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ANTICANCER COMPOSITIONS

FIELD OF INVENTION

The present invention relates generally to cancer therapeutics More particularly it is directed to novel pharmaceutical compositions of water insoluble anticancer drugs for therapeutic administration as exemplified by the taxanes which include paclitaxel, docetaxel and their derivatives and analogues.

BACKGROUND AND SUMMARY OF THE INVENTION

Paclitaxel is a taxane and a member of the terpenoid family of compounds present in very small quantities in the Taxus brevifolia species such as the pacific Yew tree. These compounds, collectively known as taxoids, taxins or taxanes, have potent anticancer properties in, among others, ovarian cancer, lymphoma, and breast cancer. Because of its poor solubility in water, the current commercial formulation of paclitaxel is prepared by dissolving 6 mg of the drug in one milliliter of a mixture of polyoxyethylated castor oil (Cremophor®(EL) and dehydrated alcohol. The commercially available paclitaxel formulation is for intravenous administration only. There exists no commercial formulation of paclitaxel, which can be administered orally. The commercial injectable formulation is physically unstable especially for treatments requiring long infusion time. The infusate may contain up to 10% each of alcohol and Cremophor®EL. The physical stability of the paclitaxel formulation may be increased by increasing the amounts of Cremophor®EL in the formulation, but may also lead to an increased incidence of adverse reactions. Yet another approach as described in U.S. patent 5,681,846 is to decrease the drug and Cremophor® concentration and increase the alcohol content in the formulation.

An undesirable effect of Cremophor®EL in paclitaxel and other drug formulations is the production of possible anaphylactoid reaction with associated dyspnea, hypotension, angioedema and uticaria. Cremophor®EL is also known to extract plasticizers such as di-ethylhexyl-phthalate from the polymers commonly used intravenous infusion tubings and infusion bags. These plasticizers are known to promote toxic reactions, such as Adult Respiratory Distress Syndrome (ARDS), in patients which have been exposed to high levels.

Various other methods have been used to increase the water solubility of paclitaxel and other anticancer drugs, for example, by conjugation of the water insoluble drug moiety with

water soluble polymers as taught by U.S. patent 5,437,055, WO 97/10849, and WO 97/33552. While WO 94/12031 teaches that a composition of paclitaxel with Cremophor®EL, absolute alcohol and citric acid increases the stability however, no mention is made if the proposed composition increases the solubility of paclitaxel. Others have used liposome preparations as a means of eliminating Cremophor®EL and reducing vehicle toxicity as described by Sharma et al (Pharm. Res. 11:889-896, 1994). An oil-in-water emulsion (U.S. patent 5,616,330) is another approach to preparing Cremophor® free paclitaxel formulation. The latter two formulation approaches have limitations in terms of low degree of drug loading. Yet another approach uses cyclodextrins to make a water-soluble formulation of paclitaxel as described in WO 94/26728.

The present invention is based on a strong need for a safer and stable injectable and oral formulation of anticancer drugs, particularly the taxanes such as paclitaxel, docetaxel and their derivatives and analogues and other anticancer drugs.

U.S. patent 5,407,683 discloses a composition containing paclitaxel in squalene as solution in absence of a surfactant and then forming a self-emulsifying glass by addition of an aqueous sucrose solution followed by evaporation of water. The resulting glass upon mixing with water forms an emulsion with a particle size in a range of 2 to 10 μm. The preparation of such glass requires the use of undesirable organic solvents, which must be completely removed before medical use.

Quay et al describe a conventional oil-in-water emulsion system (WO 98/30205) consisting of vitamin E as a carrier oil in which a drug may be dissolved, together with polyethyleneglycol and related surfactants. Conventional emulsions have limited shelf life and are often difficult to terminally heat sterilize or even filter sterilize. The particle size of conventional emulsions is usually far greater than microemulsions.

