsion compositions may be in the range of about 2% to about 40% w/w, such as about 5% to about 20% w/w and about 10% to about 15% w/w.

Generally, the emulsion formulation of the present invention can be prepared by performing one or more of the following steps:

- (1) Dissolve the drug, triglyceride oil, emulsifier(s), and stabilizer(s) in a sufficient amount of a volatile solvent, such as ethanol, methylene chloride, chloroform, isopropanol, methanol, tert-butylalcohol, etc. to form a clear solution,
- (2) Remove the solvent to a toxicologically acceptable residual level by vacuum or by blowing solution with a nitrogen or air stream to obtain an oil phase.
- (3) Dissolve the preservative(s), cryoprotectant(s), and other inactive ingredients in water to form an aqueous phase.
- (4) Optionally filter the oil and/or aqueous phase to remove particles.
- (5) Add the aqueous phase to the oil phase, and mix well to form a crude oil-in-water emulsion.
- (6) Adjust pH of the crude emulsion to the desired pH range.
- (7) Pass the crude emulsion through a high pressure homogenizer, such as a Microfluidizer 110F equipped with an emulsion interaction chamber by Microfluidics Corp, MA, operating at approximately 18000 psi pressure for 2 to 10 passages until the emulsion droplets reaches the desired average size range and the emulsion is free of droplets of greater than 5 microns in diameter.
- (8) Aseptically pass the emulsion through a sterile 0.2 micron-rated membrane filter to sterilize the emulsion.

(9) Aseptically fill the filtered emulsion into appropriate sterile containers and seal the containers with appropriate sterile stoppers.

(10) Optionally the emulsions may be freeze-dried to form the oil-in-solid dispersion system.

5 (11) Perform necessary tests on the final emulsion or the oil-in-solid dispersion system.

The compositions of the present invention may be administered to an animal in need thereof via various routes, such as intravenous, intramuscular, intra-articular, intra-peritoneal, or oral administration.

10 The present invention also provides methods for using the compositions described herein. For instance, the present invention provides methods for treating cancer that comprise administering to a patient in need thereof compositions that comprise highly water soluble anti-neoplastic drugs (*e.g.*, vinorelbine bitartrate).

15 The following examples are intended to illustrate the invention without limiting the practice thereof.

EXAMPLES

EXAMPLE 1

PH-STABILITY PROFILING OF VINORELBINE IN SELECTED PH BUFFERS

This study was to determine the range of pH in which vinorelbine is most stable. Vinorelbine solutions at 97.0 µg/mL concentrations were prepared at pH 1.95, 3.05, 5.98, 7.01 and 8.04 in sodium phosphate buffers, and at pH 4.01 and 5.01 in sodium acetate buffers.

The buffered vinorelbine solutions were stored at 40°C and analyzed by HPLC for vinorelbine concentrations at various time points. The recovery of vinorelbine over the initial concentration, representing the stability of vinorelbine in each buffer is shown in Table 1. The pH range in which vinorelbine is most stable was defined based on the maximum recovery.

Table 1

PH STABILITY STUDY OF VINORELBINE AT 97.0 µg/mL AT 40°C/75%

RELATIVE HUMIDITY (RH) -% RECOVERY OVER 0 DAY TIME POINT

Time point (day)	pH 1.95	pH 3.04	pH 4.01	pH 5.01	pH 5.98	pH 7.01	pH 8.04
3	97.9	99.9	98.2	97.5	97.9	96.1	92.1
7	97.3	99.7	100.4	98.6	105.0	94.8	88.0
14	90.2	98.1	97.0	97.6	96.9	91.4	85.1

Conclusion: vinorelbine appeared most stable at pH between 3 and 5, this acidic pH range was thus chosen for the emulsion formulations used in the other examples.

EXAMPLE 2

EFFECT OF STABILIZERS ON VINORELBINE INCORPORATION INTO
THE OIL PHASE OF AN EMULSION

Because of the high solubility of vinorelbine in water, its
5 incorporation in a normal oil phase was found minimal. In other words, without
a stabilizer, vinorelbine is present primarily in the aqueous phase.

The purpose of this study was to determine the effect of selected
stabilizers on vinorelbine incorporation into the oil phase of an emulsion. Since
vinorelbine is a weak base, a stabilizer is preferred to be an acid with lipophilic
10 property. Four stabilizers including sodium oleate, vitamin E succinate,
riboflavin-5-phosphate sodium and cholesterol sulfate were evaluated. Each
stabilizer contains an acid group head and a lipophilic tail, and is generally
considered appropriate for injection.

Emulsions used for this study contained 1.4% (w/w) vinorelbine
15 bitartrate, 10% (w/w) soybean oil, 1.2% (w/w) soy lecithin (Phospholipon® 90G
by PHOSPHOLIPID GmbH), 0.005% (w/w) disodium EDTA, and 2.25% (w/w)
glycerol. The concentration of each stabilizer added into formulation is: 1.57%
(w/w) for sodium oleate, 2.73% (w/w) for vitamin E succinate, 2.46% (w/w) for
riboflavin-5-phosphate sodium and 2.52% (w/w) for cholesterol sulfate. A
20 formulation without any stabilizer ("control") and a formulation with extra
Phospholipon 90G were also evaluated in the study.

The drug incorporation into the oil phase was tested using a
dialysis method developed specifically for this purpose. This method involved
filling 500 mg of an emulsion into a Slide-A-Lyzer dialysis cassette with 10,000
25 MW cutoffs, placing the cassette in 70 mL phosphate buffered saline (PBS), pH
7.4 and shaking the solution on a platform shaker at 100 RPM. A small volume
(1 mL) of PBS was removed at each time point and analyzed by HPLC for
vinorelbine concentration. The time profiles of vinorelbine concentration in PBS
are shown in Figure 1. The stabilizer that provided a high incorporation and

small, uniform and stable emulsion droplets was selected as the preferred stabilizer for the emulsion formulations used in other examples.