Microemulsions are thermodynamically stable and optically transparent or opaque depending on the particle size of the emulsion. Microemulsions have a mean droplet size of less than 200 nm, in general between 20-100 nm. In contrast to conventional emulsions, the microemulsions are formed in the presence of an aqueous phase by self emulsification without any energy input. In the absence of water, this self emulsifying system exists as a transparent-looking mixture of oil and surfactants in which a lipophilic drug is dissolved.

Wheeler et al describe an emulsion preparation (U.S. patent 5,478,860) containing a mixture of paclitaxel, an oil and a polyethylene glycol-linked lipid which is covered by a

monolayer of a polar lipid such as phosphatidylglycerol or phosphatidylethanolamine. This mixture, after homogenization in presence of an aqueous phase at appropriate pressure, yields an emulsion with a particle size in the range of 100 nm. It is not known if this is the mean or minimum particle size and if it is number weighted or volume weighted. The necessity of using undesirable organic solvents for initial dissolution of ingredients is not advisable even if the organic solvent is removed prior to use. In addition to an elaborate evaporation step, the method requires input of energy by way of high pressure homogenization adding to the overall cost. Because the preconcentrate of a true microemulsion is usually non-aqueous, it can provide longer shelf life than a regular emulsion which is in aqueous suspension.

Lacy et al disclose a capsule delivery system (U.S. patent 5,645.856) for oral delivery of hydrophobic drugs containing a digestible oil, and a combination of surfactants. The selection of surfactant is made such that it inhibits the in vivo lipolysis of the oil.

Eugster discloses an ultra microemulsion system (Swiss Patent CH 688 504 A5) for paclitaxel and its analogs composed of an oil and one or more surfactants providing a formulation of the drug with a mean particle size of 2.2-3 nm thus approaching a solution rather than an emulsion. It is not known if this formulation is useful for oral, injectable or topical use.

There have been attempts to enhance oral activity of taxanes by co-administration of taxanes with another drug such as cinchonine (WO 97/27855) or cyclosporin, ketoconazole etc. (WO 97/15269). Similarly, WO 97/48689 describes the use of various carbocyclic compounds in combination with anticancer drugs to enhance oral bioavailability of the drug. All three of these approaches have the drawback of combination drug therapy where a second drug with drastically different pharmacological activity is administered. In practice such a drug combination approach is the last resort taken by those familiar with the drug development process due to drastic increase in preclinical and clinical regulatory requirement for approval resulting in increasing cost and time to market.

SUMMARY OF THE INVENTION

In accordance with the present invention it has now surprisingly been found that particularly stable anticancer drug formulations, particularly the taxanes, that self emulsify in aqueous medium giving an average particle size in a range of about 10 nm to about 10 microns and that have improved bioavailability characteristics, are obtainable. Also described are self-

emulsifying preconcentrates that disperse, without the input of high energy (i.e., other than mixing energy to cause dispersion), to form droplets of average size of up to about 10 microns.

Accordingly, this invention provides a pharmaceutical composition in the form of a self-emulsifying preconcentrate comprising an anticancer drug as the active ingredient solubilized in a carrier medium comprising at least one hydrophobic component, at least one hydrophilic component and at least one surfactant.

The self-emulsifying systems and their corresponding preconcentrates described in this invention consist of a hydrophobic component, an ingredient selected from triglycerides, diglycerides, monoglycerides, free fatty acids, and fatty acid esters (such as fatty acid esters of hydroxyalkanes or of dihydroxyalkanes) and derivatives thereof, individually or in combination. Preferably the surfactant is a non-ionic surfactant or a mixture of non-ionic surfactants. The invention is also characterized as optionally including a hydrophilic component, for instance a hydroxyalkane such as ethanol and/or a dihydroxyalkane such as 1,2-propylene glycol and/or a polyethylene glycol having an average molecular weight of less than or equal to 1000.

Compositions of the current invention will include, in addition to the water insoluble drug, the hydrophobic components and the optional hydrophilic components, and at least one surfactant. Examples of suitable surfactants are:

1. Polyoxyethylene-sorbitan-fatty acid esters; e.g. mono- and tri-lauryl, palmityl, stearyl and oleyl esters; e.g. products of the type known as polysorbates and commercially available under the trade name "Tween".

2. Polyoxyethylene fatty acid esters, e.g., polyoxyethylene stearic acid esters of the type known and commercially available under the trade name Myrj.

3. Polyoxyethylene castor oil derivatives, e.g., products of the type known and commercially available as Cremophors®. Particularly suitable are polyoxyl 35 castor oil (Cremophor® EL) and polyoxyl 40 hydrogenated castor oil (Cremophor® RH40).

4. α-tocopherol, α-tocopheryl polyethylene glycol succinate (vitamin E TPGS), α-tocopherol palmitate and α-tocopherol acetate .

5. PEG glyceryl fatty acid esters such as PEG-8 glyceryl caprylate/caprate (commercially known as Labrasol), PEG-4 glyceryl caprylate/caprate (Labrafac Hydro WL 1219), PEG-32 glyceryl laurate (Gelucire 44/14), PEG-6 glyceryl mono oleate (Labrafil M 1944 CS), PEG-6 glyceryl linoleate (Labrafil M 2125 CS).

6. Propylene glycol mono- and di-fatty acid esters, such as propylene glycol laurate, propylene glycol caprylate/caprate; also diethyleneglycol-monoethylether (DGME), commercially known as Transcutol (Gattefosse, Westwood, NJ).

7. Sorbitan fatty acid esters, such as the type known and commercially available under the name Span (e.g., Span 20).

8. Polyoxyethylene-polyoxypropylene co-polymers, e.g., products of the type known and commercially available as Pluronic or Poloxamer.

9. Glycerol triacetate.

10. Monoglycerides and acetylated monoglycerides, e.g., glycerol monodicocoate (Imwitor 928), glycerol monocaprylate (Imwitor 308), and mono-and di-acetylated monoglycerides.

Suitable surfactants are not limited to those mentioned above, but may include any compound or compounds that would enhance the galenic properties of the preconcentrate.

Compositions in accordance with the present invention may include other ingredients in addition to the drug, one or more hydrophobic components, one or more hydrophilic components, one or more surfactants, inhibitors of cytochrome P450 enzymes or inhibitors of the p-glycoprotein transport system such as grapefruit extract or compounds isolated from it. The composition may include, in addition to the forgoing, one or more ingredients, additives or diluents such as pharmaceutically acceptable polymeric or inorganic materials, anti-oxidants, preserving agents, flavoring or sweetening agents and so forth.

Compositions in accordance with the present invention may be liquid or solids at ambient temperature. They may be filled in soft or hard gelatin capsules in the form of liquid composition, molten composition, or granules or powder (if composition is solid at ambient temperature and was cooled and processed before filling). Coating may be also applied to capsules or tablets. The preconcentrate may be also be diluted with water to obtain stable emulsions that may be employed as drinking formulations, or packaged as such for injection after appropriate dilution with an aqueous medium, for example.

DETAILED DESCRIPTION OF THE INVENTION

A self-emulsifying preconcentrate of the present invention comprising an anticancer drug must contain a hydrophobic component, a surfactant and optionally a hydrophilic component. The surfactant and hydrophilic component are needed for the composition to form in aqueous medium a self-emulsifying system having an average particle size of between about

10 nm and about 10 microns. They may also help enhance the solubility and stability of the anticancer drug in the formulation. The hydrophobic component is needed because if it is not incorporated in appropriate amounts in the formulation, precipitation of the drug will be observed upon mixing of the composition with an aqueous medium and/or on storage. Similar observations may be made for the hydrophilic and surfactant components.