Conclusion: The selected sterilizers were able to maintain vinorelbine in the oil droplets as demonstrated by significant reduction in the vinorelbine concentration in PBS, since the droplets are incapable of passing the dialysis membrane due to their size. Vitamin E succinate, sodium oleate and riboflavin-5-phosphate appeared to be the most effective stabilizers.

EXAMPLE 3

EFFECT OF CONCENTRATION OF SODIUM OLEATE ON THE INCORPORATION OF VINORELBINE INTO EMULSION

Having determined sodium oleate being a strong stabilizer, emulsions of the same composition as in Example 2 were prepared with sodium oleate at 1.57% (w/w), 1.18% (w/w), 0.78% (w/w) or 0.39% (w/w), which correspond to a vinorelbine-to-oleate molar ratio of 1:4, 1:3, 1:2 or 1:1, or a charge ratio of 1:1, 1:0.75, 1:0.5 or 1:0.25, respectively. A dialysis test was performed under the same conditions as described in Example 2, and eight time points were taken for twenty-eight hours. PBS samples were analyzed by HPLC for vinorelbine concentration in phosphate buffered saline at each time point. The result is shown in Figure 2. The optimal concentration of sodium oleate was determined based on the incorporation result and emulsion stability.

Conclusion: Sodium oleate added at a vinorelbine-to-oleate molar ratio of 1:4 (charge ratio of 1:1) almost completely stopped the partition of vinorelbine into the aqueous phase.

EXAMPLE 4

STABILITY OF VINORELBINE IN EMULSIONS AT A NEUTRAL PH

Having determined the preferred stabilizer and stabilizer concentration, a preliminary emulsion was prepared and tested for short-term

stability. The purpose of this study was to determine the possibility of a pH neutral emulsion based on vinorelbine stability. The emulsion prepared contained the following composition:

	% (w/w)
Vinorelbine bitartrate	1.389 (equivalent to 10 mg/mL vinorelbine freebase)
Soybean oil	10
Soy lecithin	1.2
Disodium EDTA dihydrate	0.005
Glycerol	2.25
Sodium oleate	1.57
Water QS to	100
pH	7.4

5

Both physical and chemical stability of the emulsion were tested in this experiment. The physical stability was examined by measuring the average droplet size using laser light scattering spectrometry (Malvern zetasizer 3000) and by observing the gross and microscopic appearance of the emulsion. The chemical stability is represented by the vinorelbine stability and was examined by HPLC analysis.

10

Table 2

CHEMICAL STABILITY OF THE EMULSION AT PH 7.4

Time point (day)	Storage condition	% Drug recovered over time 0
7	-20°C	100.3
	25°C	92.4
	40°C	90.9
18	-20°C	98.8
	25°C	93.0
	40°C	89.6

Table 3
PHYSICAL STABILITY OF THE EMULSION AT PH 7.4

Time point (day)	Storage condition	Appearance	AVG droplet diameter (nm)	≥ 5 micron droplets by microscopic examination
0		Uniform	179.2±1.3	None
7	-20°C		7889.7±3682.5	Many
	5°C	Not uniform, some precipitate	178.5±1.3	None
	25°C	Not uniform, some precipitate	180.5±2.0	None
	40°C	Not uniform, some precipitate	189.2±2.2	None
18	-20°C	Phase separation	2695.5±2056.5	Many
	5°C	Not uniform, some precipitate	179.9±0.4	Some
	25°C	Uniform	185.1±2.8	Many
	40°C	Uniform	196.0±0.6	Many

Conclusion: Vinorelbine emulsion at pH 7.4 is not stable both
5 physically and chemically.

EXAMPLE 5

EFFECT OF EMULSION PH ON EMULSION UNIFORMITY AND DROPLET SIZE

After the pH-neutral emulsion (Example 4) failed to show
satisfactory chemical stability, this study was performed to examine stability of
10 emulsions at various pH (Example 1).

In addition, medium chain triglyceride (MCT) was added to
replace 50% of the soybean oil. A brand product of MCT (Miglyol ®812 by
Sasol) was used. The addition of MCT was to improve the emulsion stability
and reduce the droplet size. The combination of soybean oil and MCT at 1:1

ratio has been used in a number of IV fat emulsion product and thus was considered acceptable as a carrier for vinorelbine.

The emulsion formulation contained:

Component	% (w/w)
Vinorelbine	1.39
Soybean oil	5
Miglyol 812N	5
Soy lecithin	1.2
Disodium EDTA dihydrate	0.005
Glycerol	2.25
Sodium oleate	1.57
Deionized water	QS

5 pH of the emulsions was adjusted to 7.03, 5.97, 4.97, 4.52, 3.95 and 3.74 using 0.5N HCl. The physical stability of these emulsions was examined and their appearance is summarized in Table 4. An emulsion with the most uniform in appearance and smallest and most stable droplet size was selected.

10

Table 4

PHYSICAL APPEARANCE OF EMULSIONS AT VARIOUS PH

Emulsion pH	Filterability through 0.2 micron filters	AVG droplet diameter (nm)	Appearance
7.03	Easy	169.8±2.8	Uniform
5.97	Didn't go through 0.2µm	NA	Not uniform
4.97	Easy	420.4±8.3	Not uniform
4.52	Easy	471.9±30.9	Not uniform
3.95	Easy	220.0±1.0	Not uniform
3.74	Easy	191.1±1.6	Uniform

Conclusion: Vinorelbine emulsions of uniform appearance and small droplet size (<200 nm) were only possible at pH 7.0 or pH 3.7.