Based on the above, appropriate combinations or mixtures of a hydrophobic component, a surfactant and a hydrophilic component (when used) with the water insoluble drug are necessary to obtain a stable microemulsion preconcentrate that would yield upon mixing with an aqueous medium a stable dispersion with an average particle size of between about 10 nm and about 10 microns.

Preferred as hydrophobic components are triglycerides, diglycerides, monoglycerides, free fatty acids, and fatty acid esters and derivatives thereof, individually or in combination. Examples of hydrophobic components include but are not limited to propylene glycol dicaprylate/caprate, caprilic/capric triglyceride, caprylic/capric/linoleic triglyceride, e.g. synthetic medium chain triglycerides having C8-12 fatty acid chains or other derivatized (synthetic) triglycerides of the type known and commercially available under Miglyol 810, 812, 818, 829 and 840, linoleic acid, linoleic acid ethyl ester, fish oils as free fatty acids, their esterification and their transesterification products, e.g. of the type known and commercially available under EPAX 6000 FA, EPAX 4510 TG, individually or in combination. Additional examples include vegetable oils and C12-18 fatty acid mono-, di- and triglycerides prepared by individual admixing or as transesterification products of vegetable oils (such as soybean oil, almond oil, sunflower oil, olive oil or corn oil) with glycerol.

Preferred as hydrophilic components are 1,2-propylene glycol, ethanol and polyethylene glycol having an average molecular weight of less than or equal to 1000, individually or in combination. More preferred as hydrophilic components are 1,2-propylene glycol and ethanol, individually or in combination. Especially preferred as hydrophilic components is a combination or mixture of 1,2-propylene glycol and ethanol.

The relative proportion of the drug and the other ingredients in the composition of the current invention will vary depending whether it is delivered as a self-emulsifying preconcentrate or after dilution with water, depending on the particular ingredients and the desired physical properties of the formulation. Especially desired concentration limits in the self-emulsifying preconcentrate are as follows:

1. Oil phase: from 10 to 80% w/w of the preconcentrate. The oil phase may consist of triglycerides, diglycerides, monoglycerides, free fatty acids, propylene glycol mono or diesters and free fatty acids, esters and derivatives thereof, individually or in combination.

2. Cumulative amounts of surfactants: from 20 to 80% w/w of the preconcentrate.

3. Cumulative amounts of hydrophilic components, such as 1,2-propylene glycol and/or ethanol and/or a polyethylene glycol having an average molecular weight of less than or equal to 1000 : from 0% to 40% w/w of the preconcentrate. The total of all ingredients will be 100%.

It is understood that the application of the teachings of the present invention, to the conditions described, will be evident to one skilled in the art of preparing such formulations, and to one skilled in treating such medical conditions. Additional features and advantages of the present invention are described below in preferred embodiments, which are intended as example, and not as limitation. In the following examples, the ingredients were weighed out into appropriate containers in the amounts described below. In all examples described below, a clear liquid was obtained upon appropriate mixing and heating.

EXAMPLES

The formulations represented in the following examples were prepared by mixing the oil components with surfactants and cosurfactants followed by the addition of drug powder as indicated. The composition may be prepared at room temperature or heated to 40-50°C to accelerate the solubilization process. Several mixing techniques can be used ranging from mechanical stirring and agitation to sonication. All compositions shown below give liquid or semi-solid preconcentrates at room temperature.

An experiment to test the efficiency of forming microemulsions from the preconcentrates was carried out by diluting the preconcentrate in 20-50 fold with water or simulated gastric fluid with gentle mixing or shaking. The aqueous medium temperature varied between 20 and 37°C. Particle size analysis was then carried out using a photon correlation spectroscopy based particle sizer, Nicomp 370. Data reported in the following examples correspond to volume weighted particle size.

EXAMPLE 1

Ingredients	Amount (g)
Miglyol 840	1.971
Cremophor® RH40	2.190
Imwitor 308	0.767
Labrasol	0.548
Paclitaxel	0.175
Total	5.651

Mean particle size: 31 nm

EXAMPLE 2

Ingredients	Amount (g)
Miglyol 840	4.820
Cremophor® RH40	4.990
Imwitor 308	1.750
Labrasol	1.250
Paclitaxel	0.489
Transcutol	2.000
Total	15.299

Mean particle size: 13 nm

EXAMPLE 3

Ingredients	Amount (g)
Miglyol 840	1.396
Cremophor® RH40	1.551
Imwitor 308	0.543
Labrasol	0.388
Paclitaxel	0.122
Grapefruit extract	0.400
Total	4.400

Mean particle size: 30 nm.

EXAMPLE 4

Ingredients	Amount (g)
Miglyol 840	1.560
Cremophor® RH40	1.610
Imwitor 308	0.565
Labrasol	0.405
Paclitaxel	0.285
Ethanol	0.575
Total	5.000

Mean particle size: 14 nm

EXAMPLE 5

Ingredients	Amount (g)
Miglyol 812	1.435
Tween 80	2.150
Lipoid E80	0.705
Soybean oil	0.178
Linoleic acid	0.174
Ethanol	0.305
Paclitaxel	0.068
Total	5.000

Mean particle size: 102 nm

EXAMPLE 6

Bioavailability of paclitaxel micro-emulsion preconcentrate was assessed using the formulation described in Example 1. Paclitaxel was given in doses of 2.5 mg/Kg or 5 mg/Kg to 8 male dogs of approximately 10 Kg body weight. The formulation was administered in the morning after overnight fasting in the form of a capsule followed by water. Free access to food and water was allowed two hours after dosing. Blood samples were drawn at different point (pre-dose, 0.5, 1, 2, 3, 4, 6, 8, 12, and 24 hr) and stabilized with EDTA, placed in Vacutainers, and stored at 2-8°C. The blood samples were then extracted using a liquid-liquid method and assayed by HPLC/UV. Bioavailability calculations were done by comparing the pharmacokinetic (PK) profiles obtained for orally given paclitaxel micro-emulsion preconcentrate with an intravenous commercial formulation. Bioavailability values ranging from 25 % to 60 % were obtained. Figure 1 corresponds to a typical pharmacokinetic profile obtained for paclitaxel preconcentrate.

While the invention has been described in connection with what is presently considered to be the most practical and preferred embodiment, it is to be understood that the invention is not to be limited to the disclosed embodiment, but on the contrary, is intended to cover various modifications and equivalent arrangements included within the spirit and scope of the appended claims.

WHAT IS CLAIMED IS:

1. A storage-stable, self-emulsifying, and non-aqueous preconcentrate of a taxane microemulsion comprising a taxane dissolved in a carrier system, which carrier system consists essentially of :

10 to 80% w/w of a hydrophobic component selected from a group consisting of a triglyceride, a diglyceride, a monoglyceride, a free fatty acid, a fatty acid ester, a fish oil, a vegetable oil and combinations thereof;

20 to 80% w/w of surfactant component consisting of at least one non-ionic surfactant,

up to 35% w/w diethylene glycol monoethylether, and

up to 40% w/w of a hydrophilic component selected from the group consisting a hydroxyalkane, a dihydroxyalkane, a polyethylene glycol having an average molecular weight of at most 1000, and combinations thereof;

wherein the preconcentrate, when mixed with water or simulated gastric fluid, gives an average droplet size of at most 10 microns, and a doze of preconcentrate has a taxane bioavailability of 25% to 60% of the taxane in the dose upon oral administration..