EXAMPLE 6

PH 3.7 EMULSION STABILITY

A portion of the pH 3.7 emulsion prepared in Example 5 was used to monitor the physical and chemical stability of the emulsion. The emulsion was aliquoted into 2 mL glass vial and stored at -20°C, 25°C and 40°C. Up to 4-week stability results were generated (Tables 5 and 6).

Table 5

PHYSICAL STABILITY OF THE PH 3.74 EMULSION

Time point (day)	Storage condition	AVG Droplet diameter (nm)	Appearance	Microscopic appearance
7	-20°C	121.4±2.7	Uniform	Uniform, No ≥ 5 micron droplets
	25°C	115.2±0.7	Uniform	Uniform, No ≥ 5 micron droplets
	40°C	123.6±8.5	Uniform	Uniform, Some ≥ 5 micron droplets
14	-20	149.0±0.3	Uniform	Uniform, Some ≥ 5 micron droplets
	25°C	112.8±1.3	Uniform	Uniform, No ≥ 5 micron droplets
	40°C	145.2±2.6	Uniform	Uniform, Some ≥ 5 micron droplets
28	-20°C	NA	Not uniform	NA
	25°C	115.3±5.3	Uniform	Uniform, No ≥ 5 micron droplets
	40°C	113.2±2.4	Uniform	Uniform, some small particles

Table 6
CHEMICAL STABILITY OF THE PH 3.75 EMULSION

Time point (day)	Storage condition	Vinorelbine recovered (%)	Vinorelbine Purity (%)	RRT: 1.07-1.25	
				% Impurity#1	% Impurity#2
7	-20°C		98.4	0.93	0.92
	25°C	101.8	98.0	0.68	1.64
	40°C	99.8	97.9	1.21	1.14
14	-20		98.5	0.74	0.68
	25°C	101.1	98.5	0.69	0.67
	40°C	98.2	97.8	0.91	1.06
28	-20°C		97.8	0.91	1.78
	25°C	98.5	98.1	1.03	0.79
	40°C	95.7	96.6	1.92	1.99

Conclusion: The pH 3.7 vinorelbine emulsion appeared physically
5 and chemically stable at 25°C.

EXAMPLE 7

DILUTION STUDY

This study was performed to determine the method by which the emulsion should be diluted for intravenous administration. Since the preferable
10 emulsion is acidic, a neutralization and/or dilution step was tested for purpose of intravenous infusion.

The pH 3.7 liquid emulsion was diluted to 5.0 mg/mL vinorelbine freebase with 53 mM arginine freebase or 50 mM sodium hydroxide at a 1:1 volume ratio. The pH of the diluted/neutralized emulsion was 7.08 with 53 mM
15 arginine and was 7.29 with 50 mM sodium hydroxide. The emulsions were further diluted to 0.5 mg/mL and 2 mg/mL vinorelbine freebase with 5% dextrose solution (D5W) for droplets stability monitoring at room temperature.

In another study, the pH 3.7 emulsion was diluted with D5W or lactated ringer's (LR) injection without arginine or sodium hydroxide as neutralizing agent. Again, the diluted emulsions were evaluated for droplets stability. The stability results of diluted emulsion are shown in Tables 7 and 8.

5

Table 7

EMULSION DROPLET SIZE UPON NEUTRALIZATION AND DILUTION

Sample ID	Neutralize agent	Diluted with D5W to (mg/ml)	pH	ZAve(nm) at time 0	ZAve(nm) at 7 hour
#1	NaOH	0.5	8.05	150.3±5.1	181.4±32.0
#2	NaOH	2.0	7.70	138.9±18.5	120.6±3.5
#3	Arginine	0.5	7.36	136.8±5.9	212.6±6.6
#4	Arginine	2.0	7.29	138.9±18.4	130.8±10.0
#5	NA	Diluted with LR to 0.5	5.05	151.1±10.8	265.5±21.0
#6	NA	Diluted with LR to 2.0	4.68	136.7±1.2	179.2±26.7
#7	NA	Diluted with D5W to 0.5	4.11	124.3±2.2	188.8±0.9
#8	NA	Diluted with D5W to 2.0	3.73	185.4±14.1	147.4±10.3

Table 8

EMULSION APPEARANCE UPON NEUTRALIZATION AND DILUTION

Sample ID	Neutralize agent	Diluted with D5W to (mg/ml)	pH	Microscope check at time 0	Microscope check at 7 hour
#1	NaOH	0.5	8.05	Some ≥ 5 micron droplets	Some ≥ 5 micron droplets
#2	NaOH	2.0	7.70	Some ≥ 5 micron droplets	Some ≥ 5 micron droplets
#3	Arginine	0.5	7.36	Some ≥ 5 micron droplets	Some ≥ 5 micron droplets
#4	Arginine	2.0	7.29	Some ≥ 5 micron droplets	Some ≥ 5 micron droplets
#5	NA	Diluted with LR to 0.5	5.05	No ≥ 5 micron droplets	Some ≥ 5 micron droplets
#6	NA	Diluted with LR to 2.0	4.68	Some ≥ 5 micron droplets	Some ≥ 5 micron droplets

Sample ID	Neutralize agent	Diluted with D5W to (mg/ml)	pH	Microscope check at time 0	Microscope check at 7 hour
#7	NA	Diluted with D5W to 0.5	4.11	No $\geq$ 5 micron droplets	No $\geq$ 5 micron droplets
#8	NA	Diluted with D5W to 2.0	3.73	No $\geq$ 5 micron droplets	No $\geq$ 5 micron droplets

Conclusion: The pH 3.7 vinorelbine emulsion of Example 5 may be diluted with D5W prior to intravenous infusion.