2. The preconcentrate of claim 1, wherein the carrier contains 15 to 75% w/w of the hydrophobic component.

3. The preconcentrate of claim 1, wherein the carrier contains up to 30% w/w of the hydrophilic component.

4. A storage-stable, self-emulsifying, and non-aqueous preconcentrate of at least one taxane in a composition consisting essentially of :

10 to 80% w/w of a hydrophobic component selected from the group consisting of a triglyceride, a diglyceride, a monoglyceride, a free fatty acid, a fatty acid ester, a fish oil, a vegetable oil and combinations thereof ;

20 to 80% w/w of a surfactant component consisting of at least one non-ionic surfactant; and

up to 40% w/w of a hydrophilic component selected from the group consisting of a hydroxy alkane, a dihydroxyalkane, a polyethylene glycol having an average molecular weight of at most 1000, 1,2-propylene glycol, ethanol, and combinations thereof;

wherein said preconcentrate, when mixed with water or simulated gastric fluid, gives an average droplet size of at most 10 microns, and a dose of preconcentrate has a taxane bioavailability of 25% to 60% of the taxane in the dose upon oral administration.

5. The preconcentrate of claim 4, wherein the hydrophilic component comprises a combination of 1,2-propylene glycol and ethanol.

6. The preconcentrate of claim 4 or claim 5, wherein the preconcentrate also comprises an inhibitor of P-glycoprotein transport system or an inhibitor of cytochrome P450 enzyme.

7. The preconcentrate of claim 6, wherein the composition comproses grapefruit extract of a component thereof.

8. The preconcentrate of any one of claims 4 – 7, wherein the taxane is paclitaxel or docetaxel.

9. An orally administrable pharmaceutical composition consisting essentially of the preconcentrate of claim 4 in a pharmaceutically acceptable carrier or diluent.

10. A parenterally injectable pharmaceutical composition comprising the preconcentrate of claim 4 in a pharmaceutically acceptable diluent.

11. The preconcentrate of claim 4 filled in a soft or hard gelatin capsule.

12. A storage stable, self-emulsifying, and non-aqueous preconcentrate of a taxane in a microemulsion comprising the taxane dissolved in a carrier system, which carrier system consists essentially of:

10 to 80% w/w of a hydrophobic component;

20 to 80% w/w of a surfactant component consisting of one or more non-ionic surfactants; and

up to 40% of a hydrophilic component.

13. the preconcentrate of claim 12, wherein the preconcentrate forms an emulsion having droplets with an average size of 10 microns or less when mixed with an aqueous medium.

14 the preconcertrate of claim 12 or claim 13, wherein a dose of the preconcentrate has a taxane bioavailability of 25% to 60% upon oral administration.

15. the preconcentrate of claim 13 or claim 14, wherein the preconcentrate forms a liquid having an average droplet size of at most 10 microns when mixed with an aqueous medium and heated to 20-37°C.

16. The preconcentrate of any one of claims 12 – 15, wherein at least a portion of the hydrophilic component consists of ethanol, such that the carrier system contains at least 6% w/w ethanol.

17. The preconcerntrate of any one of claim 12 – 16, wherein the preconcentrate, upon oral adminsitration, forms a microemulsion *in situ* in the gastrointestinal tract.

18. An orally administrable pharmaceutical composition comprising the preconcentrate, of any one of claims 12 – 17 in a pharmaceutically acceptable carrier or diluent.

19. A parentally injectable pharmaceutical composition comprising the preconcentrate of any one of claims 12-17 in a pharmaceutically acceptable carrier or diluent.

20. A storage-stable, self-emulsifying, non-aqueous preconcentrate of a taxane in a microemulsion comprising a taxane dissolve in a carrier system, which carrier system consists essentially of:

10 to 80% w/w of a hydrophobic component;

20 to 80% w/w of a surfactant component; and

up to 40% of a hydrophilic component, at least a portion of which hydrophilic component consists of ethanol, such that the carrier system contains at least 6% w/w ethanol.

21. The preconcentrate of claim 20, wherein the preconcentrate forms an emulsion having droplets with an average size of 10 microns or less when mixed with an aqueous medium.

22. The preconcentrate of clam 20 or 21, wherein a dose of the preconcentrate has a taxane bioavailibilityof 25% to 60% upon oral administration.