EXAMPLE 8

5 PREPARATION OF VINORELBINE EMULSION FOR STABILITY, VEIN IRRITATION AND ACUTE TOXICITY STUDY

In this example, an emulsion with the following composition was produced:

Component	% (w/w)
Vinorelbine bitartrate	1.4
Miglyol 812 N	15
Soy lecithin	7.5
Disodium EDTA dihydrate	0.005
Oleic acid	1.5
Sucrose	15
Deionized water to QS	100
HCl to adjust pH to	3.5 +/- 0.2

The batch size was 108 mL. The following describes the method of preparation:

A. An oil phase was prepared by dissolving vinorelbine tartrate, Miglyol 812N, Soy lecithin, and oleic acid in a sufficient quantity of dehydrated ethanol to form a clear solution. The ethanol was removed

using a Rotavapor (BÜCHI R-114) to a residual ethanol concentration of <1%.

B. An aqueous phase was prepared by dissolving sucrose and disodium EDTA dihydrate in water for Injection.

5 C. The oil and aqueous phases were mixed together using a Silverson homogenizer (Model L4RT with a 2" head) at 5,000-10,000 RPM for about 5 minutes to form a crude emulsion.

D. The pH of this crude emulsion was adjusted from 3.5 using 1N HCl.

10 E. The crude emulsion was homogenized for six passes in the Microfluidizer Model 110S.

F. In a laminar flow hood, the emulsion was filtered through a 0.45µm filter and then a 0.2µm sterile filter (Sartorius, MiniSart).

15 G. The filtered emulsion was dispensed in 5 mL aliquots into 5 mL pre-sterilized glass vials. These vials were sealed with pre-sterilized rubber stoppers.

EXAMPLE 9

STABILITY OF VINORELBINE EMULSION

The chemical stability of vinorelbine in the emulsion prepared in
20 Example 8 was studied using reverse-phase HPLC method. This method allows determination of concentration and purity of vinorelbine in the emulsion. The vinorelbine chemical stability data of the emulsion are shown in the table below:

Chemical stability

Time point	Storage Condition	Conc. (µg/mL)	% Recovery	% Purity
0	NA	14.0	100.0	98.5
Wk 1	-20°C	14.4	100.0	98.4

Time point	Storage Condition	Conc. ($\mu\text{g/mL}$)	% Recovery	% Purity
	2-8°C	14.6	101.2	98.7
	25°C	14.3	98.9	98.9
	40°C	14.5	100.7	98.6
Wk 2	-20°C	14.3	100.0	99.0
	2-8°C	13.9	97.5	99.3
	25°C	14.1	98.6	98.9
	40°C	14.2	99.5	98.5
Wk 4	-20°C	14.8	100.0	99.0
	2-8°C	14.7	99.4	99.1
	25°C	14.4	97.3	98.7
	40°C	14.6	98.5	97.2
Wk 12	-20°C	14.0	100.0	99.1
	2-8°C	14.0	100.1	99.1
	25°C	14.3	102.7	99.0
	40°C	13.7	98.2	98.9

The physical stability of the emulsion prepared in Example 8 was also evaluated. The physical stability was measured by the ability of the emulsion to maintain its average droplet size and the absence of large droplet (>5 micron in diameter). The average droplet size was determined by laser light scattering using a Malvern Zetasizer 5000, and the presence of large droplet (>5 micron in diameter) was examined by observing the undiluted emulsion using an optical microscope at 400X magnifications. The physical stability data of the emulsion are provided in the table below:

Physical stability

Time point	Storage Condition	AVG Droplet size (nm)	Large droplets (>5 micron in diameter)	Visual appearance
0		115	None	Uniform
Wk 1	-20°C	117	Many	Uniform
	2-8°C	113	None	Uniform
	25°C	120	Some	Uniform
	40°C	146	Some	Uniform
Wk 2	-20°C	126	Many	Uniform
	2-8°C	115	None	Uniform
	25°C	122	Some	Uniform
	40°C	162	Many	Uniform
Wk 4	-20°C	137	Many	Uniform
	2-8°C	110	Some	Uniform
	25°C	123	Some	Uniform
	40°C	169	Some	Uniform
Wk 12	-20°C	201	Many	N/A
	2-8°C	116	None	Uniform
	25°C	143	Some	Uniform
	40°C	117	Some	Yellowish, viscous

Conclusion: Vinorelbine is chemically stable in the emulsion prepared in Example 8 at 2-8°C or 25°C, and the emulsion is physically stable

5 at 2-8°C for at least 12 weeks (3 months).

EXAMPLE 10

FREEZE-DRYING VINOIRELBINE EMULSION

This study was to demonstrate the feasibility of converting a liquid emulsion to a freeze-dried emulsion or the "oil-in-solid dispersion system",

5 which is believed more stable than the liquid emulsion.

Both low pH (pH 3.75) and neutral pH (pH 7.14) emulsion formulations were designed and prepared for freeze-drying or lyophilization study.

The formulation contained:

10

	% (w/w)
Vinorelbine bitartrate	1.39
Miglyol 812 N	15
Soy lecithin	7.5
Disodium EDTA dihydrate	0.005
Oleic acid	1.45
Sucrose	15
Deionized water to QS	100

To prepare the freeze-dried emulsions, vinorelbine bitartrate, Miglyol 812, phospholipon 90G and oleic acid were first dissolved in sufficient amount of dehydrated ethanol to form a clear solution. The ethanol was
15 removed using a rotary evaporator under vacuum at room temperature overnight to obtain an oil phase. The oil phase was mixed with an aqueous phase, which contained sucrose and sodium EDTA to form a crude emulsion using a high shear homogenizer. The oleic acid used in this formulation resulted in a low pH emulsion naturally. The crude emulsion was then
20 microfluidized for 6 passages to form a pH 3.75 final emulsion.

A portion of the pH 3.75 crude emulsion was adjusted to pH 7.14 using 0.5N sodium hydroxide. The crude emulsion was then microfluidized for 6 passages to form a pH 7.14 final emulsion. Both emulsions were filtered

through 0.2-micron filters and filled into as 1 mL each in 5 mL vials or 0.2 mL each in 2 mL vials. The height of the fill was about 3-4 mm. The vials were partially stoppered with lyophile stoppers and freeze-dried using a freeze-dryer (Dura-Stop™ mp by FTS System).

- 5 At the completion of the freeze-drying cycle, the freeze-dryer chamber was back filled with nitrogen gas NF to about 95% of atmospheric pressure and then fully stoppered by collapsing the shelves. The stoppered vials were sealed with aluminum crimp seals.

- 10 The dried emulsion or the oil-in-solid dispersion system was white "cakes" with uniform appearance. Prior to testing, the lyophile was reconstituted with deionized water and mixed for 1-2 minutes to re-form the liquid emulsion. The appearance (gross and microscopic) was recorded and the droplet size was determined (Table 9).

Table 9

- 15 VINORELBINE EMULSIONS PREPARED BY RECONSTITUTION OF THE OIL-IN-SOLID DISPERSION SYSTEM WITH DEIONIZED WATER

Emulsion pH	Appearance	Microscopic appearance	Particle size (nm)
3.75	Uniform	No ≥ 5 micron droplets	120.1 $\pm$ 3.0
7.14	Not uniform	A lot of ≥ 5 micron droplets	NA

- 20 Conclusion: The pH 3.7 vinorelbine emulsion may be freeze-dried to form the oil-in-solid dispersion system, and such oil-in-solid dispersion system can form an oil-in-water emulsion with size characteristics similar to the initial emulsion, upon dilution in water.

EXAMPLE 11

VEIN IRRITATION TEST

- 25 The objective of this test was to compare vein irritation of a vinorelbine emulsion of the present invention with a marketed vinorelbine solution product. The vinorelbine emulsion used was prepared as in Example 8

(without the freeze-drying step). The marketed vinorelbine solution (MINNUOBIN[®] marketed by Sino-Sanofi in China) contains 1% vinorelbine tartrate (equivalent to 1% vinorelbine freebase) in water at pH 3.5. This product has the same composition as Navelbine[®], which is marketed in the U.S by
5 GlaxoSmithKline.

Six white rabbits were divided into three groups (two for each group, one male and one female). Each rabbit received daily an intravenous bolus injection through the marginal ear veins consecutively for 5 days. To Group I, 5% dextrose solution (D5W) was injected daily as a negative control.

10 To Group II, MINNUOBIN[®] was administered at a dose of 1.68 mg/kg/day after dilution to 0.3 % vinorelbine freebases in D5W as a positive control.

To Group III, the vinorelbine emulsion prepared as in Example 8 was administered at a dose of 1.68 mg/kg/day either at 1% (undiluted) or
15 after dilution to 0.3 % vinorelbine freebase in D5W.

In Group III treated with the vinorelbine emulsion prepared as in Example 8, no drug-related signs of vein irritation were observed by appearance examination and pathology histology. All observed changes were due to mechanical punctuation during injection. Rabbits in Group II (treated
20 with MINNUOBIN[®]) exhibited signs of mild to medium level vein irritation.

Conclusion: The vinorelbine emulsion of this invention did not cause vein irritation, while the solution formulation resulted in significant vein irritation at the same intravenous dose.

EXAMPLE 12

25 ACUTE TOXICITY TEST

The objective of this test was to compare acute toxicity of a vinorelbine emulsion prepared as in Example 8 with a marketed vinorelbine solution product (MINNUOBIN[®]) in mice.

Mice (100) were divided into five (5) dose groups with 20 animals in each group (10 males and 10 females). MINNUOBIN[®] and the vinorelbine emulsion as described in Example 8 were administered intravenously.

Immediate reactions were observed and acute toxicity was calculated using the

5 Bliss method.

The iv × 1 acute toxicity LD50 values were calculated as:

Vinorelbine solution (MINNUOBIN [®]):	37.56 (34.24-41.21) mg/kg
Vinorelbine emulsion (Example 8):	40.93 (37.75-44.38) mg/kg

Conclusion: There was no statistical difference between these two groups.

10

EXAMPLE 13

ANTI-CANCER EFFICACY TEST

The objective of this test was to compare anticancer activities of a vinorelbine emulsion prepared as in Example 8 with a marketed vinorelbine solution product (MINNUOBIN[®], Example 11) in mice bearing mouse Lewis
15 lung cancer or xenografted with human breast cancer (Bcap-37). The animals were injected intravenously through tail veins with a vinorelbine formulation at two equal toxicity doses, *i.e.*, 1/5 and 1/10 of the LD50 dose as determined in Example 12. Dextrose solution (5%, D5W) was used as negative control. Each formulation was given daily for 6 days. The efficacy was determined by
20 measuring in the tumor size, and average tumor inhibition values were calculated. For each cancer model, two studies were conducted with 10 mice in each treatment group. Figure 3 shows the average tumor growth curves of all treatment groups in the xenografted model. The tables below list the average tumor inhibition values.