23. The preconcentrate of claim 21 or claim 22, wherein the preconcentrate forms a liquid having an average droplet size of at most 10 microns when mixed with an aqueous medium and heated to 20-37°C.

24. The preconcentrate of any one of claims 20-23, wherein the preconcentrate, upon oral administration, forms a microemulsion *in situ* in the gastrointestinal tract.

25. An orally administrable pharmaceutical composition comprising the preconcentrate of any one of claims 20-24 in a pharmaceutically acceptable carrier or diluent.

26. A parenterally injectable pharmaceutical composition comprising the preconcentrate of any one of claims 20-24 in a pharmaceutically acceptable carrier or diluent.

27. A storage-stable, self-emulsifying, and non-aqueous preconcentrate of a taxane in a microemulsion comprising a taxane dissolve in a carrier system, which carrier system consists essentially of:

10 to 80% w/w of a hydrophobic component selected from the group consisting of a triglyceride, a diglyceride, a monoglyceride, a free fatty acid, a fatty acid ester, a fish oil, a vegetable oil, and combination thereof;

20 to 80% w/w of a surfactant component consisting of one or more surfactants selected from the group consisting of a polyoxyethelene-sorbitan-fatty acid ester, a polyoxyethylene fatty acid ester, a polyoxyethylene castor oil derivative, α-tocopherol, α-tocopheryl polyethelene glyco succinate, α-tocopherol palmitate, α-tocopherol acetate, a PEG glyceryl fatty acid ester, a propylene glycol mono- or di-fatty acid ester, a sorbitan fatty acid ester, a polyoxyethelene-polyoxypropylene co-polymer, glycerol triacetate, a monoglyceride, an acetylated monoglyceride, and combinations thereof; and

up to 40% of a hydrophilic component, at least a portion of which hydrophilic component consisting of ethanol, such that the carrier system contains at least 6% w/w ethanol.

28. The preconcentrate of claim 27, wherein the preconcentrate forms an emulsion having droplets with an average size of 10 microns or less when mixed with an aqueous medium.

29. The preconcentrate of claim 27 or claim 28, wherein a dose of the preconcentrate has a taxane bioavailability of 25% to 60% upon oral administration.

30. The preconcentrate of claim 28 or claim 29, wherein the preconcentrate forms a liquid having an average droplet size of at most 10 micron when mixed with an aqueous medium and heated to 20-37°C.

31. The preconcentrate of any one of claims 27-30, wherein the preconcentrate, upon oral administration, forms a microemulsion *in situ* in the gastrointestinal tract.

32. An orally administrable pharmaceutical composition comprising the preconcentrate of any one of claims 27-31 in a pharmaceutically acceptable carrier or diluent.

33. A parentally injectable pharmaceutical composition comprising the preconcentrate of ane one of claims 27-31 in a pharmaceutically acceptable carrier or diluent.

34. A method of orally or parenterally administering a taxane to a subject in need of same comprising administrating a dose of a storage-stable, self-emulsifying, and non-aqueous

preconcentrate of a taxane in a microemulsion comprising a taxane dissolved in a carrier system, which carrier system consists essentially of :

10 to 80% w/w of a hydrophobic component selected from the group consisting of a triglyceride, a diglyceride, a monoglyceride, a free fatty acid, a fatty acid ester, a fish oil, a vegetable oil and combinations thereof ;

20 to 80% w/w of a surfactant component consisting of one or more non-ionic surfactants, and

up to 40% w/w of a hydrophilic component selected from the group consisting of a hydroxyalkane, a dihydroxyalkane, a polyethylene glycol having an average molecular weight of at most 1000, and combinations thereof

wherein the preconcentrate, when mixed with water or simulated gastric fluid, gives an average droplet size of at most 10 microns, and a dose of the preconcentrate has a taxane bioavailability ranging from 25% to 60% of the taxane in the dose upon oral administration.