25

Model No. 1 - Mice subcutaneously inoculated with mouse Lewis lung cancer

Average Tumor Inhibition (%)					
Vinorelbine emulsion			Vinorelbine solution		
Dose (iv×6qd)	Study #1	Study #2	Dose (iv×6qd)	Study #1	Study #2
7.63mg/kg	89.07% n = 10	85.91% n = 10	8.19mg/kg	87.07% n = 9	84.23% n = 10
3.81mg/kg	61.46% n = 10	60.07% n = 10	4.10mg/kg	57.32% n = 10	56.38% n = 10

Model No. 2 - Nude mice xenografted with human breast cancer (Bcap-37)

Average Tumor Inhibition (%)					
Vinorelbine emulsion (SDP-012)			Vinorelbine solution		
Dose (iv×5qd)	Study #1	Study #2	Dose (iv×5qd)	Study #1	Study #2
7.63mg/kg	86.97% n = 6	88.33% n = 6	8.19mg/kg	81.13% n = 6	82.20% n = 6
3.81mg/kg	67.25% n = 6	70.33% n = 6	4.10mg/kg	64.08% n = 6	67.33% n = 6

5 Conclusion: Vinorelbine emulsion prepared according to Example 8 formulation exhibited a dose-dependent anti-tumor activity against both mouse and human cancers. Its anticancer efficacy appeared statistically equivalent to the vinorelbine solution formulation.

EXAMPLE 14

PHARMACOKINETICS STUDY

The objective of this test was to obtain pharmacokinetics properties of the vinorelbine emulsion prepared as in Example 8 and compare it with a marketed vinorelbine solution product (MINNUOBIN[®], Example 11) in rats.

In Study No. 1, the vinorelbine emulsion was dosed to rats intravenously through tail veins at 2, 4 and 8 mg/kg, blood samples were taken and analyzed using a validated LC-MS method. A total of 36 rats were used with 12 animals for each dose where six were used for blood sampling from 0-6 hr and the other 6 for 6-12 hr after dosing. Blood concentration curves are shown in Figure 4 and pharmacokinetic parameters are listed in the table below:

Dose mg/kg	$t_{1/2\alpha}$ (h)	$t_{1/2\beta}$ (h)	V1 (L/kg)	Vd (L/kg)	CL (L/h/kg)	AUC ⁰⁻²⁴ (ig×h /L)	AUC ^{0-∞} (ig×h /L)	MRT (h)
2	0.32	9.75	4.59	20.98	1.49	1133.74	1404.72	13.77
4	0.28	11.22	3.0	24.14	1.49	2271.48	2717.59	12.36
8	0.38	9.09	3.84	21.89	1.67	4396.36	5070.47	10.1

In Study No. 2, both vinorelbine emulsion (Example 8) vinorelbine solution (MINNUOBIN[®], Example 11) were dosed to rats intravenously through tail veins at 4 mg/kg, blood samples were taken and analyzed using a validated LC-MS method. A total of 24 rats were used with 12 animals for each formulation where six were used for blood sampling from 0-6 hr and the other 6 for 6-12 hr after dosing. Blood concentration curves are shown in Figure 5 and pharmacokinetic parameters are listed in the table below:

5	ID/dose	$t_{1/2\alpha}$ (h)	$t_{1/2\beta}$ (h)	V1 (L/kg)	Vd (L/kg)	CL (L/h/kg)	AUC ⁰⁻²⁴ ($\mu\text{g}\cdot\text{h}/\text{L}$)	AUC ^{0-\infty} ($\mu\text{g}\cdot\text{h}/\text{L}$)	MRT (h)
	Emulsion 4mg/kg	0.28	11.22	3.00	24.14	1.49	2271.48	2717.59	12.36
	Solution 4mg/kg	0.28	11.70	3.88	25.68	1.52	2106.23	2891.99	17.96

Conclusion: Based on a two-compartment model, the estimated elimination half-life of vinorelbine in rats after receiving the vinorelbine emulsion at 2, 4 and 8 mg/kg dose was 9.75, 1.22 and 9.09 h, respectively, and the AUC₀₋₂₄ were 1133.74, 2271.48 and 4396.36 $\mu\text{g}\cdot\text{h}/\text{L}$, respectively. A linear relationship between the AUC and dose was observed. Based on the elimination half-life and AUC values, the emulsion and solution are very comparable pharmacokinetically.

15

EXAMPLE 15

BIODISTRIBUTION STUDY

The objective of this test was to obtain biodistribution properties of the vinorelbine emulsion prepared as in Example 8 and compare it with a marketed vinorelbine solution product (MINNUOBIN[®], Example 11) in rats.

20

In this study, both vinorelbine emulsion (Example 8) vinorelbine solution (MINNUOBIN[®], Example 11) were dosed to rats intravenously through tail veins at 4, various organ samples were taken at 15 min, 2h and 8h after dosing and analyzed using a validated LC-MS method. For each formulation, a total of 18 rats were used with six animals used for each time point.

25

Concentrations of vinorelbine in the organs are listed in the tables below:

15 min after dosing

		Plasma	Heart	Liver	Spleen	Lung	Kidney	Stomach	Skin	Fat	Testis	Ovary	Thymus	Lymph	Muscle	Intestine	Brain
Emulsion	C(μ g/ml or g)	0.85	23.65	15.41	13.87	17.92	34.58	8.26	4.06	2.00	0.28	11.62	4.93	4.71	3.27	6.04	0.22
	$\pm$ SD	0.11	5.41	1.13	2.75	2.74	5.56	1.51	1.11	0.91	0.03	1.91	0.97	0.95	0.52	1.59	0.06
Solution	C(μ g/ml or g)	0.61	27.70	21.69	13.87	29.00	43.75	10.20	3.10	2.16	0.25	17.22	4.31	5.17	5.14	6.03	5.33
	$\pm$ SD	0.13	6.34	7.83	3.37	7.43	10.89	0.92	0.35	0.72	0.07	2.39	0.62	1.25	0.71	1.39	0.51
	P value	0.0079	0.2613	0.0802	0.9996	0.0065	0.0959	0.0229	0.0710	0.7463	0.5364	0.0339	0.2147	0.4933	0.0004	0.9917	3.15E-10

2 hr after dosing

		Plasma	Heart	Liver	Spleen	Lung	Kidney	Stomach	Skin	Fat	Testis	Ovary	Thymus	Lymph	Muscle	Intestine	Brain
Emulsion	C(μ g/ml or g)	0.24	10.12	10.21	17.96	14.74	20.15	6.70	4.80	2.16	0.18	9.03	7.77	5.53	4.08	3.72	0.21
	$\pm$ SD	0.01	2.90	1.87	3.50	2.42	1.46	1.67	0.79	1.33	0.02	0.74	2.27	1.57	0.60	0.28	0.14
Solution	C(μ g/ml or g)	0.16	9.50	10.92	18.43	19.16	17.07	7.76	3.87	1.53	0.22	10.97	5.50	7.79	5.18	4.17	4.36
	$\pm$ SD	0.02	1.28	3.17	4.26	4.48	4.00	1.72	0.77	0.52	0.10	2.49	1.03	2.18	0.78	0.62	0.63
	P value	1.63E-05	0.6390	0.6465	0.8383	0.0596	0.1070	0.3036	0.0678	0.3050	0.5777	0.2659	0.0492	0.0655	0.0205	0.1435	2.20E-08

5

8 hr after dosing

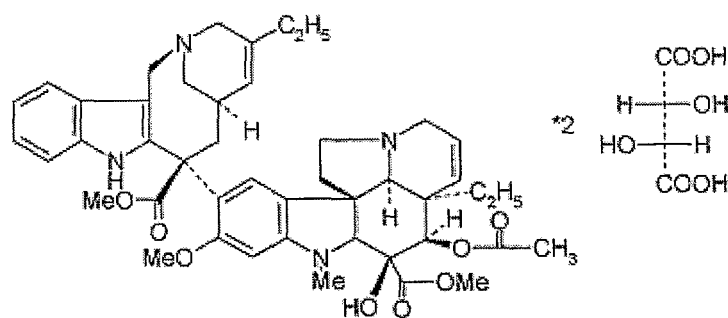
		Plasma	Heart	Liver	Spleen	Lung	Kidney	Stomach	Skin	Fat	Testis	Ovary	Thymus	Lymph	Muscle	Intestine	Brain
Emulsion	C(μ g/ml or g)	0.11	2.91	7.89	18.71	10.69	8.98	5.87	3.81	0.71	0.29	7.36	10.20	6.20	5.61	2.79	0.23
	$\pm$ SD	0.02	0.73	1.44	3.17	3.02	2.18	0.90	0.92	0.07	0.04	1.03	2.09	1.77	1.19	0.50	0.08
Solution	C(μ g/ml or g)	0.10	2.55	4.49	16.42	9.49	8.12	4.85	1.91	0.81	0.27	4.81	7.04	7.57	2.91	2.49	2.13
	$\pm$ SD	0.01	0.45	1.56	2.38	1.87	3.24	0.50	0.51	0.22	0.07	2.02	1.92	2.25	0.89	0.93	0.86
	P value	0.1605	0.3362	0.0029	0.1864	0.4297	0.6028	0.0354	0.0013	0.3450	0.7104	0.1230	0.0212	0.2655	0.0013	0.5023	3.06E-04

Conclusion: The emulsion and solution exhibited generally comparable biodistribution profiles. The emulsion appeared to prevent vinorelbine from entering into brain, which may suggest a possibility of a reduced neurotoxicity such as peripheral neuropathy and constipation commonly associated to the solution formulation.

CLAIMS

What is claimed is:

1. A composition comprising a triglyceride oil, an emulsifier, a stabilizer, water, and a highly water soluble drug, wherein
 - (a) the composition is an emulsion having an oil and an aqueous phase, and
 - (b) the drug is substantially in the oil phase.
2. The composition of claim 1 wherein the emulsion is at a pH range of 3 to 7.
3. The composition of claim 1 wherein the drug alone is venous toxic.
4. The composition of claim 1 wherein the drug is weakly basic.
5. The composition of claim 1 wherein the drug is selected from the group consisting of dopamine, ciprofloxacin, vancomycin, norvancomycin, doxorubicin, daunorubicin, vinca alkaloids, and pharmaceutically acceptable salts thereof.
6. The composition of claim 1 wherein said drug is vinorelbine bitartrate having the chemical name of 3',4'-didehydro-4'-deoxy- C '-norvincal leukoblastine [*R* -(*R* *, *R* *)-2,3-dihydroxybutanedioate (1:2)(salt)] and the following structure:



or another pharmaceutically acceptable salt of vinorelbine.

7. The composition of claim 1 wherein the triglyceride oil is a triglyceride having long chain fatty acids, a triglyceride having medium chain fatty acids, or a mixture thereof.

8. The composition of claim 1 wherein the emulsifier is egg lecithin, soy lecithin, a synthetic phospholipid, or a mixture thereof.

9. The composition of claim 1 wherein the stabilizer is a fatty acid, riboflavin-5-phosphate, vitamin-E succinate, cholesterol sulfate, or a mixture thereof.

10. The composition of claim 9 wherein the fatty acid is oleic acid or a pharmaceutically acceptable salt thereof.

11. The composition of claim 1 wherein the drug has an aqueous solubility of over 50 mg/ml.

12. The composition of claim 1 wherein the drug has an aqueous solubility of over 100 mg/ml.

13. The composition of claim 1 wherein the drug has an aqueous solubility of over 500 mg/ml.

14. The composition of claim 1 wherein the drug has an aqueous solubility of over 1000 mg/ml.
15. The composition of claim 1 wherein the drug-to-stabilizer charge ratio is in a range of 1:1 to 1:10
16. The composition of claim 1 wherein no less than 90% of the drug is present in the oil phase of the emulsion.
17. The composition of claim 1 wherein no less than 95% of the drug is present in the oil phase of the emulsion.
18. The composition of claim 1, wherein the drug in the composition is in a concentration range of 1 to 50 mg/mL.
19. A composition for delivering a highly-water soluble, venous toxic and weakly basic drug, comprising:
 - (a) drug (in freebase) at a concentration of about 0.01% to about 5.0% (w/v),
 - (b) soybean oil at a concentration of about 5% to about 10% (w/v),
 - (c) medium chain triglyceride at a concentration of about 5% to about 10% (w/v),
 - (d) oleic acid at a concentration of about 1.5% to 7.5% (w/v),
 - (e) egg lecithin at a concentration of about 1.2% to 10% (w/v),
 - (f) sucrose at a concentration of about 10% to 15% (w/v),
 - (g) sodium edetate dehydrate at a concentration of about 0.005% (w/v),
 - (h) sodium hydroxide/hydrochloric acid to adjust pH to pH 3-5,and

- (i) water.

20. A composition for delivering a highly-water soluble, venous toxic and weakly basic drug, comprising:

- (a) drug (in freebase) at a concentration of about 0.01% to about 5.0% (w/v),
 - (b) soybean oil at a concentration of about 5% to about 20% (w/v),
 - (c) oleic acid at a concentration of about 1.5% to 7.5% (w/v),
 - (d) soy lecithin at a concentration of about 1.2% to 10% (w/v),
 - (e) sucrose at a concentration of about 10% to 15% (w/v),
 - (f) sodium edetate dehydrate at a concentration of about 0.005% (w/v),
 - (g) sodium hydroxide/hydrochloric acid to adjust pH to pH 3-5,
- and
- (h) water.

21. A composition of matter for delivering a highly-water soluble, venous toxic and weakly basic drug, comprising:

- (a) drug (in freebase) at a concentration of about 0.01% to about 5.0% (w/v),
 - (b) soybean oil at a concentration of about 5% to about 20% (w/v),
 - (c) oleic acid at a concentration of about 1.5% to 7.5% (w/v),
 - (d) soy lecithin at a concentration of about 1.2% to 10% (w/v),
 - (e) glycerin at a concentration of about 2.5% (w/v),
 - (f) sodium edetate dehydrate at a concentration of about 0.005% (w/v),
 - (g) sodium hydroxide/hydrochloric acid to adjust pH to pH 3-5,
- and

(h) water.

22. The composition for delivering a highly-water soluble, venous toxic and weakly basic drug, comprising:

- (a) vinorelbine bitartrate at a concentration of about 1.0% (w/v),
- (b) medium chain triglyceride at a concentration of about 10% (w/v),
- (c) oleic acid at a concentration of about 1.5% (w/v),
- (d) soy lecithin at a concentration of about 7.5% (w/v),
- (e) sucrose at a concentration of about 15% (w/v),
- (f) sodium edetate dehydrate at a concentration of about 0.005% (w/v),
- (g) sodium hydroxide/hydrochloric acid to adjust pH to about 3.75, and
- (h) water.

23. A composition of matter for delivery by intravenous, intramuscular or intraarticular routes of a highly-water soluble, venous toxic and weakly basic drug, comprising:

- (a) vinorelbine bitartrate at a concentration of about 1.0% (w/v),
- (b) soybean oil at a concentration of about 5% (w/v),
- (c) medium chain triglyceride at a concentration of about 5% (w/v),
- (d) oleic acid at a concentration of about 1.5% (w/v),
- (e) soy lecithin at a concentration of about 7.5% (w/v),
- (f) sucrose at a concentration of about 15% (w/v),
- (g) sodium edetate dehydrate at a concentration of about 0.005% (w/v),

- (h) sodium hydroxide/hydrochloric acid to adjust pH to about 3.75, and
- (i) water.

24. A composition for delivering a highly water-soluble, venous toxic and weakly basic drug, comprising an oil-in-solid dispersion prepared by freeze-drying the liquid emulsion composition according to any one of claims 19-23.

25. A lyophilized formulation of a highly water-soluble, venous toxic and weakly basic drug, wherein the formulation, when hydrated, produces the composition according to any one of claims 19-23.

26. The lyophilized formulation of claim 25 wherein the formulation is reconstituted in a liquid medium to provide particles and wherein the particles increase in size by less than one-fold as compared to particles before lyophilization.

27. A method for treating cancer comprising administering to a patient in need thereof the composition according to any one of claims 1 and 19-21 wherein the drug is a vinca alkaloid or a pharmaceutically acceptable salt thereof.

28. The method according to claim 27 wherein the pharmaceutically acceptable salt of vinca alkaloid is vinorelbine bitartrate.

29. A method for treating cancer comprising administering to a patient in need thereof the composition according to claim 22 or claim 23.

30. The method according to any one of claims 27-29 wherein the composition is administered intravenously.

31. The method according to any one of claims 27-29 wherein the composition is administered intramuscularly or intraarterially.